Kuwait National Action Plan On Antimicrobial Resistance 2022

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Abbreviations and acronyms

AMC	Antimicrobial consumption								
AMR	Antimicrobial resistance Antimicrobial Stewardship								
AMS	Antimicrobial Stewardship College of American Pathologists								
САР	College of American Pathologists								
CDC	Centre for Disease Control Continuous Education								
CE	Continuous Education								
CLSI	Clinical and Laboratory Standards Institute								
СМЕ	Continuous medical education								
CMS	Central Medical Stores								
CRE	Carbapenem resistant Enterobacteriaceae								
EPA	Environment Public Authority								
ESBL	extended-spectrum beta-lactamases								
FAO	Food and Agriculture Organization								
GLASS	Global Antimicrobial Resistance and Use Surveillance System								
HAI	Healthcare -associated infection								
HCWs	Healthcare workers								
HIS	Hospital Information System								
ICD	Infection Control Directorate								
IPC	Infection prevention and control								
КЕРА	Kuwait Environment Public authority								
KIMS	Kuwait Institute of Medical Specializations								
KISR	Kuwait Institute of Scientific Research								
LIS	Laboratory Information System								
MDR	Multidrug resistant								
MDROs	Multidrug resistant organisms								
MOE	Ministry of Education								
МОЕ	Ministry of Education								

МОН	Ministry of Health								
МОНЕ	nistry of Higher Education								
MRSA	Methicillin resistant Staphylococcus aureus								
NAP	National Action Plan								
NFP	National Focal point								
NGOs	Non-Governmental Organizations								
NMCG	National Multisectoral Coordinating Group								
OIE	World Organization of Animal Health								
ToR	Terms of reference								
тот	Training of trainers								
PAAET	The Public Authority of Applied Education and Training								
PAAF	The Public Authority of Agriculture Affairs and Fish Resources								
PAFN	The Public authority for Food and nutrition								
PSD	Pharmaceutical Services Directorate								
WHO	World Health Organization								

Forward

This national action plan for Kuwait to combat antimicrobial resistance has been formulated in the line of the WHO strategic objectives. It addresses the involvement of all the related sectors including human and veterinary medicine, food chain, finance, environment, and the general public consumers. Therefore, the National Committee for the Proper Use of Antimicrobials was involved and technical subcommittees were established to prepare this plan.

Executive summary

This national action plan for Kuwait to combat antimicrobial resistance has been formulated in the line of the WHO strategic objectives. It addresses the involvement of all the related sectors including human and veterinary medicine, agriculture, finance, environment, and the general public consumers.

The AMR NAP was prepared over a period of three months covering the governance of the plan and the structure of the work to be done in AMR awareness, surveillance, infection prevention and control, rational antimicrobial use. Research and sustainability regarding AMR were included in the corresponding pillars.

The preparation of the plan was done through meetings held by the WHO consultant and the WHO national AMR focal person at MOH (Dr. Abeer Aly) with stakeholders from the different ministries and authorities including the MOH, PAAF, and KEPA. The approach to the plan development was based on mapping the current situation regarding each pillar and then building on what is available a new plan to be executed over the coming five years.

Regarding the situation analysis, Kuwait had already started its response regarding the AMR pandemic. Regarding country response and governance, an NMCG has already been formed and an AMR focal person and IPC focal person have already been appointed and are working.

Major work on AMR prior to the plan development was concentrated on the its control in humans.

AMR Awareness

The MOH has already started few activities that improve general public awareness about AMR in human health. Short films, advertisements, SMS, posters, brochures and leaflets have already been prepared and distributed during the activities of National campaigns of proper use of antibiotics performed by the IPC Directorate. During the past 2 years, this material is broadcasted during the World Antimicrobial Awareness Week. Education about AMR is included in some human health-related specialties but not in all and not in veterinary or agriculture professionals' continuous education.

Besides the organization of the focal persons' functions and the inter-ministerial communications, the plan targets general public awareness campaigns and yearlong awareness messages, regarding human and non-human health. AMR-related education is another target of this plan in basic primary education, undergraduate university education, postgraduate studies, as well as continuous

education of professionals. Collaboration with FAO, OIE and other regional/international NGOs that already have AMR education-related programs is strongly encouraged. Drug companies play a major role in promoting the use of antimicrobials, where both MOH and PAAF would demand in the future that these companies provide impartial education about AMR, along with its causes and consequences. The country will actively participate in the World Antimicrobial Awareness Week, however the efforts targeting AMR awareness will be spread throughout the yearly calendar.

AMR Surveillance

Public and private hospitals have well-equipped microbiology laboratories that provide reliable microbiology data. Individual facility-based AMR surveillance is being performed in some hospitals, that do issue their own antibiograms, yet this practice is not available throughout the country. A focus group has already been created at MOH and has started gathering data to be reported to WHO GLASS platform and the AMR data bank is increasing gradually. A central AMR laboratory is not available, yet a CDC laboratory is in the process of being built and its budget has already been assigned. In non-human health sectors, there is no real tangible data about AMR except for few research projects that cannot be counted as representative to these fields.

The NAP started with an organizational effort to link hospitals to the central AMR surveillance unit at MOH, and to empower and enlarge this unit (that is the "GLASS Unit" now) to become the national AMR surveillance unit. This unit would enable hospitals to issue their own local antibiograms to be compiled at a later stage in a national AMR report, that will be periodically reviewed and updated. A central AMR reference laboratory is planned to be present within the new CDC, where confirmation of AMR as well as training and quality control of the laboratories in the country will be organized. Broader and more active participation in GLASS platform data feeding is also anticipated in the NAP, not to mention CAP accreditation for the laboratories in the country. AMR detection and surveillance represent in the non-human health sector is a major priority of the upcoming NAP. Mapping of available AMR data in these fields is planned, as well as improving the veterinary/agriculture laboratory capacity at PAAF to detect and ultimately report and surveil AMR. AMR-related research projects in agriculture, food safety and environment will be prioritized in universities or in grants and funds through KEPA research office.

<u>Infection Prevention and Control</u>

The work in the IPC pillar has been quite advanced in the country with the presence of a central IPC directorate at MOH that coordinates IPC among the different hospitals. All hospitals (public and private) already have local functioning IPC offices with the supervision and coordination by the IPC directorate at MOH. IPC guidelines are available, but some need updating. IPC training is performed centrally and locally in the hospitals. Regarding the structural basis of the IPC in Kuwait, adequate hygiene and sanitation of drinking water is available in the country as well as proper waste management is under the authority of the Ministry of Public Works. Surveillance of healthcare associated infections is well established in hospitals, however, this work is still mostly manual and thus requires important resources in terms of time and workforce. IPC in the veterinary world is applied through biosecurity measures in farms under the authority of PAAF that makes sure that animals are vaccinated according to OIE guidelines. There is an infrastructure related to the safety and quality control of products in the food chain.

Regarding the Governance in this pillar, more organization is required, whereby IPC focal persons in PAAF, KEPA and food safety authorities will be appointed. These functions will be assigned to existing employees and will be added to their TOR. This strategy was proposed by the stakeholders, since it is feasible considering the actual workload and it minimizes extra costs. Although a lot has already been done at MOH, the plan targets further improvement in IPC. In order to improve compliance with IPC principles in healthcare facilities, the plan suggests implementation of IPC through decrees and mandates. On the other hand, an update of the national IPC guidelines was suggested based on the latest WHO guidelines on core components of IPC programmed at the national and acute health care facility level. Since digitalization of hospital data and medical records is currently in process in Kuwaiti hospitals, it was suggested that HAI surveillance be linked directly from the hospitals to IPC directorate. Additionally, availability of a digital surveillance system has been suggested to minimize paper work, improve efficiency and save time. IPC education has already been touched upon in the plan at all levels in the human and non-human health sectors including basic undergraduate and continuous postgraduate education. National IPC guidelines in non-human health sectors including animal health, agriculture and environment will be developed.

Rational Use of Antimicrobials

The regulatory authorities of proper procurement legislation and control of importation and sale of registered medicines are well established in the country and well controlled by MOH authorities. There is a legislation that prohibit selling antimicrobials without a prescription of a certified health

professional. On the inpatient level, post-prescription review for appropriateness of antimicrobial agents 48 to 72 hours after administration is being performed in public hospitals, yet only depending on the presence of qualified personnel and ID physicians. In general, national authorities work with close communication with OIE, FAO and WHO, in addition to being part of GCC.

The organization of the governance of this pillar is one of the priorities in the upcoming NAP through assigning its responsibilities to specific focal persons in MOH, PAAF and KEPA. Other priorities include establishing inter-ministerial communication and putting a high-priority list of antimicrobials that should not be used in veterinary medicine or as crop growth promoters. In human health, antimicrobial stewardship programs in hospitals will be implemented and they are planned to be controlled at a central unit in MOH with clear national antimicrobial usage guidelines that are regularly updated according to national antibiograms and AMR epidemiologic data. In order to achieve these aims, a national task force will be appointed to include professionals from the public and private sectors with the needed academic and scientific expertise. Likewise, antimicrobial use guidelines in the agriculture and veterinary fields are to be put and implemented, and their sale would be exclusively based upon veterinary prescriptions. Finally, training courses and workshops targeting rational antimicrobial use in human and non-human health sectors should be provided periodically.

Background

Introduction

Antimicrobial (including antibiotic, antiviral, antifungal and antiprotozoal) agents are critical tools for fighting infectious diseases in humans, terrestrial and aquatic animals and plants, but they are becoming ineffective.¹

Alarming levels of resistance have been reported in countries of all income levels, with the result that common diseases are becoming untreatable, and lifesaving medical procedures riskier to perform. Drug-resistant infections already cause at least 700,000 deaths globally a year, including 230,000 deaths from multidrug-resistant tuberculosis, a figure that could increase to 10 million deaths globally per year by 2050 under the most alarming scenario if no action is taken. Around 2.4 million people could die in high-income countries between 2015 and 2050 without a sustained effort to contain antimicrobial resistance (AMR).¹

The emergence of AMR is a complex problem driven by many interconnected factors, in particular the use and misuse of antimicrobials. Antimicrobial use, in turn, is influenced by an interplay of the knowledge, expectations and interactions of prescribers and patients, economic incentives, characteristics of the health system and the regulatory environment.²

AMR will affect everybody, regardless of where they live, their health, economic circumstances, lifestyle or behavior. It will affect sectors beyond human health, such as animal health, agriculture, food security and economic development.³

In the light of this complexity, coordinated interventions are needed that simultaneously target the behavior of healthcare providers and patients and change important features of the environments in which they interact.²

Alert to this crisis, the May 2015 World Health Assembly endorsed a global action plan to tackle AMR. All Member States are urged to have in place, national action plans on antimicrobial resistance that are aligned with the global action plan with Whole-of-society engagement including a one health approach.³

Situation analyses and Assessment

AMR in Kuwait is a serious problem resulting in increases in morbidity, mortality and healthcare costs. Similar to other countries, Kuwait faces challenges to combat AMR. Resistant organisms are implicated in several outbreaks in hospitals for example Methicillin resistant *Staphylococcus aureus* (MRSA), Multi drug resistant (MDR) *Acinetobacter* spp, MDR. *Pseudomonas. aeruginosa* and Carbapenem resistant Enterobacteriaceae (CRE).

There is no nationwide surveillance of AMR, data are available from individual studies from some hospitals. The prevalence of extended spectrum beta lactamase (ESBL) & CRE isolates in positive blood cultures of patients in a teaching hospital in Kuwait over a 2-year period were studied. 24.9% of Enterobacteriaceae were ESBL producers and 5.2% were CRE. ⁴

There is an alarming high prevalence (11.3%) of colistin-resistant Enterobacteriaceae among healthy food handlers in community while the fecal carriage rate of CRE was (5.3%)⁵. According to single center study, there was a high burden of *Acinetobacter. baumannii* (*A. baumannii*) related infections among Intensive care units (ICU) patients and multiple outbreaks were detected during the study period.⁶

Data from Kuwait National Healthcare-associated infections surveillance system (KNHSS) ^{7,8} reported alarming results. The most common organisms isolated from Healthcare associated infections (HAIs) in 2019 and 2020 were *Klebsiella pneumoniae*(K.pneumoniae). In 2019 CR K.pneumoniae represented 26.7 % from all K.pneumoniae isolated from different types of HAIs. However, in 2020 it is doubled to reach 50.1 %. *A. baumannii* which is the 4th common organisms causing HAIs in both 2019 and 2020 shows increasing MDR rates. In 2019, MDR A.baumannii represented 82.9 % and increased to reach 88.5 % in 2020.

Kuwait is a low tuberculosis (TB) incidence (~ 24 cases/100 000 population) country with low (~ 1.1%) incidence of MDR-TB. Nearly 85% of all TB cases and > 90% of MDR-TB cases occur in expatriate population. These cases mainly arise due to reactivation of latent TB infection acquired previously by TB patients in their respective countries. ^{9,10}Until recently, rifampicin-resistant TB/MDR-TB among Kuwaiti subjects were infrequently detected and transmission of DR-TB was rarely reported in Kuwait. ¹¹⁻¹⁴ A fourfold increase in the detection of MDR-TB among Kuwaiti subjects was noted in recent years (2014–2017); however, the factors responsible for this sudden increase remained unknown and transmission of MDR-TB within Kuwait was not apparent from routine surveillance studies. ¹⁵

Kuwait has a low-prevalence of HIV-1 infection. However, the detection of major mutations conferring resistance to nucleoside and non-nucleoside reverse transcriptase inhibitors (NNRTIs) in 12.5% of ART-naive patients, and about 30% of treatment-experienced patients. However, the antibiotics in the healthcare setting is a big challenge. In 2008, a national study to assess the physicians, adherence to the local hospital antibiotic guidelines was conducted in 9 Ministry of Health (MOH) hospitals. The results revealed that 25% of the prescriptions were unnecessary (not indicated) while only 30.4% of the prescriptions were fully adherent to the guidelines. However, the detection of major mutations conferring mutations and successful to the prescriptions were fully adherent to the guidelines.

of resistance. Understanding patients' knowledge and practices concerning antibiotics can help to mitigate antibiotic resistance and sustain antibiotic effectiveness. ¹⁸Communication to public on the association between unnecessary antimicrobial use and the emergence and spread of AMR seems an important component of strategies to control AMR. 19-20 A cross-sectional survey was performed using a pretested self-administered questionnaire on a sample of 770 randomly selected Kuwaiti individuals. Nearly three-quarters (72.8%) of respondents had been prescribed antibiotics within 12 months prior to the study period, and 36% of them had not finished the course of treatment. Over one-quarter (27.5%) were self-medicated with antibiotics to treat mainly common cold, sore throat and cough. Self-medication was more prevalent among those who were prescribed antibiotics and those who had attitudes towards using and accessing antibiotic inappropriately. Almost 47% of participants had low knowledge regarding action, use, safety and resistance of antibiotics. Forty one percent of respondents had attitudes towards using and accessing antibiotic inappropriately.²¹ During the activities of Kuwait National Campaign for the Proper Use of Antimicrobials 2019, assessment of the level of knowledge, attitudes and practices towards antibiotic use among public attending Kuwait MOH Healthcare facilities was performed by using a questionnaire. The results revealed insufficient knowledge regarding the antibiotic utilization (median knowledge score was 60%, IQR=80-40) with poor attitude (median attitude score was 40%, IQR=60-20) however, the practice was better (median practice score was 77.8%, IQR= 88.9-55.6)²² The environmental dimension of antimicrobial resistance has received comparatively less focus than AMR in human or animal health. However, the natural environment is an important reservoir of antimicrobial resistance. Drug-resistant microbes are in people, animals, food, and the environment (in water, soil and air)²³. Due to demographic and environmental factors, the marine environment of the Gulf Cooperation Council (GCC) region may be particularly susceptible to the threat of antimicrobial resistance. However, there is currently little information on the presence of AMR in the GCC marine environment to inform the design of appropriate targeted surveillance activities. To date no studies have addressed the issue of AMR in marine systems, which represents a key data gap in Kuwait. A baseline survey that is the first to obtain information on the prevalence of AMR within bacterial isolates collected from Kuwait's marine environment. In GCC, the studies have identified the presence of AMR bacteria in fish and seawater collected from locations close to sewage discharges ²⁴ and within the effluent itself ²⁵. The presence of AMR bacteria has also been used as an indicator to monitor the exposure of green turtles (Chelonia mydas) to different marine pollutants ²⁶

Misuse of antibiotics by public plays an important role in driving the emergence and dissemination

Recently, a study objective was to develop²⁷, implement and conduct a rapid regional baseline monitoring survey of the presence of AMR in the GCC marine environment, through the analysis of seawater collected from high-risk areas across four GCC states: (Bahrain, Oman, Kuwait, and the United Arab Emirates). 560 Escherichia coli strains were analyzed as part of this monitoring programme between December 2018 and May 2019. Multi-drug resistance (resistance to three or more structural classes of antimicrobials) was observed in 32.5% of tested isolates. High levels of reduced susceptibility to ampicillin (29.6%), nalidixic acid (27.9%), tetracycline (27.5%), sulfamethoxazole (22.5%) and trimethoprim (22.5%) were observed. Reduced susceptibility to the high priority critically important antimicrobials: azithromycin (9.3%), ceftazidime (12.7%), cefotaxime (12.7%), ciprofloxacin (44.6%), gentamicin (2.7%) and tigecycline (0.5%), was also noted. A subset of 173 isolates was whole genome sequenced, and high carriage rates of qnrS1 (60/173) and blaCTX-M-15 (45/173) were observed, correlating with reduced susceptibility to the fluoroquinolones and third generation cephalosporins, respectively.

A base line survey study ²⁸ had been conducted. In total, 598 isolates of Escherichia coli (351 seawater; 247 bivalves) were isolated from seawater and biota (Venus clam, Circenita callipyga) across Kuwait coastline, and screened for their potential for resistance against an array of commonly deployed frontline antibiotics frontline (23 antibiotics) by micro-dilution (48 h incubation) onto the custom dehydrated 96-well Sensititre™ GN2F panels (GN2F, Thermo Scientific, UK). Results demonstrate the resistant was widespread across all sites (seawater: summer 89 - 64%; winter 90-57% and biota: summer 77%; winter 88%). Ranking the resistance profiles for seawater and biota across both summer and winter periods suggests that the profile of resistance may be influenced by seasonal factors. For example, in strains of E. coli isolated from winter biota samples displayed a high-level of resistance to FOX (51.4%), which then dropped substantially in samples screened from the summer (7.7%). Likewise, the resistance profiles between seawater and biota samples didn't always mirror each other and could point different drivers, within each matrix, being responsible for the promotion and maintenance of AMR. The dataset available doesn't allow for definitive statements to be made about either of these subjects, but does point to future research lines to follow. This spread of observed resistances to older as well as new antibiotics, encompassing almost all tested classes and including antimicrobials used for a variety of clinical and veterinary applications is of some concern.

National SWOT analysis as per each pillar

Governance and multisectoral "One Health" coordination

Strengths	 A national multisectoral coordinating group (NMCG) is already established that includes high-rank influential employees and professionals of the Kuwaiti government from all sectors. It has strong political support, authority to act. It is accountable to the government. It is formed of and supported by technical experts. This group is responsible to facilitate and coordinate development of the national AMR action plan, to facilitate and oversee implementation, monitoring and evaluation of the AMR action plan, and to ensure regular data collection and information sharing among all relevant sectors and stakeholders. There are a few tools and data to form a basis for preparation of a national action plan on AMR such HAI surveillance in hospitals, reporting AMR to GLASS platform, as well as other audit tools for hand hygiene and other issues related to IPC in hospital settings.
Weaknesses	 Technical working groups are not available in all sectors except for a few in human health. Relevant groups will be formed at a later stage as per this NAP on AMR. Focal persons for the coordination between different ministries and authorities regarding the "One Health" approach are not available except in the MOH.
Opportunities	 A preliminary draft of situation analysis (SWOT), strategic objectives, interventions and activities for the NAP on AMR was put by the NMCG, yet, the operational and monitoring plans are to be developed.
Threats	None

AMR Awareness

Strengths	 Regarding activities that target national awareness of AMR, only the Infection Control Directorate in MOH already prepared AMR-relevant information focusing on human health and this material is diffused to the general public only during the AMR/ABX awareness week. On the public level, there was a study done by the Infection Control Directorate about the estimation of knowledge, attitudes and practice of the general public regarding antibiotic use in humans and the sample was randomly selected individuals from outpatient clinics.
	 Among professional in human health, workshops and educational activities targeting AMR in human health are sometimes offered by MOH, scientific societies, drug companies, hospitals, and universities, but it is not regular or structured. Regarding the inclusion of AMR and related topics in quality assurance programs, only IPC is included in accreditation standards of healthcare facilities.
Weaknesses	 Regarding activities that target national awareness of AMR, public communication programs targeting the general audiences in human health practice, animal health practice, plant production and crops, food chain and in the environmental sector are not available. Material prepared by the Infection Control Directorate MOH targeting the general public (mentioned above) is not diffused all yearlong and does not include AMR information based in the One Health approach. Education and information on AMR is not available in school curricula. There is a weak communication between ministries and authorities regarding the importance of AMR, where each one works alone. There are no undergraduate studies/specialties in animal health and agriculture, in Kuwait. These professionals receive their degrees and training outside the country. For those who should be licensed to work in Kuwait, they should undergo an interview and licensing exam that do not include questions related to AMR. AMR and related topics in special targeted modules are only included in undergraduate curricula in Medical Schools and not in other related fields. AMR and related topics are not mandatory in continuing education programs in any field. CE is not mandatory for obtaining work licenses for newcomers to Kuwait or for license renewal. However, CME is needed for promotion of physicians and dentists. In the veterinary/agriculture fields, there is no central structured training for professionals by PAAF regarding AMR. Some activities related to antibiotic use are offered by drug companies for promotional purposes.
Opportunities	 A general CE program is going to be built soon for pharmacists (needs time to expand and encompass aspects of AMR) In the veterinary/agriculture fields, PAAF closely communicates with FAO/OIE and concerned professionals attend few educational activities/workshops organized by FAO/OIE.

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	 Regarding the Country participation in international or regional AMR awareness campaigns, the IPC directorate regularly participate with the WHO regarding human health. Otherwise in non-human health sector, the country participates in some meetings organized by FAO or OIE however; this is not structured or systematic. Kuwait is part of the GCC AMR group and participates in all its activities. Now AMR is recognized a national priority where there will be interministerial collaboration and commitment through this national action plan to combat AMR. Relation between PAAF, MOH, FAO, OIE, WHO and KEPA is an opportunity to enhance AMR awareness in different fields based on the One Health approach, in addition to being part of GCC.
Threats	None

AMR Surveillance

Strengths	 Surveillance of AMR in isolates from humans has already started in one hospital through reporting its AMR data to GLASS platform. Now reporting to WHO GLASS will extend to 8 hospitals from the country. As for the incidence and prevalence of AMR in humans, some hospitals report their own AMR data on an individual basis. On the hospital level, a strength is the presence of national healthcare-associated infections surveillance system with the capability of detection of MDROs isolated from HAIs. Surveillance of infectious diseases such as brucellosis, TB, etc. is present. There is an existing tailored program for routine monitoring of the environment, as well as a chemical and biological focused marine monitoring program for environmental pollutants. Monitoring of antibiotic residues is done for milk, meat, poultry, eggs and vegetables A strength is the availability of a well-equipped microbiology laboratories in MOH and in other healthcare facilities, as well as new technologies to detect and identify AMR. Some of the functions of a national reference laboratory exist but in different places and not in one single place. For example, external Quality Control (QC) schemes are done by the central entity of medical labs services at MOH, and this central entity plays role in distributing external QC samples and getting the results. The Lab council requested from MOH for accrediting the labs in the country. This council takes into consideration the presence of a qualified microbiologist in the lab as well as the success in QC schemes, in addition to abiding by CLSI guidelines. Despite that there is no training for lab personnel regarding AMR surveillance, but all personnel should be qualified (education/degrees). All labs in the country unanimously follow CLSI guidelines in the country, and thus there is no need to put new lab practice guidelines. Sporadic research on AMR exists

	 National authorities work with close communication with OIE, FAO and WHO, in addition to being part of GCC. 						
Weaknesses	 National AMR surveillance and antimicrobial use monitoring systems are not available. As for the incidence and prevalence of AMR in humans, there are no national reports on this issue. Regarding surveillance of AMR in isolates from animals, plants, food and environment, it is unavailable, thus the incidence and prevalence of AMR in these fields is unavailable too. Monitoring of use of antimicrobial agents in humans is almost not available except for a national study that is starting and did not expand yet. Monitoring of use of antimicrobial agents in animals, plants and environment does not exist except for monitoring antibiotic residue in food products like milk, meat, poultry, eggs and vegetables. In addition, data on the extent and impact of AMR such as human morbidity, mortality and other health outcomes in relation to AMR, as well as data on economic impact of AMR in humans, animals, plants, food, and the environment are not available. A national mechanism coordinates the different national AMR surveillance and antimicrobial use monitoring systems is not available. A National Reference Lab for antimicrobial resistance for human and animal sectors is not available. A national research agenda for AMR surveillance is not available. 						
Opportunities	 An opportunity is the enrolment of Kuwaiti MOH in GLASS- AMC and utilization of this platform. It is economically feasible to increase manpower to perform AMR surveillance activities and report data in all sectors. There is one laboratory that is assigned for reporting AMR data to GLASS platform and this can be extended for national AMR surveillance. There is a proposal for the accreditation of 2 laboratories in the country by The College of American Pathologists (CAP) and if this pilot project succeeds, this will extend to other labs in the country. There is an opportunity to conduct AMR-related research in these fields: a decree regarding environment protection in Kuwait exists to support the recommendation from KEPA and PAAF for the need to conduct AMR research in these fields. The presence of this decree facilitates submitting research proposals and projects and getting funded. Another opportunity is the ppresence of research centres that are interested in performing studies on AMR and expanding it throughout the country (KEPA, KISR and Kuwait University) Relation between PAAF, MOH, FAO, OIE, WHO is an opportunity to enhance AMR surveillance in different fields based on the One Health approach 						
Threats	• None						

IPC

Human Health

- On the national level in Kuwait, there is a national central IPC directorate/national office in the MOH, established since 1980. This national office communicates with all IPC teams in public and private healthcare facilities across the country. This directorate has national IPC guidelines that are being regularly updated, plus checklists and audit tools.
- On the hospital level, each hospital has an IPC office that consists of at least 1 Medical Doctor (MD) and around 5 to 10 nurses depending on the hospital size/number of hospital beds, and this office is in close communication with the central IPC directorate.
 IPC policies and procedures are included in hospital accreditation standards and are being regularly audited by accreditation bodies.
- Regarding IPC professionals in healthcare facilities, they have
 usually an MD degree and an advanced degree in public health or
 microbiology. Their IPC/AMR training is mandatory for a period of 2
 months before joining the job. It is condensed and well-structured
 prepared by the MOH central IPC directorate. Training for the HAI
 surveillance is done at the IPC directorate while the IPC training is
 done in MOH hospitals. Regular in-service training is done for the
 IPC professionals in the Central IPC directorate. Additionally,
 targeted training is performed when there is any updates or new
 policies/guidelines.
- Upon hiring in hospitals, nurses who will work at IPC offices receive mandatory training for 3 months in the form of in-hospital training by IPC professionals as per specific material prepared and supplied by the central IPC directorate. Recently, they receive HAI surveillance training in the IPC directorate. IPC nurses receive regular refreshing training while targeted training is performed when there is any updates or new policies/ guidelines.
- Regarding general HCWs training, there is a training program in each hospital for HCWs and the IPC team in the hospital is responsible for it. All newly employed HCWs receive an IPC training and nurses should receive an IPC training on a yearly basis, but this issue is not very well structured or followed, and this should be worked on and improved.
- Healthcare-associated infections (HAI): National HAI surveillance is being implemented in all MOH hospitals. It is prospective patientbased facility wide includes all types of infections in all inpatient locations with calculation of risk adjusted rates. Dialysis surveillance for chronic outpatient hemodialysis patients is also done. Surveillance is based on CDC/NHSN methodology. For MDROs,

Strengths

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- incidence density/1000 patient days and % of the total isolates of 8 groups of MRDOs are calculated. Reports and feedback are disseminated and discussed with the relevant stockholders. National data base and benchmarks of HAI are prepared in the IPC directorate.
- Healthcare waste management: There is a structured management program of the hazardous waste generated from healthcare systems (solid and liquid waste). The MOH has guidelines for both types of wastes. There are incinerators with KEPA's authority to oversee the functioning of these incinerators. (Responsible entities: Hotel service Directorate, Health Licence Directorate, special committee in KEPA)
- Waste water management system: There is a structured management program under the authority of the Ministry of public works and being monitored from KEPA & Ministry of Public Works.
- On the healthcare facility level, there is a presence of safe sufficient drinking water and adequate sanitation and hygiene, where there is regular testing of the quality of water used in healthcare.
- Food handling in healthcare facilities and community:
 - regular testing of food handlers by MOH,
 - o presence of health cards;
 - guidelines for food handling in healthcare facilities depending on department plus general guidelines for cooking, storage.... etc.
- Vaccination programs:
 - availability of up-to-date guidelines and vaccines for the pediatric and adult population
 - o robust vaccination coverage for the pediatric population
 - adult vaccination guidelines stratified by age, occupation, comorbidities especially respiratory diseases, seasonal events like Pilgrimage.

Non-human health sectors (food production/animal husbandry/ veterinary/ agriculture/ environmental fields)

- <u>Animal Husbandry/Agriculture/Food Production:</u>
 - Biosecurity measures are present in the farms under the responsibility of PAAF
 - There are guidelines for vaccinating, isolating and treating animals against infectious diseases (based on OIE guidelines) monitored and audited by PAAF.
 - All animals should be registered and there is a veterinary laboratory at PAAF.
 - Infrastructure related to the safely securing/auditing/quality control of products in the food chain is present, whether

being the food produced in the country or imported. The responsible entities are the PAFN and PAAF. Food is examined chemically and microbiologically according to GSO standards in public health labs in MOH. **Environment:** o Disposal of diseases animals and waste is well organized and controlled and closely monitored by PAAF. **Human Health** Regarding IPC education in the country in general, there are no opportunities of learning about IPC outside the formal teaching provided by the central IPC directorate. There no courses given at universities or masters degrees or structured continuous education programs. Regarding HAI surveillance, HAI data is collected manually intrahospital and reported to MOH IPC directorate manually and this represents a gap. Responsible authorities believe that it is better that this work becomes computerized. This issue should be imposed on hospitals to shift from manual to electronic work. Yet, even in hospitals that already have electronic medical records, IPC data is not linked to these records and there is still a lot of manual work to link between the two. Healthcare waste management: disposal is not optimized similar to the public sector; it needs monitoring and better implementation. Long term care facilities (LTCF): There isn't a proper structure or specialized manpower and no specific IPC guidelines for LTCF. Weaknesses Vaccination Programs: vaccination uptake in adults is weak and needs to be improved, thus raise adult awareness regarding the importance of vaccination for the prevention of many vaccinepreventable diseases. Non-human health sectors <u>Animal Husbandry/Agriculture/Food Production:</u> o Absence of National guidelines for infection control biosecurity measures in animal husbandry. There are no research projects directly related to IPC/AMR in animal husbandry or agriculture **Veterinary Education and Training:** • There are no veterinary schools or education in Kuwait • Veterinarians receive their degrees and training outside the country. Absence of centrally oriented courses or education for veterinary professionals regarding IPC/AMR. Research: A gap is the absence of IPC/AMR research in these fields. **Opportunities** Human Health

- In healthcare facilities, IPC bundles of care are already available and implemented but need updating by including new evidence-based recommendations from the medical literature.
- Healthcare-associated infections: this data should become automatically retrievable from the computerized system to be implemented in healthcare facilities similar to other information in medical records, not to mention to link all relevant lab data to IPC data. And from another side, this IPC data should be digitally linked and available in real time on the network of central IPC directorate.

Long term care facilities (LTCF):

- Under the responsibility of Public Authority of Disability Affairs which is a separate authority from MOH and which is recently supervised by the Minister of Health.
- There are two large centers and several satellite centers linked together.
- The team responsible for supervision of the IPC program in LTCF centers constitutes of a part time doctor and four full time nursing staff with basic knowledge of IPC.

Non-human health sectors:

- Animal Husbandry/Agriculture/Food Production:
 - Surveillance of infectious diseases such as brucellosis, TB, etc. is present.
 - There are veterinary testing laboratories at PAAF.

Veterinary Education and Training:

- In order to be licensed to work in Kuwait, professionals who are hired from outside the country should undergo an interview and licensing exam. This procedure should be repeated every 2 years to renew the license. The exam and interview do not include questions related to IPC/AMR.
- The continuous education on IPC and AMR is only provided by drug companies.

• Environment:

 Presence of sewage effluent monitoring guidelines but not targeting the spread of AMR. This represents a potential or opportunity to perform studies aiming at monitoring the presence of resistant organisms in sewage effluent.

Research:

 There is an opportunity to conduct AMR-related research in these fields: a decree regarding environment protection in Kuwait exists to support the recommendation from KEPA and PAAF for the need to conduct IPC/AMR research in these fields. The presence of this decree facilitates submitting research proposals and projects and getting funded.

	 Relation between PAAF, MOH, FAO, OIE, WHO is an opportunity to enhance IPC in different fields based on the
	One Health approach, in addition to being part of GCC.
Threats	None

Rational Use of Antimicrobials

<u>Rational Ose of </u>	7 THE PROPERTY OF THE PROPERTY
Strengths	 There is a national human/animal/plant drug regulatory authority (Pharmaceutical and Herbal Medicines Registration and Control Administration in MOH) that works using international standards and guidelines, not to mention that they have their own protocols for drug marketing authorization to ensure that antimicrobial agents are quality assured, safe and effective. Pharmaceutical and Herbal Medicines Registration and Control Administration in MOH is responsible for quality management system for the antimicrobial agents supply chain as well as the preservation of antimicrobial agents. Pharmaceutical and Herbal Medicines Registration and Control Administration and The Dug Inspection Control in MOH are responsible for detecting and combating counterfeit antimicrobial agents. Pharmaceutical and Herbal Medicines Registration and Control Administration in MOH is responsible for regulating and controlling drug promotional practices (humans/animals). This entity does not allow drug promotion outside what is written in the brochure. But in the veterinary field, this issue is not being audited. As for the reimbursement lists for human health, this is under the authority of Pharmaceutical and Herbal Medicines Registration and Control Administration and no reimbursement occurs outside the protocols of this entity. There is a legislation that prohibit selling of antimicrobials without a prescription of a certified health professional. Veterinarians prescribe mass antimicrobial prophylactic therapy for herds or for treatment purposes, in addition to mass vaccination as prophylaxis when necessary according to guidelines of OIE Antibiotic residue is detected in milk, meat, poultry, eggs and vegetables. There is a washout period during which the animal does not receive any antimicrobials (prior to the milking process or slaughtering). Post-prescription review for appropriateness of antimicrobial agents 48–72 h aft

Weaknesses	 Pharmaceutical and Herbal Medicines Registration and Control Administration in MOH) does not systematically audit drug promotional practices in the animal field. Identifying/addressing economic incentives that encourage-inappropriate use of antimicrobial agents does not exist, as well as that optimize the use of antimicrobial agents in all sectors do not exist. Regarding the purchasing and prescribing of antimicrobial agents, a national essential medicine list is not available. On the institutional level, some healthcare facilities have their own essential medicine lists and others don't. There are standard treatment guidelines for use of antimicrobial agents in humans, but they need to be expanded and updated. These guidelines are present in public and private hospitals. These guidelines and their application are not properly audited. Standard treatment guidelines for use of antimicrobial agents in animals do not exist. Veterinary practitioners follow OIE guidelines and practice from what they have learned during their basic education. Responsible governmental authorities don't teach these guidelines and audit its proper practice or not; guidelines workshops do not exist. There is no follow-up on updated versions of these guidelines in animal health. There are no legislations to prevent the use of antibiotics used in humans and animals' therapy as growth promotors in animals and for crop protection There is no list or policies for the use of critically important antimicrobials. These legislations need to be activated. Policies on reduction in nontherapeutic use of antimicrobial agents in animal health are not available. Well-established Antimicrobial stewardship programs are not available in healthcare facilities (primary, secondary and tertiary care, private and public). However, Some activities are available in some healthcare settings. Qualified human resources are very scarce; the available
	A national research agenda for antimicrobial use and stewardship is not available.
	available.
Opportunities	 Authorities are working on formulating an essential medicine list. Relation between PAAF, MOH, FAO, OIE, WHO is an opportunity to enhance antimicrobial stewardship in different fields based on the One Health approach
Threats	• None

Country response

Governance

A National Committee for the proper use of antimicrobials was formulated by a ministerial decree signed by His Excellency the Minister of Health. This committee has members representing different specialties and sectors all over the country and headed by the Assistant Undersecretary of technical affairs and coordinated by the Director of Infection Control Directorate with the following as members:

- 1. WHO National Focal point of AMR-MOH
- 2. Director of Public Health Directorate -MOH
- 3. Director of Pharmaceutical Services Directorate -MOH
- 4. Director of Central Medical Stores Directorate-MOH
- 5. Director of Technical Directorate-MOH
- 6. Director of Primary Healthcare Directorate-MOH
- 7. Director of Medical Licensing Department-MOH
- 8. Director of Health Promotion Department-MOH
- 9. Director of Drug Inspection Control-MOH
- 10. Director of Pharmaceutical and Herbal Medicines Registration and Control Administration-MOH
- 11. Head of Public Health Laboratories-MOH
- 12. Head of Microbiology and Virology Committee- Council of Medical Laboratories-MOH
- 13. Director of Research and Studies Office- Kuwait Environment Public authority (KEPA)
- 14. Representative from The Public authority for Food and nutrition (PAFN)
- 15. Representative of Animal Health Department-The Public Authority of Agriculture Affairs and Fish Resources (PAAF)
- 16. Representative of Agriculture Sector Public Authority of Agriculture Affairs and Fish Resources (PAAF)

The purpose of this committee is to oversee and, when necessary, to coordinate AMR-related activities in all sectors

Governance Operational and Budget Plan

Activity	Sub- activity	Sub-sub- activity	Sub-sub- sub- activity	Unit	Quantity	Date	Location	Responsib le entity	Cost	Source Of funding	Indicator
Strategic Obje G1.1 Identify the needed AMR focal points in	ective G1. N	ational coord		een focal poi	nts in the dif	ferent sectors	s and differen	t pillars			
productio n, food safety and security, Plants and	G1.1.1 Define the TOR of focal points in each of these sectors			G1.1.1 TOR	G1.1.1 5	G1.1.1 Done	-	G1.1.1 NMCG	G1.1.1 None	G1.1.1 None	G1.1.1 TOR are put

authoritie s along with a yearly meeting of the different focal persons.											
	G1.1.2 Employ and assign the functions of a focal person for each pillar and for AMR in general	G1.1.2.1 Assign AMR focal points for each pillar in human health	G1.1.2.1.1 one focal point for each of the following pillars: IPC, Surveillan ce, ABX, Awarenes s	G1.1.2.1 Focal persons assigned	G1.1.2.1 4	G1.1.2.1 3 months	G1.1.2.1 MOH	G1.1.2.1 NMCG	G1.1.2.1 Assigned from existing ministry employee s	G1.1.2.1 None	G1.1.2 Focal persons for each pillar at MOH are assigned. Focal points in PAAF, and KEPA, and food safety are assigned
		G1.1.2.2 Assign the general focal person in MOH	G1.1.2.2.1 Dr. Abeer Aly pending assignmen t/ New employee	G1.1.2.2 Focal person assigned	G1.1.2.2 1	G1.1.2.2 3 months	G1.1.2.2 MOH	G1.1.2.2 NMCG	G1.1.2.2 Assigned from existing ministry/e mployees	G1.1.2.2 None	
		G1.1.2.3 Assign 1 focal	G1.1.2.3.1	G1.1.2.3 Letter from	G1.1.2.3 1	G1.1.2.3 3 months	G1.1.2.3 PAAF	G1.1.2.3 NMCG	G1.1.2.3 Assigned from	G1.1.2.3 None	

person in animal health ative in NMCG is the agricultur e fields for all pillars including IPC, Surveillan ce, ABX, Awarenes s	NMCG through MOH Undersecr etary General					existing authority employee s		
G1.1.2.4 Assign 1 focal person in KEPA for all pillars including IPC, Surveillan ce, ABX, Awarenes s		G1.1.2.4 1	G1.1.2.4 3 months	G1.1.2.4 KEPA	G1.1.2.4 NMCG	G1.1.2.4 Assigned from existing authority employee s	G1.1.2.4 None	
G1.1.2.5 Assign 1 Focal person in food safety IPC: Surveillan ce: G1.1.2.5.1 leave the decision to ministries /authoriti es	G1.1.2.5 Focal person assigned	G1.1.2.5 1	G1.1.2.5 3 months	G1.1.2.5 Food safety	G1.1.2.5 NMCG	G1.1.2.5 Assigned from existing ministry /authority employee s	G1.1.2.5 None	

		ABX Awarenes s									
	G1.1.2 Include inter- ministerial communic ation and coordinati on with the general AMR focal person in the TOR of the focal points			G1.1.2 TOR	G1.1.2 5	G1.1.2 3 months	G1.1.2 MOH	G1.1.2 NMCG	G1.1.2 None	G1.1.2 None	G1.1.2 None
Strategic O	G2.1.1	rovide legal a	and ministeria	al support for	the realization	on of the acti	vities listed ir	the plan			
G2.1 Form inter- ministerial NMCG committe e	Nominate the members and have a ministerial decree about their appointm ent			G2.1.1 One Health Committe e	G2.1.1 1	G2.1.1 Already available					G2.1.1 Already available
	G2.1.2 Write the TOR of the NMCG			G2.1.2 TOR	G2.1.2 1	G2.1.2 Already available					G2.1.2 Already available

	that include: to facilitate and oversee implemen tation, monitorin g and evaluation of the AMR action plan G2.1.3 The NMCG has dedicated funds for administr ative	G2.1.3.1 NMCG puts a budget for administr ative costs		G2.1.3.1 Budget	G2.1.3.1 1	G2.1.3.1 3 months	G2.1.3.1 MOH	G2.1.3.1 NMCG	G2.1.3.1 To be assigned later	G2.1.3.1 MOH	G2.1.3.1 A budget is put and provided
	costs.										
	bjective G3. T	he plan is bas	sed on evider	ce-based me	dicine and ρι	it by professi	onals				
G3.1 Technical groups for specific activities are created when needed from all sectors	G3.1.1 Mapping of profession als throughou t the country from public and private			G3.1.1 Mapping of profession als	G3.1.1 1	G3.1.1 3 months	G3.1.1 MOH/PAA F/KEPA/P AFN	G3.1.1 NMCG/ MOH Undersecr etary general /Director General in PAAF/KEP A/PAFN	G3.1.1 None	G3.1.1 None	G3.1.1 Technical groups are formed for each pillar, including represent ative of One

Strategic Ok	sectors that could be part of technical groups when needed.	rovide guida	nce and tools	to form a bas	sis for prepar	ation of a na	tional action	plan on AMR			Health approach
G4.1 Formulate or endorse guidelines for the different topics, along with tools	G4.1.1 Each technical group for each pillar will agree on existing guidelines or producing new guidelines or endorsing internatio nal guidelines and make them national			G4.1.1 Guidelines as per each pillar need	G4.1.1 4	G4.1.1 1 year	G4.1.1 MOH/PAA F/KEPA/P AFN	G4.1.1 Technical groups/ Focal persons of each pillar	G4.1.1 None (Professio nals are from the ministries /authoriti es and will perform these consultan cies as part of their jobs)	G4.1.1 None	G4.1.1 Guideline s are available when needed

N.B. Refer to Annex 1 for Governance Axis Strategic and Monitory plans

AMR Awareness Operational and Budget Plan

Global action plan strategic objective 1: Improve awareness and understanding of antimicrobial resistance through effective communication, education and training.

Activity	Sub- activity	Sub-sub- activity	Sub-sub- sub- activity	Unit	Quantity	Date	Location	Responsib le entity	Cost	Source Of funding	Indicator
	bjective 1.1 O	rganize the a	ctivities of th	is axis and en	sure a prope	r follow up			1		
1.1.1 To designate a specific employee or post that is responsibl e of realizing the plan, do the proper communic ations and follow up on the progress of the plan	1.1.1 Assign one employee /post to overlook these activities in MOH, KEPA, PAAF, food safety			1.1.1 Employee or position	1.1.1	1.1.1 3 months	1.1.1 MOH	1.1.1 MOH, KEPA, PAAF, PAFN	1.1.1 New Employee Or assign the function to existing employee	1.1.1 MOH	1.1.1 Awarenes s Focal person assigned
	1.1.2 Put TOR of employm ent			1.1.2 -TOR for MOH focal person -PAAF focal person	1.1.2	1.1.2 2 months	1.1.2 MOH	1.1.2 NMCG	1.1.2 None	1.1.2 None	1.1.2 TOR is put for all 4 focal persons.

				-KEPA Focal person -PAFN Focal person							
	1.1.3 Letter of employm ent or assignmen t of Responsib ilities of focal persons in the 4 ministries /authoriti es			1.1.3 Letter	1.1.3 1 per ministry/a uthority	1.1.3 3 months	1.1.3 MOH PAAF KEPA PAFN	1.1.3 MOH, KEPA, PAAF, PAFN	1.1.3 None	1.1.3 None	1.1.3 Letter of employm ent or assignme nt of duties to specific focal persons in the 4 ministries /authoriti es is done
1.2.1 Establish an evidence- based public communic ations program targeting the general public in the	1.2.1.1 Estimate the knowledg e, attitude and practice of general public regarding AMR in humans, animals, agricultur e and	crease nation	nal public awa	1.2.1.1 Already done on a random sample. in outpatient clinics on the use of antibiotics in humans	in as per One	1.2.1.1 Done	oacii uirough	Communicat	ion program		1.2.1.1 Done

communit y	environm ent										
	1.2.1.2 Conduct national public awareness campaigns based on the One health approach	1.2.1.2.1 Preparatio n of material regarding the awareness campaign based on the One health approach	1.2.1.2.1.1 Assign a team to formulate and design the public communic ation material regarding AMR	1.2.1.2.1.1 Already done		1.2.1.2.1.1 Done					1.2.1.2.1. 1 Done
			1.2.1.2.1.2 Produce the public communic ation material regarding AMR	1.2.1.2.1.1 Already available for AMR in humans		1.2.1.2.1.1 Done					1.2.1.2.1. 1 Done
			1.2.1.2.1.3 Update the communic ation material to cover all aspects of the One	1.2.1.2.1.3 Updated communic ation material	1.2.1.2.1.3	1.2.1.2.1.3 6 months	1.2.1.2.1.3 MOH/ PAAF/ KEPA/ PAFN	1.2.1.2.1.3 The awarenes s technical task force from MOH/PAA F/KEPA/P AFN	1.2.1.2.1.3 To be determine d by the responsibl e authoritie s	1.2.1.2.1.3 MOH/ PAAF/ KEPA/ PAFN	1.2.1.2.1. 3 The communic ation material covers all fields of One Health approach (Yes/No)

	health approach: Animal Health, Agricultur e, food safety and environm ent								
Put a yearly plan on broadcasti ng and posting the public communic ation material	1.2.1.2.2.1 Broadcasti ng is all year long, not only during the AMR awareness week (specific timing, place)	1.2.1.2.2.1 Broadcasti ng schedule (TV, Bill boards, advertise ment in malls,)	1.2.1.2.2.1	1.2.1.2.2.1 6 months	1.2.1.2.2.1 MOH Media Office	1.2.1.2.2.1 MOH/ Media Office	1.2.1.2.2.1 To be determine d by the responsibl e authoritie s	1.2.1.2.2.1 MOH/ PAAF/ KEPA/ PAFN	1.2.1.2.2. 1 Schedule is put (Yes/No)
	1.2.1.2.2.2 Prepare costing of the plan	1.2.1.2.2.2 Detailed costing of the plan	1.2.1.2.2.2	1.2.1.2.2.2 8 months	1.2.1.2.2.2 MOH/MO Informati on	1.2.1.2.2.2 MOH/ Media Office /MO Informati on	1.2.1.2.2.2 None	1.2.1.2.2.2 None	1.2.1.2.2. 2 Costing plan is finalized
1.2.1.2.3 Launch the broadcasti ng plan		1.2.1.2.3 Advertise ments launched according to a yearly schedule	1.2.1.2.3 1 plan/year	1.2.1.2.3 1 year	1.2.1.2.3 MOH/MO Informati on	MOH/ Media Office /MO Informati on	1.2.1.2.3 The budget that was put while costing all the	1.2.1.2.3 MOH/PAA F/KEPA/P AFN	1.2.1.2.3 Proportio n of the plan that was put on schedule

1.2.1.2.4 Engage different types of media including social media in AMR awareness such as putting material/a ds periodicall y on ministry/g overnmen tal websites, social media accounts like Facebook, Twitter, Instagram , etc.	1.2.1.2.4 Media message	1.2.1.2.4	1.2.1.2.4 1 year	1.2.1.2.4 MOH/MO Informati on	MOH/ Media Office /MO Informati on	advertise ments 1.2.1.2.4 To be determine d by the responsibl e authoritie s	1.2.1.2.4 MOH	1.2.1.2.4 Types of media or social media that mention AMR/mon th
1.2.1.2.5 Involve a public figure like Dr. Abdullah	1.2.1.2.5 Involvement of a public figure (Dr.	1.2.1.2.5	1.2.1.2.5 3 months	1.2.1.2.5 MOH/ TV/ Social Media Networks/	1.2.1.2.5 MOH/Me dia Office/Aw areness	1.2.1.2.5 None	1.2.1.2.5 None	1.2.1.2.5 Public figure (Dr. Abdullah Al-Sanad) sending

Al-Sanad (spokespe rson MOH) in delivering AMR messages, like tweets on twitter or messages in Talk Shows or through TV ads/News bar		Abdullah Al-Sanad)			Platforms	focal person			message regarding AMR
1.2.1.2.6 Discuss AMR in TV talk shows	1.2.1.2.6.1 MOH sends a letter to Ministry of Informati on/asking to host profession als in AMR to discuss this issue on TV and thus raise public awareness	1.2.1.2.6.1 Letter	1.2.1.2.6.1	1.2.1.2.6.1 2 months	1.2.1.2.6.1 MOH	1.2.1.2.6.1 Awarenes s focal person/ MOH undersecr etary general	1.2.1.2.6.1 None	1.2.1.2.6.1 None	1.2.1.2.6. 1 Proportio n of talk shows episodes that discuss AMR
1.2.1.2.7	1.2.1.2.7.1	1.2.1.2.7.1	1.2.1.2.7.1	1.2.1.2.7.1 2 months	1.2.1.2.7.1 PAAF	1.2.1.2.7.1 PAAF	1.2.1.2.7.1 None	1.2.1.2.7.1 None	1.2.1.2.7. 1

Strategic Ol	piective 1.3 In	Involve Non- Governme ntal Organizati ons (NGOs) in AMR awareness campaign	Contact FAO and OIE and ask about their communic ations program regarding AMR and communic ate with them the NAP on AMR ducation as p	Letter to FAO and OIE/Kuwai t offices	n approach a	t an early age	KEPA PAFN	KEPA PAFN			None
1.3.1 Include basic AMR education in school curricula	1.3.1.1 MOH sends a letter with key facts about AMR, antibiotics , hygiene and IPC basics through the School Health Administr ation asking to include material	iciude Aivik e	uucation as p	1.3.1.1 Letter	1.3.1.1 1	1.3.1.1 2 months	1.3.1.1 MOH	1.3.1.1 MOH/Sch ool Health Administr ation	1.3.1.1 None	1.3.1.1 None	1.3.1.1 Letter sent Yes or No

related to these topics in school curricula									
1.3.1.2 Prepare a module/A dd material related to basic AMR knowledg e, antibiotics , hygiene and IPC in the curricula	1.3.1.2.1 Create a committe e of comprisin g of AMR profession al and education profession als to create a module about basic AMR knowledg e, antibiotics , hygiene and IPC to be included in school curricula	1.3.1.2.1 Committ e	112171	1.3.1.2.1 2 months	1.3.1.2.1 MOH/Min istry of Education	1.3.1.2.1 MOH/ Awarenes s Focal person/ awarenes s technical task force	1.3.1.2.1 Rewardin g of the committe e members	1.3.1.2.1 MOH/ MOE	1.3.1.2.1 Module prepared Yes/No
	1.3.1.2.2 Include this module in secondary	1.3.1.2.2 Letter from MOH	1.3.1.2.2	1.3.1.2.2 8 months	1.3.1.2.2 MOH/ Ministry of Education	1.3.1.2.2 MOH/ School Health Administr ation/	1.3.1.2.2 None	1.3.1.2.2 None	1.3.1.2.2 Percent of schools that teach the AMR module

	school education			Ministry of Education	in their secondary school curricula
Strategic Objective 1	1.3.1.2.3 Yearly AMR module workshop s involving teachers	1.3.1.2.3 1.3.1.2.3 Workshop 1/year	1.3.1.2.3 MOH/ Ministry of Education	Directorat e/ Ministry of Education	1.3.1.2.3 Proportio n of teachers
Strategic Objective 1 1.4.1 Establish an evidence- based communic ation program targeting the healthcare providers for human health, animal health,	survey to estimate this knowledg e among healthcar e workers and identify	1.4.1.1.1 Already done 2011	1.4.1.1.1 Done	pproach	1.4.1.1.1 Done

e and environm ent sector										
	1.4.1.2 Conduct awareness program within hospitals, primary Healthcar e, among physicians , nurses, pharmacis ts, dentists, physiothe rapists, and other HC providers	1.4.1.2.1 MOH mandates to all hospitals and primary Health care centers to provide mandator y sessions about AMR to all employee s who are subject to contract/li cense renewal	1.4.1.2.1 Letter from MOH to hospitals and to primary Health care centers	1.4.1.2.1	1.4.1.2.1 2 months	1.4.1.2.1 MOH	1.4.1.2.1 MOH/Awa reness Focal person	1.4.1.2.1 None	1.4.1.2.1 None	1.4.1.2.1 Proportio n of hospitals and primary Health care centers that provide and request yearly attendanc e of AMR conferenc es/session s
	1.4.1.3 AMR mandator y courses for pharmacis ts in private practice who are subject to	1.4.1.3.1 Letter from MOH to Drug Inspection Control to request evidence of AMR course attendanc	1.4.1.3.1 Letter	1.4.1.3.1	1.4.1.3.1 6 months	1.4.1.3.1 MOH	1.4.1.3.1 MOH	1.4.1.3.1 None	1.4.1.3.1 None	1.4.1.3.1 Letter is issued

contract/ license renewal	e about AMR								
renewai	upon license renewal								
1.4.1.4 Link of AMR course attendanc e to hospital pharmacis ts' and primary Healthcar e pharmacis ts' promotio n.	1.4.1.4.1 MOH recomme nds yearly attendanc e for promotio n	1.4.1.4.1 Mandate	1.4.1.4.1	1.4.1.4.1 1 year	1.4.1.4.1 MOH	1.4.1.4.1 MOH Undersecr etary general/ AMR Focal person/	1.4.1.4.1 None	1.4.1.4.1 None	1.4.1.4.1 Mandate is issued
1.4.1.5 Preparatio n of a bank of presentati ons, films, articles, etc. diffused to these syndicates and hospitals to be used as	1.4.1.5.1 Toolkit for AMR communic ation program to be used in hospital and syndicates	1.4.1.5.1 Toolkit (Slide sets, films, articles, etc)	1.4.1.5.1 1	1.4.1.5.1 1 year	1.4.1.5.1 NMCG/A wareness Technical task force	1.4.1.5.1 NMCG/A wareness Technical task force	1.4.1.5.1 To be determine d by MOH	1.4.1.5.1 MOH	1.4.1.5.1 Toolkit available

	material in their AMR communic ation programs.										
	1.4.1.6 Syndicates and hospitals send yearly report to MOH about this activity			1.4.1.6 Report to MOH from hospitals	1.4.1.6 Once/ year/ hospital	1.4.1.6 Yearly starting 1 year from starting the plan	1.4.1.6 Hospitals	1.4.1.6 AMR Focal person	1.4.1.6 None	1.4.1.6 None	1.4.1.6 Proportio n of hospitals that send report about AMR Awarenes s activities to MOH
Strategic Ob approach	ojective 1.5 Ra	aise awarene	ss of veterina	rians, agricul	ture professi	onals and env	vironmental v	workers abou	t AMR based	on the One F	lealth
1.5.1 Conduct an awareness program among veterinari ans, agricultur e profession als, environm ental workers, in addition	1.5.1.1 Prepare AMR awareness toolkit including slide sets, short films, lectures, posters for these profession als			1.5.1.1 AMR toolkit	1.5.1.1 1	1.5.1.1 6 months	1.5.1.1 PAAF KEPA PAFN	1.5.1.1 Awarenes s Technical Task force	1.5.1.1 To be determine d by the responsibl e authoritie s	1.5.1.1 PAAF KEPA PAFN	1.5.1.1 AMR toolkit is available

to profession als in PAAF, KEPA, and food safety.										
	1.5.1.2 Integrate AMR awareness in continuou s education programs of veterinari ans, agricultur e profession als, environm ental workers, in addition to profession als in PAAF, KEPA, and food safety	1.5.1.2.1 The focal person in each ministry/a uthority checks education al activities in their facilities and Includes AMR awareness as a topic of continuous education	1.5.1.2.1 AMR education sessions in PAAF KEPA PAFN	1.5.1.2.1 2 per year in each specialty	1.5.1.2.1 1 year	1.5.1.2.1 PAAF KEPA PAFN	1.5.1.2.1 Focal person in each ministry/a uthority	1.5.1.2.1 None	1.5.1.2.1 None	1.5.1.2.1 Number of AMR Awarenes s sessions per year in each Ministry/a uthority
		1.5.1.2.2	1.5.1.2.2	1.5.1.2.2	1.5.1.2.2	1.5.1.2.2	1.5.1.2.2	1.5.1.2.2	1.5.1.2.2	1.5.1.2.2

		Mandate that all drug companie s' education al activities should include AMR awareness	Mandate	1	6 months	PAAF	- Awarenes s focal person in PAAF/Dire ctor General of PAAF	None	None	Proportio n of promotio nal activities by drug companie s in these fields that include messages about AMR
1.5.2 in regional/i nternation al AMR awareness campaign	L.5.2.1 Participat e actively n WHO regional AMR campaign and FAO and DIE activities related to	1.5.2.1.1 Register Kuwait authoritie s (KEPA, PAAF, MOH) to participat e in the yearly schedule of AMR activities in WHO, FAO, OIE And encourage individual healthcar e profession als,	1.5.2.1.1 Participati on in Internatio nal activities organized by WHO FAO OIE		1.5.2.1.1 1 year	1.5.2.1.1 MOH PAAF KEPA	1.5.2.1.1 Focal persons in MOH PAAF KEPA/ Awarenes s Technical Task force	1.5.2.1.1 None	1.5.2.1.1 None	1.5.2.1.1 Proportio n of events related to AMR in which the country participat es

		veterinari ans, agricultur e profession als, environm ental workers to participat e									
	1.5.2.2 Participat e in GCC activities related to AMR	1.5.2.2.1 Kuwait is part of GCC AMR Group		1.5.2.2.1 Done		1.5.2.2.1 Done					1.5.2.2.1 Done
		1.5.2.2.2 Kuwait participat es in all the AMR activities GCC		1.5.2.2.2 Done		1.5.2.2.2 Done					1.5.2.2.2 Done
_	-	clude AMR to		her educatio	n/post-gradu	ate studies in	all human h	ealth specialt	ies, agricultu	re, veterinary	food
1.6.1 Include AMR and related topics based on the One Health approach	1.6.1.1 Include AMR and related topics based on the One Health approach	1.6.1.1.1 preparatio n of modules for these specialties	aities.	1.6.1.1.1 Module for each specialty	1.6.1.1.1 1 per specialty	1.6.1.1.1 6 months	1.6.1.1.1 MOH KEPA PAFN	1.6.1.1.1 Awarenes s focal persons in all ministries /authoriti es /Awarene	1.6.1.1.1 None	1.6.1.1.1 None	1.6.1.1.1 Modules prepared yes/no

as a core compone nt of profession al education, training, certificati on	in undergrad uate curricula for human health (medical, nursing, dental and pharmace utical), food safety, and environm ent related specialties .						ss technical task force			
		1.6.1.1.2 Letter from MOH, PAFN KEPA to Ministry of Higher Education to impose on universitie s/authoriti es to include these	1.6.1.1.2 Letter	1.6.1.1.2 1	1.6.1.1.2 6 months	1.6.1.1.2 MOH KEPA PAFN	1.6.1.1.2 Awarenes s focal persons in all ministries /authoriti es / Awarenes s task force	1.6.1.1.2 None	1.6.1.1.2 None	1.6.1.1.2 Letter sent Yes/No

	modules in the correspon ding education programs.								
1.6.1.2 AMR and related topics based on the One Health approach included in the licensing procedure for nurses, dentists, physicians , pharmacis ts, veterinari ans and other profession als for which licensing is required.	1.6.1.2.1 Letter from MOH, PAAF, PAFN and KEPA to licensing bodies of these profession als to include AMR and its related topics in licensing exams and interviews or in the evaluation procedure for license renewal	1.6.1.2.1 Letter from, MOH, PAAF, PAFN and KEPA, to the licensing bodies of each of these profession s	1.6.1.2.1 1 per profession /per licensing body	1.6.1.2.1 3 months	1.6.1.2.1 MOH PAAF KEPA PAFN	1.6.1.2.1 MOH, PAAF, PAFN and KEPA /AMR focal persons in all authoritie s	1.6.1.2.1 None	1.6.1.2.1 None	1.6.1.2.1 Letters sent Yes/No

Strategic Objective Include 1.7 AMR and related topics in quality assurance and accreditation programs in human health, animal health, agriculture, food chain and environment

1.7.1 IPC, AMR surveillanc e and AMS are included in the accreditati on standards of healthcare facilities	1.7.1.1 Letter from MOH to accreditati on bodies to include AMR surveillan ce and implemen tation of antimicro bial stewardsh ip programs as part of accreditati on standards in all healthcar e facilities as well as in the licensing renewal procedure s of private healthcar e facilities			1.7.1.1 Letter	1.7.1.1	1.7.1.1 3 months	1.7.1.1 MOH	on1.7.1.1 MOH/AM R awarenes s focal person/ General AMR focal person	1.7.1.1 None	1.7.1.1 None	1.7.1.1 AMR surveillan ce and AMS programs are included in the accreditat ion and license renewal standards for healthcar e facilities
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	part of these standards		_								
1.7.2 IPC and rational use of antimicro bials is checked in the procedure of licensing farms, animal husbandry, agricultur e, aquatic and marine fields	1.7.2.1 Review of large animal/po ultry farms licensing policies and procedure s, to include IPC and proper ABX use and preventio n of ABX use as growth promoters	1.7.2.1.1 Mandate that ABX are not to be used as growth promoters		1.7.2.1.1 Mandate	1.7.2.1.1	1.7.2.1.1 3 months	1.7.2.1.1 PAAF	1.7.2.1.1 PAAF Director General /PAAF awarenes s focal person	1.7.2.1.1 None	1.7.2.1.1 None	1.7.2.1.1 Mandate issued
		1.7.2.1.2 Letter from PAAF Director General to licensing office to include these points in		1.7.2.1.2 Letter	1.7.2.1.2	1.7.2.1.2 3 months	1.7.2.1.2 PAAF	1.7.2.1.2 PAAF Director General	1.7.2.1.2 None	1.7.2.1.2 None	1.7.2.1.2 AMR- related topics are included in licensing conditions

Kuwait national action plan on antimicrobial resistance 2022

the					
standards					

N.B. Refer to Annex 2 for AMR Awareness Axis Strategic and Monitory plans

AMR Surveillance Operational and Budget Plan

Global action plan strategic objective 2: Strengthen the knowledge and evidence base through surveillance and research.

Strategic intervent ions	Activity	Sub- activity	Sub-sub- activity	Sub-sub- sub- activity	Unit	Quantity	Date	Location	Responsi ble entity	Cost	Source Of funding	Indicator
Strategic C		: Set up a n	ational surv	eillance syste	em for antin	nicrobial res	istance base	d on One He	ealth Approa	ch		1
2.1.1 Establish a national coordina tion structure for AMR surveilla nce in humans, veterinar y agricultu re and environ ment	2.1.1.1 Write and approve terms of referenc e for a national coordina ting committ ee for AMR surveilla nce with the mandate to oversee the AMR surveilla nce program, including AMR surveilla nce program, including AMR surveilla nce in Humans,				2.1.1 Committ ee that includes focal for human, vets, agricultu re, environ ment and Food safety	2.1.1	2.1.1 6 months	2.1.1 MOH	2.1.1 MOH	2.1.1 None	2.1.1 None	2.1.1 Committ ee is formed yes/No

Veterinar										
у,										
agricultu										
re,										
environm										
ent and										
Food										
safety.										
2.1.1.2.										
Write										
and										
approve										
terms of										
referenc										
e for a										
central										
office that										
executes	2.1.1.2.1									
the	Write		2.1.1.2.1							
National	terms of		TOR of a		2.1.1.2.1					2.1.1.2.1
surveilla	referenc		central	2.1.1.2.1	3	2.1.1.2.1	2.1.1.2.1	2.1.1.2.1	2.1.1.2.1	TOR
nce in	e of the		office at	1	months	MOH	MOH	None	None	written
humans	AMR		MOH		1110111113					Wilten
and	office at		IVIOIT							
receives	МОН									
data										
from										
other										
AMR										
offices in										
PAAF,										
KEPA and										
food										
safety										

2.1.1.2.2 Create the AMR coordina ting office at MOH.	2.1.1.2.2. 1 Include the already existing Glass team into the MOH National AMR office for Human sector at MOH. 2.1.1.2.2. 1.1 Add employ es with support from IT at MOH to be able to handle nation de data (to do the function of the central office).	2.1.1.2.2. 1.1 Addition al employe es	2.1.1.2.2. 1.1 2	2.1.1.2.2. 1.1 6 months	2.1.1.2.2. 1.1 MOH	2.1.1.2.2. 1.1 MOH/Inf ormation system administr ation	2.1.1.2.2. 1.1 2 extra employe es	2.1.1.2.2. 1.1 MOH	2.1.1.2.2. 1.1 2 extra employe es are employe d. Yes/No
	2.1.1.2.2. 2 Mandate that each MOH hospital has an AMR surveilla nce focal person that coordina tes with Central AMR surveilla	2.1.1.2.2. 2 Mandate	2.1.1.2.2. 2 1	2.1.1.2.2. 2 3 months	2.1.1.2.2. 2 MOH	2.1.1.2.2. 2 MOH	2.1.1.2.2. 2 None	2.1.1.2.2. 2 None	2.1.1.2.2. 2 Mandate issued Yes/No

		nce office.								
I F F k a a t f s r	2.1.1.2.3 Inform PAAF,PA FN and KEPA about the need for AMR surveilla nce in their respectiv e fields.		2.1.1.2.3 Letter	2.1.1.2.3	2.1.1.2.3 1 month	2.1.1.2.3 MOH	2.1.1.2.3 MOH	2.1.1.2.3 None	2.1.1.2.3 None	2.1.1.2.3 Letter sent
e f ii k	2.1.1.2.4 Nominat e AMR focal persons in PAAF, KEPA, and Food safety		2.1.1.2.4 Nominati on of focal persons	2.1.1.2.4	2.1.1.2.4 3 months	2.1.1.2.4 PAAF and KEPA and PAFN	2.1.1.2.4 PAAF/ KEPA/ PAFN	2.1.1.2.4 None	2.1.1.2.4 None	2.1.1.2.4 AMR surveilla nce focal persons appointe d in all 3 authoriti es.
j c c t f f	2.1.1.2.5 Write the job descripti on of these focal persons. The job descripti on would be to		2.1.1.2.5 TOR	2.1.1.2.5	2.1.1.2.5 3 months	2.1.1.2.5 PAAF and KEPA and PAFN	2.1.1.2.5 PAAF/ KEPA/ PAFN	2.1.1.2.5 None	2.1.1.2.5 None	2.1.1.2.5 TOR are set

	collect AMR data and supply it to the Central AMR office and full coordina tion with the Central AMR office at MOH.									
2.1.1.3 Identify priority organism s for the purpose of national AMR surveilla nce in human sector, animal sector, plants and Environm ent.			2.1.1.3 List of priority organism s for AMR surveilla nce in humans, animals, agricultu re and food safety	2.1.1.3 1 list	2.1.1.3 3 months	2.1.1.3 MOH	2.1.1.3 Central committ ee for AMR surveilla nce	2.1.1.3 None	2.1.1.3 None	2.1.1.3 List is available and includes all listed sectors.

2.1.1.4 Systemat ically collect, analyze and report data on AMR in humans in order to inform decision-making at national and internati onal levels (Choose sentinel group of labs)	2.1.1.4.1 Choose a list of labs (dealing with specimen from humans) that can provide reliable AMR data will form a sentinel for starting AMR data collectio n that will be submitte d to AMR central office.		2.1.1.4.1 List of labs that potential ly will submit AMR data Already exist for 2021 GLASS Data	2.1.1.4.1	2.1.1.4.1 3 months	2.1.1.4.1 MOH / GLASS team	2.1.1.4.1 MOH	2.1.1.4.1 None	2.1.1.4.1 None	2.1.1.4.1 List is chosen (8 Labs submitte d GLASS data for 2021 -Amiri -Adan - Mubarak Alkhabee r -Jaber - Farwania -Jahra -Chest -Ibn Sina)
	2.1.1.4.2 Increase the number of labs that are included in the list on yearly	2.1.1.4.2. 1 Establish close communi cation between central AMR	2.1.1.4.2. 1 New Labs in the list	2.1.1.4.2. 1 10 /year	2.1.1.4.2. 1 1 year	2.1.1.4.2. 1 MOH/ Central AMR Surveilla nce office	2.1.1.4.2. 1 MOH/ Central AMR Surveilla nce office	2.1.1.4.2. 1 None	2.1.1.4.2. 1 None	2.1.1.4.2. 1 Proportio n of labs in the country that are included in the list

	basis until it becomes represen tative of the national data.	OFFICE that also collects data for glass with the lab council to have a list of labs that are								
		are accredite d by the council, and to include them in the list of labs eligible to submit data to GLASS and National AMR surveilla								
2.1.1.5 Establish an antimicr obial resistance	all MOH laborator ies in	nce 2.1.1.5.1. 1 Training of hospital AMR Surveilla	2.1.1.5.1. 1 TOT Worksho ps	2.1.1.5.1. 1 1/year	2.1.1.5.1. 1 1 year	2.1.1.5.1. 1 MOH/ Central AMR Surveilla	2.1.1.5.1. 1 MOH/ Central AMR Surveilla	2.1.1.5.1. 1 30,000 KD/year	2.1.1.5.1. 1 MOH	2.1.1.5.1. 1 Yearly worksho p performe d

	surveilla nce network for human health	national referenc e Lab.	nce focal persons for data cleaning and data entry into the MOH surveilla nce system				nce office	nce office			Yes/No
		2.1.1.5.2 Generate yearly report about AMR in humans in general and mainly priority organism s in humans by the MOH central office.		2.1.1.5.2 Report	2.1.1.5.2 1/year	2.1.1.5.2 2 years	2.1.1.5.2 MOH/ Central AMR Surveilla nce office	2.1.1.5.2 MOH/ Central AMR Surveilla nce office	2.1.1.5.2 None	2.1.1.5.2 None	2.1.1.5.2 Yearly human AMR report is issued
2.1.2 Participa te in Global Antimicr obial	2.1.2.1 Nominat e a hospital ad a team for			2.1.2.1 Team ad hospital nominat ed		2.1.2.1 Already done					2.1.2.1 Already done

Resistanc e Surveilla nce System (GLASS- AMR)	GLASS reporting with TOR											
,	2.1.2.2 Give the GLASS team the authority to request and collect data from labs nationwi de				2.1.2.2 Letter from MOH	2.1.2.2 1 to each lab or hospital	2.1.2.2 Already done					2.1.2.2 Already done
		2.1.2.2.1 The team initiates data collectio n from a sentinel of labs			2.1.2.2.1 Report submitte d to glass	2.1.2.2.1 1/year	2.1.2.2.1 Started 2021					2.1.2.2.1 Started 2021
	2.1.2.3 Send yearly report to glass with increasin g	2.1.2.3.1 Central GLASS focal group (Part of AMR central	2.1.2.3.1 1 Yearly training of GLASS focal persons on GLASS	2.1.2.3.1. 1.1 Central worksho ps yearly	2.1.2.3.1. 1.1 Training worksho p	2.1.2.3.1. 1.1 1/year	2.1.2.3.1. 1.1 2 years	2.1.2.3.1. 1.1 MOH/ Central AMR Surveilla nce office	2.1.2.3.1. 1.1 MOH/ Central AMR Surveilla nce office	2.1.2.3.1. 1.1 20,000 KD	2.1.2.3.1. 1.1 MOH	2.1.2.3.1. 1.1 Worksho p done, yes/No have 10- 20 %

	number of contribut ing labs	office), chooses from the list of eligible accredite d labs submitte d by central entity of medical labs to AMR office, 2.1.2.3.2 Yearly lab report is sent to GLASS with increasin g number of contribut ing labs every year.	data collectio n, manage ment and entry		2.1.2.3.2 Report sent to GLASS	2.1.2.3.2 1/year	2.1.2.3.2 Started already	2.1.2.3.2 MOH/ Central AMR Surveilla nce office	2.1.2.3.2 MOH/ Central AMR Surveilla nce office	2.1.2.3.2 None	2.1.2.3.2 None	increase in number of participa ting labs in GLASS annually 2.1.2.3.2 Data is available in the yearly GLASS report
Strategic O activities in		: Build labo	ratory capac	ity to produ	ce high-qual	ity microbio	logical data	for patient r	managemen	t and suppor	rt surveilland	ce
2.2.1 Lab in Hospitals become	2.2.1.1 Pilot with 2 labs				2.2.1.1 Lab CAP affiliation	2.2.1.1	2.2.1.1 1 year	2.2.1.1 Labs	2.2.1.1 Medical Laborato	2.2.1.1 To be determin ed	2.2.1.1 MOH	2.2.1.1 Started already

CAP certified (College of American pathologi sts)								ries Council			
	2.2.1.2 Increase the number of labs that are CAP members	2.2.1.2.1 Lab council puts a plan for sequenti al recruitm ent of labs into the CAP		2.2.1.2.1 Plan	2.2.1.2.1	2.2.1.2.1 1 year	2.2.1.2.1 Lab council	2.2.1.2.1 MOH	2.2.1.2.1 None	2.2.1.2.1 None	2.2.1.2.1 List is put already
		2.2.1.2.2 Sequenti al recruitm ent of labs into CAP		2.2.1.2.2 Labs recruited	2.2.1.2.2 2 labs/year	2.2.1.2.2 5 years	2.2.1.2.2 Labs	2.2.1.2.2 Medical Laborato ries Council	2.2.1.2.2 To be determin ed	2.2.1.2.2 MOH	2.2.1.1 Proportio n of Labs that are CAP affiliated
Improve private labs capacity to be eligible for submitti ng data to GLASS	2.2.2.1 Reinforce the accredita tion activity of the lab council that audits labs	2.2.2.1.1 Mandate from MOH that each private lab should pass the accredita tion by		2.2.2.1.1 Mandate	2.2.2.1.1	2.2.2.1.1 12-24 months	2.2.2.1.1 MOH	2.2.2.1.1 MOH/Me dical lab. council/ Medical Licensing Dep.	2.2.2.1.1 None	2.2.2.1.1 None	2.2.2.1 Proportio n of labs that are functiona l and that have passed the accredita

and National AMR	accordin g to availabilit y of qualified microbiol ogist, results of External QC and abidance to guideline s	the Lab council										tion testing
		2.2.2.1.2 Accredita tion of the lab council includes the availabili ty of qualified microbiol ogist, results of External QC and abidance to guideline s		s	2.2.2.1.2 Update of accredita tion standard s of labs	2.2.2.1.2	2.2.2.1.2 6 months	2.2.2.1.2 Labs	2.2.2.1.2 Medical Lab council	2.2.2.1.2 None	2.2.2.1.2 None	Updated accredita tion standard s of Labs prepared
Strategic o	bjective 2.3	Collect data		Agriculture a			22111	22111	22111	22111	22111	22111
2.3.1	2.3.1.1	2.3.1.1.1	2.3.1.1.1. 1		2.3.1.1.1. 1	2.3.1.1.1. 1	2.3.1.1.1. 1	2.3.1.1.1. 1	2.3.1.1.1. 1	2.3.1.1.1. 1	2.3.1.1.1. 1	2.3.1.1.1. 1

Collect AMR data from Vets	Map available data On AMR in vet and agricultu re in the country.	Agricultu re and Environ ment focal persons collect available info about AMR in the country and issue a report	Map PAAF labs and the type of AMR data that is generate d.	Report about PAAF Labs	1	6 months	PAAF Vet labs	PAAF	None	None	Report is Submitte d to central AMR surveilla nce office
			2.3.1.1.1. 2 Collect available data about AMR in PAAF/PA FN	2.3.1.1.1. 2 Report from PAAF and food safety focal persons	2.3.1.1.1. 2	2.3.1.1.1. 2 6 months	2.3.1.1.1. 2 PAAF/PA FN	2.3.1.1.1. 2 PAAF/PA FN	2.3.1.1.1. 2 None	2.3.1.1.1. 2 None	2.3.1.1.1. 2 AMR Vets /food reports are submitte d to central AMR Surveilla nce office
	2.3.1.2 Improve vet lab capacity in a way they generate	2.3.1.2.1 Establish quality control of Vet Microbio logy labs	2.3.1.2.1. 1 Assign expert to establish a system for	2.3.1.2.1. 1 professio nal assigned	2.3.1.2.1. 1	2.3.1.2.1. 1 1 year	2.3.1.2.1. 1 PAAF	2.3.1.2.1. 1 PAAF	2.3.1.2.1. 1 Salary	2.3.1.2.1. 1 PAAF	2.3.1.1.1. 1 Professio nal is assigned

6 6 6	data about AMR vets and agricultu re		quality control and put a plan	/employ ed							
			2.3.1.2.1. 2 Execute the plan of quality control in Vet labs	2.3.1.2.1. 2 qc of vet labs	2.3.1.2.1. 2 1	2.3.1.2.1. 2 2 year	2.3.1.2.1. 2 vet labs	2.3.1.2.1. 2 PAAF	2.3.1.2.1. 2 to be determin ed by PAAF	2.3.1.2.1. 2 PAAF	2.3.1.2.1. 2 Proportio n of VET Labs that Participat e in external quality control activities
		2.3.1.2.2 Extend Vet Micro Labs to identify priority organism s antibiogr am	2.3.1.2.2. 1 Increase man power	2.3.1.2.2. 1 Employe es	2.3.1.2.2. 1 To be determin ed Per lab	2.3.1.2.2. 1 6 months	2.3.1.2.2. 1 Vet Labs	2.3.1.2.2. 1 PAAF	2.3.1.2.2. 1 to be determin ed by PAAF	2.3.1.2.2. 1 PAAF	2.3.1.2.2. 1 none
			2.3.1.2.2. 2 Road map and full project	2.3.1.2.2. 2 Project	2.3.1.2.2. 2 1	2.3.1.2.2. 2 1 year	2.3.1.2.2. 2 PAAF/Ve t labs	2.3.1.2.2. 2 PAAF	2.3.1.2.2. 2 none	2.3.1.2.2. 2 none	2.3.1.2.2. 2 The project is put with specified budget

									and ready for executio n
	2.3.1.2.2. 3 Worksho ps for training about antibiogr am	2.3.1.2.2. 3 Worksho p	2.3.1.2.2. 3 1/year	2.3.1.2.2. 3 1 year	2.3.1.2.2. 3 PAAF Central Lab, or assigned lab	2.3.1.2.2. 3 PAAF	2.3.1.2.2. 3 Worksho p cost	2.3.1.2.2. 3 PAAF	2.3.1.2.2. 3 Worksho p is performe d.
2.3.1.2.3 Impleme nt national surveilla nce for antibiogr am priority animal pathogen s, zoonotic and commen sal bacterial isolates.	2.3.1.2.3. 1 Choose a number of Vet labs that could submit AMR data to PAAF for AMR surveilla nce	2.3.1.2.3. 1 List of labs	2.3.1.2.3. 1 1 list/year	2.3.1.2.3. 1 1 year	2.3.1.2.3. 1 PAAF	2.3.1.2.3. 1 PAAF AMR surveilla nce focal Person	2.3.1.2.3. 1 None	2.3.1.2.3. 1 None	2.3.1.2.3. 1 List of labs is available yearly
	2.3.1.2.3. 2 Collect data from a Sentinel	2.3.1.2.3. 2 Yearly data collectio n	2.3.1.2.3. 2 1/year	2.3.1.2.3. 2 2 years	2.3.1.2.3. 2 PAAF	2.3.1.2.3. 2 PAAF/ AMR surveilla	2.3.1.2.3. 2 None	2.3.1.2.3. 2 None	2.3.1.2.3. 2 Data is yearly collected

			of vet hospitals or labs that submit data to the departm ent of surveilla nce. (They belong to the list).					nce focal person			
		2.3.1.2.4 The focal AMR entity at PAAF submits AMR data to the Central AMR surveilla nce lab.		2.3.1.2.4 Data Submitte d to Central AMR office at MOH	2.3.1.2.4 1/year	2.3.1.2.4 1 year	2.3.1.2.4 PAAF/ MOH	2.3.1.2.4 AMR Surveilla nce focal person at PAAF	2.3.1.2.4 None	2.3.1.2.4 None	2.3.1.2.4 Data submitte d to MOH Central Office.
2.3.2 Collect AMR data from Environ ment.	2.3.2.1 Map available data from KEPA	2.3.2.1 Check KEPA Labs and publicati ons about any AMR		2.3.2.1 Search	2.3.2.1	2.3.2.1 3 months	2.3.2.1 KEPA labs	2.3.2.1 KEPA focal person for AMR surveilla nce	2.3.2.1 None	2.3.2.1 None	2.3.2.1 Report is issued

2.3.2.2 Initiate AMR data collectio n from Environm ent By research projects.	data in environ ment 2.3.2.2.1 The director of the office of research and studies puts AMR surveilla nce as one of the priority subjects in	2.3.2.2.1. 1 The plan of prioritizi ng AMR surveilla nce in Environm ent and marine environm ent re and is being discussed with universiti	2.3.2.2.1. 1 Minutes of discussio ns in meetings with universiti es//instit utes official letters	2.3.2.2.1. 1 4/5	2.3.2.2.1. 1 6 months	2.3.2.2.1. 1 KEPA Research office	2.3.2.2.1. 1 KEPA Research office	2.3.2.2.1. 1 None	2.3.2.2.1. 1 None	2.3.2.2.1. 1 % of research meetings with universiti es//instit utes where AMR is requeste d to be apriority in research projects.
	research	es/institu tes 2.3.2.2.1. 2 Call for proposal from universiti es//instit utes	2.3.2.2.1. 2 Letter to universiti es/institu tes from KEPA research director to send projects related to AMR	2.3.2.2.1. 2 1/univers ity	2.3.2.2.1. 2 1 year	2.3.2.2.1. 2 KEPA research office directora te	2.3.2.2.1. 2 KEPA research office directora te	2.3.2.2.1. 2 None	2.3.2.2.1. 2 None	2.3.2.2.1. 2 Proportio n of universiti es /institute s that send projects related to AMR in

		in environ ment							environm ent to KEPA research office
3 Th dii of of re an sti ap fo fre KE pr re to su nc Ve ag re En	he irector f the ffice of esearch	2.3.2.2.1. 3 Projects and funds	2.3.2.2.1. 3 Multiple	2.3.2.2.1. 3 6 months	2.3.2.2.1. 3 KEPA research office	2.3.2.2.1. 3 KEPA research office	2.3.2.2.1. 3 None	2.3.2.2.1. 3 None	2.3.2.2.1. 3 % of proposed projects related to AMR in environm ent that have been funded
Increase 1 Le capacity of KEPA KE Lab that Di	etter om EPA irector eneral	2.3.2.2.2. 1 Letter from KEPA General director to	2.3.2.2.2. 1 1	2.3.2.2.2. 1 3 months	2.3.2.2.2. 1 KEPA	2.3.2.2.2. 1 KEPA	2.3.2.2.2. 1 None	2.3.2.2.2. 1 None	2.3.2.2.2. 1 KEPA Lab starts reporting AMR data

quality of coastal water to include AMR identifica tion and surveilla nce in water and sediment	Technical affairs sector of micro labs asking to prepare a project with a budget that aims at expandin g their work from only identifica tion to antibiogr am		technical affairs sector							
	testing 2.3.2.2.2. 2 KEPA analytical central lab KEPA prepare road map for the labs to be able to do antibiogr	2.3.2.2.2. 2.1 Identify the needed machiner y and their budget	2.3.2.2.2. 2.1 List and budget	2.3.2.2.2. 2.1 1	2.3.2.2.2. 2.1 6 months	2.3.2.2.2. 2.1 KEPA Analytica I Lab center	2.3.2.2.2. 2.1 KEPA Analytica I Lab center	2.3.2.2.2. 2.1 None	2.3.2.2.2. 2.1 None	2.3.2.2.2. 2 Road map project with budget is put to extend KEPA microbiol ogy lab from

	am testing									bacterial identifica tion to include also antibiogr am testing
		2.3.2.2.2 2.2 Put a plan of needed extra staffing	2.3.2.2.2. 2.2 Plan	2.3.2.2.2. 2.2 1	2.3.2.2.2. 2.2 6 months	2.3.2.2.2. 2.2 KEPA Lab council	2.3.2.2.2. 2.2 KEPA Lab council	2.3.2.2.2. 2.2 None	2.3.2.2.2. 2.2 None	See above
	2.3.2.2.2. 3 KEPA provides the related budget	2.3.2.2.2. 3 Communication between KEPA and ministry of finance	2.3.2.2.2. 3 Letters	2.3.2.2.2.3	2.3.2.2.2. 3 2 years	2.3.2.2.2. 3 KEPA MO finance	2.3.2.2.2. 3 KEPA MO finance	2.3.2.2.2. 3 None	2.3.2.2.2. 3 None	2.3.2.2.2. 3 None
	2.3.2.2.2. 4 KEPA microbiol ogy labs start generati ng antibiogr am results for		2.3.2.2.2. 4 Antibiogr am KEPA LABS	2.3.2.2.2. 4 1/LAB	2.3.2.2.2. 4 3 years	2.3.2.2.2. 4 KEPA Micro lab	2.3.2.2.2. 4 KEPA	2.3.2.2.2. 4 Will be determin ed in the Road map project	2.3.2.2.2. 4 KEPA	2.3.2.2.2. 4 KEPA Marine lab actually generate s AMR data

		priority organism s								
	2.3.2.2.3 Generate report about AMR surveilla nce in Environ ment		2.3.2.2.3 Report	2.3.2.2.3	2.3.2.2.3 4 years	2.3.2.2.3 KEPA	2.3.2.2.3 KEPA	2.3.2.2.3 None	2.3.2.2.3 None	2.3.2.2.3 Yes/No
2.3.3 The national AMR Surveilla nce 2.3.3.1 Office at KEPA, MOH PAAF and gathers PAFN 3 data submit available KEPA, PAAF and Central PAFN AMR Scompiles surveilla	2.3.3.1.1 Create a digital network for data sharing with MOH		2.3.3.1.1 Digital system	2.3.3.1.1	2.3.3.1.1 5 years	2.3.3.1.1 MOH, KEPA, PAAF PAFN	2.3.3.1.1 IT team at MOH, KEPA, PAAF PAFN	2.3.3.1.1 none	2.3.3.1.1 none	2.3.3.1.1 Digital System is put

surveilla nce report.	2.3.3.1.2 Add this function to the TOR of the focal person at PAAF and KEPA and food safety.			2.3.3.1.2 TOR	2.3.3.1.2	2.3.3.1.2 3 years	2.3.3.1.2 MOH/ KEPA/ PAAF/ PAFN	2.3.3.1.2 MOH/ KEPA/ PAAF/ PAFN	2.3.3.1.2 None	2.3.3.1.2 None	2.3.3.1.2 TOR includes surveilla nce function detail.
2.4.1 To have A Kuwaiti Center of Disease Control that will be with its overseei ng bacterial viral and AMR referenc e labs	t d	ational Refe	rence Lab for	2.4.1.1 Project endorsed by MOH	2.4.1.1	2.4.1.1 Already available	2.4.1.1 Kuwait CDC Lab	2.4.1.1 MOH	2.4.1.1 Included in the project, already approved by Minister of Health	2.4.1.1 MOH	2.4.1.1 Kuwait CDC will oversee the needs and functions of all referenc e laborator ial, including viral, bacterial and AMR.

2.4.1.2 Kuwait CDC becomes functiona		2.4.1.2 Function al Kuwait CDC	2.4.1.2	2.4.1.2 5 years	2.4.1.2 Kuwait CDC	2.4.1.2 MOH/ Minister of Health	2.4.1.2 included in the approved project approved by minister of health	2.4.1.2 Council of Ministers /MOH and Ministry of Finance.	2.4.1.2 Kuwait CDC is establish ed
2.4.1.3 Write and approve terms of referenc e for a national AMR referenc e laborator y with expertise in method s for confirmi ng and characte izing specific pathoger s, putting guideline s, SOP for		2.4.1.3 TOR OF Referenc e Lab	2.4.1.3	2.4.1.3 3 months	2.4.1.3 Kuwait CDC lab	2.4.1.3 Kuwait CDC/Min ister of health/M edical Lab.coun cil	2.4.1.3 none	2.4.1.3 none	2.4.1.3 TOR of AMR referenc e Lab are put. They include: expertise in method s for confirmi ng and character izing specific pathogen s, putting guideline s, SOP for identifica tion and antibiogr am

identifition an antibio am testing priority organis s for surveill nce training according to guideling s, and organize g quality assurare e scheme	of m e n									testing of priority organism s for surveilla nce training accordin g to guideline s, and organizin g quality assuranc e schemes
2.4.1.4 Prepare a full project and propos for the referen e lab	ı		2.4.1.4 Referenc e Lab project as part of Kuwait CDC	2.4.1.4	2.4.1.4 5 years	2.4.1.4 Kuwait CDC	2.4.1.4 Kuwait CDC	2.4.1.4 project already approved	2.4.1.4 Kuwait CDC budget already approved	2.4.1.4 Yes/No
2.4.1.5 Prepare a budg for the building equipm nt and	t ,		2.4.1.5 Budget	2.4.1.5	2.4.1.5 5 years	2.4.1.5 Kuwait CDC	2.4.1.5 Kuwait CDC	2.4.1.5 Budget within the budget of Kuwait CDC	2.4.1.5 MOH/ Council of ministers /Ministry	2.4.1.5 Budget for AMR lab is approved

man power of the referenc e Lab							of Finance.	
2.4.1.6 Executio n of the Referenc e Lab		2.4.1.6 1	2.4.1.6 5 years	2.4.1.6 Kuwait CDC	2.4.1.6 Kuwait CDC	2.4.1.6 Budget within the budget of Kuwait CDC	2.4.1.6 MOH/ Council of ministers /Ministry of Finance.	2.4.1.6 Referenc e lab is set and functioni ng

Delegate the function of surveilla nce of the reference Lab pending its readiness	2.4.2.1 Identify institutio n affiliated to MOH that will host the manpow er and activities of the AMR Surveilla nce office for the actual work on the National and GLASS AMR reports pending the finalizati on of the	2.4.2.1.1 A letter from MOH to all labs that asks them to coordina te with the GLASS lab by deliverin g AMR data for the purpose of AMR Surveilla nce		2.4.2.1.1 Letter from MOH to hospitals through Medical lab council	2.4.2.1.1	2.4.2.1.1 Already available	2.4.2.1.1 MOH	2.4.2.1.1 MOH	2.4.2.1.1 None	2.4.2.1.1 None	2.4.2.1.1 Already available

	referenc e lab. bjective 2.5	: Unify AMR	identification	on in all Micr	o labs							
2.5.1 Lab audit by lab accredita tion body accordin g to microbiol ogist consulta nt in charge, external quality control and use of CLSI guideline s	2.5.1.1 Accredita tion of the medical Laborato ries by Medical Laborato ries Council	2.5.1.1.1 Involve all laborator ies in external quality assuranc e programs	2.5.1.1.1. 1 Mandate that all labs including all private should be accredite d and all perform external QC.	2.5.1.1.1. 1.1 List of accredita tion bodies accepted in Kuwait, all labs should be accredite d by one of them	2.5.1.1.1. 1.1 List	2.5.1.1.1. 1.1 1	2.5.1.1.1. 1.1 6 months	2.5.1.1.1 1.1 Central Labs/MO H/ Medical Lab.coun cil	2.5.1.1.1 1.1 Central Labs/MO H/Medic al Lab.coun cil	2.5.1.1.1. 1.1 None	2.5.1.1.1. 1.1 None	2.5.1.1.1 1.1 Number of internati onal accredita tion bodies that are recomme nded in Kuwait for the labs to get their accredita tion

N.B. Refer to Annex 3 for AMR Surveillance Axis Strategic and Monitory plans

<u>Infection Prevention and Control Operational and Budget Plan</u>

Global action plan strategic objective 3: Reduce the incidence of infection through effective sanitation, hygiene and prevention measures.

Strategic intervent ions	Activity	Sub- activity	Sub-sub- activity	Sub-sub- sub- activity	Unit	Quantity	Date	Location	Responsi ble entity	Cost	Source Of funding	Indicator
Strategic C	bjective 3.1	Emphasize	the national	infection pr		d control pro	ogram in hea	althcare at the	he National	and Health	are facility	levels.
3.1.1 Create a formal organizat ional structure to ensure proper develop ment and use of infection preventi on and control policies and strategie s	3.1.1.1 Infection Control Organiza tional structure is already establish ed and functioni ng.	-	-	-	3.1.1.1 IPC Directora te in MOH that is a central, and fulfills the criteria of National IPC body that coordina tes and organizes IPC activities in HCF.	3.1.1.1	3.1.1.1 Done 1980	3.1.1.1 MOH	3.1.1.1 MOH /IPC Directora te	3.1.1.1 Done	3.1.1.1 none	3.1.1.1 Already available
	3.1.1.2 To make official that IPC is part of the organogr	3.1.2.1 Decree from MOH about IPC departm			3.1.2.1 Decree from MOH	3.1.2.1	3.1.2.1 2 Months	3.1.2.1 MOH	3.1.2.1 MOH /IPC Directora te	3.1.2.1 None	3.1.2.1 None	3.1.2.1 The decree was issued.

am of each hospital with Terms of Referenc e and allocated staff	ent in each hospital with TOR, and staffing organizat ion										
3.1.1.3 IPC departm ents in hospitals and they work closely with the central IPC Directora te	3.1.1.3.1 To impose IPC program s in all hospitals	3.1.1.3.1. 1 IPC program availabilit y is checked by accredita tion standard s		3.1.1.3.1. Recomm endation availabilit y in accredita tion standard s	3.1.1.3.1. 1	3.1.1.3.1. 1 Already available	3.1.1.3.1. 1 MOH/ Quality and Accredita tion Directora te	3.1.1.3.1. 1 MOH / Quality and Accredita tion Directora te	3.1.1.3.1. 1 None	3.1.1.3.1. 1 No need	3.1.1.3.1. 1 Exists already
	3.1.1.3.2 IPC program s are available in all hospitals and they work under the umbrella of the	-	-	3.1.1.3.2 IPC Program/ Departm ent	3.1.1.3.2 One in each hospital	3.1.1.3.2 Already available	3.1.1.3.2 IPC directora te/Hospit als	3.1.1.3.2 MOH/IPC directora te/Hospit als	3.1.1.3.2 None	3.1.1.3.2 None	3.1.1.3.2 Number of Hospitals with IPC program s and Total number of hospital departm ents

		Central IPC directora te.										
3.1.2 Impleme nt laws for mandato ry complian ce of the facilities with IPC standard s	3.1.2.1 Decree from MOH	3.1.2.1.1 Decree that recomm ends that all HCF have to abide by IPC Recomm endation s.	3.1.2.1.1. 1 IPC Directora te puts the detailed IPC measures that should be required from each HCF, and sent to MOH to be included in ministry	-	3.1.2.1.1. 1 List of measure s	3.1.2.1.1. 11	3.1.2.1.1. 1 2 months	3.1.2.1.1. 1 MOH/IPC directora te	3.1.2.1.1. 1 MOH/IPC directora te	3.1.2.1.1. 1 No cost	3.1.2.1.1. 1 No cost	3.1.2.1.1. 1 List is sent from IPC Directora te to MOH undersec retary
			3.1.2.1.1. 2 Decree is issued by MOH	-	3.1.2.1.1. 2 Decree	3.1.2.1.1. 2 1	3.1.2.1.1. 2 3 months	3.1.2.1.1. 2 MOH	3.1.2.1.1. 2 MOH	3.1.2.1.1. 2 None	3.1.2.1.1. 2 None	3.1.2.1.1. 2 Decree availabili ty and dissemin ation
		3.1.2.1.2 Medical responsi bilities.	3.1.2.1.2. 1 Add complian ce to IPC	-	3.1.2.1.2. 1 Update of medical	3.1.2.1.2. 1 1	3.1.2.1.2. 1 3 months	3.1.2.1.2. 1 MOH	3.1.2.1.2. 1 IPC Directora te/MOH	3.1.2.1.2. 1 None	3.1.2.1.2. 1 None	3.1.2.1.2. 1 The Medical responsi

		Add details IPC standard complian ce in hospitals	measures and recomme ndations to the list of medical responsi bilities of hospitals as listed by MOH		responsi bilities list issued by MOH							bility list is updated and it includes the statemen t about IPC measure s.
3.1.3 WHO IPC Core compone nt based guideline s are establish ed in all HCF	3.1.3.1 Availabili ty of endorsed IPC based on WHO IPC core compone nts	-	-	-	3.1.3.1 1 available guideline s	3.1.3.1 1 Central IPC Guideline	3.1.3.1 Done	3.1.3.1 MOH	3.1.3.1 MOH/ Directora te of IPC	-	3.1.3.1 MOH	3.1.3.1 Available
	3.1.3.2 IPC policies and procedur es are included in hospital accredita tion standard s	-	-		3.1.3.2 1 Available list	3.1.3.2 1 list	3.1.3.2 Done	3.1.3.2 MOH	3.1.3.2 MOH/ Directora te of IPC	-	3.1.3.2 MOH	3.1.3.2 Available

3.1.3.3 IPC policies and procedur es are being audited regularly by accredita tion bodies	-	-	-	3.1.3.3 1 audit tool Available	3.1.3.3 1 audit tool	3.1.3.3 Done	3.1.3.3 MOH	3.1.3.3 MOH / Quality and Accredita tion Directora te	-	3.1.3.3 MOH	3.1.3.3 Available
3.1.3.4 National infection control policies and guideline s are distribut ed and periodica lly updated	3.1.3.4.1 Regular update of national IPC guideline s	3.1.3.4.1. 1 On going, but needs to be speeded up by increasin g manpow er in IPC nationwi de in central office and in hospitals	-	3.1.3.4.1. 1 Employin g new Inspector s, MDs and IPC professio nals with data entry personne I	3.1.3.4.1. 1 -10 MD -4 data entry personne I -15 inspector s	3.1.3.4.1. 1 2 years	3.1.3.4.1. 1 MOH	3.1.3.4.1. 1 IPC directora te/MOH	3.1.3.4.1. 1 MOH	3.1.3.4.1. 1 MOH	3.1.3.4.1. 1 Number employe d/Numb er needed
	3.1.3.4.2 A direct channel with	3.1.3.4.2. 1 Communi cate this	3.1.3.4.2. 1.1 Letter from	3.1.3.4.2. 1.1 Letter	3.1.3.4.2. 1.1 1	3.1.3.4.2. 1.1 1 month	3.1.3.4.2. 1.1	3.1.3.4.2. 1.1 WCO	3.1.3.4.2. 1.1	3.1.3.4.2. 1.1 WHO	3.1.3.4.2. 1.1 Number of

WHO consulta nt regardin g few queries and new recomm endation s	need to WHO office in Kuwait	MOH/IPC directora te to WHO office in Kuwait asking for direct channel with a consulta nt for queries wheneve r they occur				WHO office Kuwait		Will be costed by WHO		answere d queries /Number of needed consultat ions
	3.1.3.4.2. 2 Communi cate this need to Regional WHO office	3.1.3.4.2. 2 Request from WCO to Regional office.	3.1.3.4.2. 2 Letter	3.1.3.4.2. 2 1	3.1.3.4.2. 2 2 months	3.1.3.4.2. 2 WHO office Kuwait	3.1.3.4.2. 2 WCO	3.1.3.4.2. 2 Will be costed by WHO	3.1.3.4.2. 2 WHO	
3.1.3.4.3 Regular update of National IPC policies and procedur es	3.1.3.4.3. 1 Need updating + speeding up the process		3.1.3.4.3. 1 provide necessar y manpow er as in 3.1.3.4.1.	3.1.3.4.3. 1 As above Manpow er 3.1.3.4.1.	3.1.3.4.3. 1 ongoing	3.1.3.4.3. 1 MOH	3.1.3.4.3. 1 Plan of manpow er Above As in 3.1.3.4.1.	3.1.3.4.3. 1 Employe es salaries	3.1.3.4.3. 1 MOH	3.1.3.4.3. 1 Number of hospitals with updated policies/ Total number of hospitals

3.1.3.5 National IPC Guideline s and policies are being taught	3.1.3.5.1 TOT periodica lly performe d nationall y	-	-	3.1.3.5.1 Training sessions	3.1.3.5.1 Started years ago, and running	3.1.3.5.1 Already being performe d	3.1.3.5.1 MOH/IPC directora te/ Hospitals	3.1.3.5.1 IPC Directora te	3.1.3.5.1 Already budget is available and being performe d	3.1.3.5.1 MOH	3.1.3.5.1 Number of IPC professio nals trained annually /total number of IPC professio nals.
	3.1.3.5.2 Continuo us training of HCW about IPC policies in hospitals is available	3.1.3.5.2. 1 Mandate from MOH that it is mandato ry to all HCW to attend at least 1 yearly and 1 upon employment		3.1.3.5.2. 1 Mandate	3.1.3.5.2. 1 1	3.1.3.5.2. 1 2 months	3.1.3.5.2. 1 MOH	3.1.3.5.2. 1 IPC Directora te/ MOH	3.1.3.5.2. 1 None	3.1.3.5.2. 1 None	3.1.3.5.2. 1 Decree is done and sent to hospitals
		3.1.3.5.2. 2 Improve attendan ce to these	3.1.3.5.2. 2.1 Attendan ce is reported to hospital	3.1.3.5.2. 2.1 Report presente d to hospital directors	3.1.3.5.2. 2.1 report/h ospital/y ear	3.1.3.5.2. 2.1 1 year	3.1.3.5.2. 2.1 MOH/ Hospitals	3.1.3.5.2. 2.1 Hospital IPC office	3.1.3.5.2. 2.1 None	3.1.3.5.2. 2.1 No	3.1.3.5.2. 2.1 Number of HCW attendee s /year

			general HCW IPC training sessions	administr ators and action taken to improve attendan ce								
				3.1.3.5.2. 2.2 Attendan ce evidence is available in yearly Hospital IPC report and monitore d by IPC Directora te	3.1.3.5.2. 2.2 Report presente d to IPC directora te	3.1.3.5.2. 2.2 1 Report/h ospital/y ear	3.1.3.5.2. 2.2 1 year	3.1.3.5.2. 2.2 IPC Directora te	3.1.3.5.2. 2.2 MOH/IPC directora te/Hospit al administr ators	3.1.3.5.2. 2.2 None	3.1.3.5.2. 2.2 No	3.1.3.5.2. 2.2 none
3.1.4 Monitor and evaluate complian ce with the IPC guideline s	3.1.4.1 Checklist s and audit tools are prepared and applied	-	-	-	3.1.4.1 documen ts (checklist s and audit tools)	3.1.4.1	3.1.4.1 Started years ago, and ongoing for new policies.	3.1.4.1 MOH/IPC Directora te	3.1.4.1 IPC Directora te	3.1.4.1 none	3.1.4.1 none	3.1.4.1 Number of policies with audit tools/Tot al number of policies

3.1.4.2 Regular update of these audit tools	tools prepared and dissemin ated and check old tools if they need update	-	-	3.1.4.2.1 Audit tool	3.1.4.2.1 One/new policy. Then total review once/yea r	3.1.4.2.1 Ongoing	3.1.4.2.1 IPC Directora te/ MOH	3.1.4.2.1 IPC Directora te/ MOH	3.1.4.2.1 None	3.1.4.2.1 None	3.1.4.2.1 Number of new policies with audit tool/Tota I number of new policies.
3.1.4.3 Audit plan availab in hospita	tion	-	-	3.1.4.3.1 checking	3.1.4.3.1 1	3.1.4.3.1 done ongoing	3.1.4.3.1 IPC Directora te/ MOH	3.1.4.3.1 IPC Directora te/ MOH	3.1.4.3.1 None	3.1.4.3.1 None	3.1.4.3.1 Done and ongoing

D	Directora										
3. Co au pl bo pr ar ar di	is.1.4.3.2 Central audit blan is being brepared annually and listribut ed to all appropriate to spitals	-	-	3.1.4.3.2 Audit plan	3.1.4.3.2	3.1.4.3.2 Done and ongoing yearly	3.1.4.3.2 IPC Directora te	3.1.4.3.2 IPC Directora te /MOH	3.1.4.3.2 none	3.1.4.3.2 none	3.1.4.3.2 Done and ongoing
3. Al re se ce ce IP di te cc e	3.1.4.3.3 Audit eport is ent to entral PC lirectora e with orrectiv	-	-	3.1.4.3.3 Report	3.1.4.3.3 one/ hospital	3.1.4.3.3 Done and ongoing	3.1.4.3.3 Hospitals /IPC directora te	3.1.4.3.3 Hospitals /IPC directora te/MOH	3.1.4.3.3 None	3.1.4.3.3 None	3.1.4.3.3 Done and ongoing
3.1.4.4 Establish national process indicator s	3.1.4.4.1 st set of sprocess indicator in the control of the contr	3.1.4.4.1. 1 Put the basic training material	-	3.1.4.4.1. 1 Process indicator s of the Bundles of care	3.1.4.4.1. 13	3.1.4.4.1. 1Already available	3.1.4.4.1. IPC directora te	3.1.4.4.1. 1 IPC directora te	3.1.4.4.1. 1No extra budget	3.1.4.4.1. 1 IPC directora te for both	3.1.4.4.1. 1 Each one of the 3 procedur es has its own bundle establish ed.
		3.1.4.4.1. 2Update	3.1.4.4.1.	3.1.4.4.1. 2New	3.1.4.4.1. 2	3.1.4.4.1. 2	3.1.4.4.1. 2 IPC	3.1.4.4.1. 2MOH/	3.1.4.4.1. 2None	3.1.4.4.1. 2None	3.1.4.4.1. 2Availabi

	of the already available bundle of care in each of the mentione d bundles, by including new evidence-based items to the bundle	Prepare the updated bundles	bundle definitio n and checklist	3	2 years	directora te/MOH	IPC directora te			lity of 3 updated bundles
	3.1.4.4.1. 3 Train the IPC professio nals nationwi de on the updated bundles	3.1.4.4.1. 3.1 TOT worksho ps	3.1.4.4.1. 3.1 TOT sessions	3.1.4.4.1. 3.1 2/year centrally Total=4	3.1.4.4.1. 3.1 2 years	3.1.4.4.1. 3.1 MOH/IPC directora te	3.1.4.4.1. 3.1 IPC Directora te /MOH	3.1.4.4.1. 3.1 No need for additiona I at MOH	3.1.4.4.1. 3.1 MOH	3.1.4.4.1. 3.1 % IPC professio nals trained about the new bundles
	3.1.4.4.1. 4 Engage all HCW with the bundles by training,	3.1.4.4.1. 4.1 Prepare the ground in the hospitals:	3.1.4.4.1. 4.1 Letter for CLABSI bundle	3.1.4.4.1. 4.1 1 dissemin ated to all hospital directors	3.1.4.4.1. 4.1 CLABSI months Each bundle 1 y	3.1.4.4.1. 4.1 IPC Directora te/MOH	3.1.4.4.1. 4.1 IPC Directora te/MOH	3.1.4.4.1. 4.1 None	3.1.4.4.1. 4.1 None	3.1.4.4.1. 4.1 Letter sent or not

audit and feedback.	Letter to hospital directors about the bundles project its importance and its processe s								
	3.1.4.4.1. 4.2 Update the material to be taught to HCW by IPC professio nals, accordin g to the updated bundles.	3.1.4.4.1. 4.2 Training package for each bundle	3.1.4.4.1. 4.2 3 (1/bundl e)	3.1.4.4.1. 4.2 2 years	3.1.4.4.1. 4.2 IPC Directora te/MOH	3.1.4.4.1. 4.2 IPC Directora te/MOH	3.1.4.4.1. 4.2 None	33.1.4.4. 1.4.2 None	3.1.4.4.1. 4.2 % of material directed to HCW that has been updated.
	3.1.4.4.1. 4.3 Inhospital training sessions by IPC professionals to all	3.1.4.4.1. 4.3 Training sessions in hospitals to HCW performe d by each	3.1.4.4.1. 4.3 Multiple in each hospital	3.1.4.4.1. 4.3 4 years	3.1.4.4.1. 4.3 MOH/Ho spitals	3.1.4.4.1. 4.3 MOH/IPC directora te/Hospit als	3.1.4.4.1. 4.3 None	3.1.4.4.1. 4.3 None	3.1.4.4.1. 4.3 Complian ce with performa nce indicator for each bundle

concerne d HCW. Include Bundle training in the yearly IPC training and in peri- employm ent training.	hospital 's IPC team							21441
3.1.4.4.1. 4.4 Auditing complian ce to these bundles on monthly basis and report to IPC directora te monthly	3.1.4.4.1. 4.4 Audit activity/A udit report. (about the updated bundles and updated checklists)	.1.4.4.1.4 .4 3/month s/hospita	3.1.4.4.1. 4.4 5 years	.1.4.4.1.4 .4 MOH/Ho spitals	.1.4.4.1.4 .4 MOH/Ho spitals/IP C Directora te	.1.4.4.1.4 .4 None	.1.4.4.1.4 .4 None	3.1.4.4.1. 4.4 Process Indicator for each of the 3 bundles: For each bundle: % Complian ce by the specific bundle /opportu nities of audit for each bundle

3.1.4.4.2 Update outcome indicator s for the 3 bundles above (CLABSI, CAUTI, VAP) accordin g to new CDC/ guideline s	3.1.4.4.2. 1 Review updated CDC outcome indicator s and update already measure d bundles outcome indicator s according ly	-	3.1.4.4.2. 1 Reviewe d outcome indicator, calculatio n and benchma rking	3.1.4.4.2. 1 3	3.1.4.4.2. 1 6 months	3.1.4.4.2. 1 MOH/IPC directora te	3.1.4.4.2. 1 MOH/IPC directora te	3.1.4.4.2. 1 None	3.1.4.4.2. 1 MOH budget	3.1.4.4.2. 1 Number of outcome indicator measure ments and benchmark is updated.
3.1.4.4.3 Integrate process and outcome indicator s for each bundle, and use in the analysis and corrective actions	3.1.4.4.3. 1 Put a road map for integrati ng analysis of outcome and process indicator s related to the bundles	-	3.1.4.4.3. 1 Protocol	3.1.4.4.3. 1 3	3.1.4.4.3. 1 1 year	3.1.4.4.3. 1 MOH/IPC directora te	3.1.4.4.3. 1 MOH/IPC directora te	3.1.4.4.3. 1 None	3.1.4.4.3. 1 MOH budget	3.1.4.4.3. 1 % of bundles whose analysis integrate s process and outcome data
3.1.4.4.4	3.1.4.4.4. 1	-	3.1.4.4.4. 1	3.1.4.4.4. 1	3.1.4.4.4. 1	3.1.4.4.4. 1	3.1.4.4.4. 1	3.1.4.4.4. 1	3.1.4.4.4. 1	3.1.4.4.4. 1

Introduc e SSI bundle	preparati on of material		Protocol, Checklist, teaching material for IPC professio nals and HCW	1 Central	4 years	MOH/IPC directora te	MOH/IPC directora te	None	MOH budget	% of material that is prepared
	3.1.4.4.4. 2 Impleme ntation of SSI bundle	3.1.4.4.4. 2.1 Teaching of IPC professionals by central IPC directorate	3.1.4.4.4. 2.1 Teaching sessions and worksho ps	3.1.4.4.4. 2.1 Central	3.1.4.4.4. 2.1 5 years	3.1.4.4.4. 2.1 IPC directora te/MOH	3.1.4.4.4. 2.1 MOH/IPC directora te /Hospital s	3.1.4.4.4. 2.1 None	3.1.4.4.4. 2.1 MOH	3.1.4.4.4. 2.1 % hospitals start submitti ng process and outcome indicator s for SSI.
		3.1.4.4.4. 2.2 Teaching of healthcar e providers inside the hospitals by IPC professio nals	3.1.4.4.4. 2.2 Teaching sessions and worksho ps	3.1.4.4.4. 2.2 2/y in hospitals	3.1.4.4.4. 2.2 5 years	3.1.4.4.4. 2.2 MOH/Ho spitals	3.1.4.4.4. 2.2 MOH/IPC directora te /Hospital s	3.1.4.4.4. 2.2 None	3.1.4.4.4. 2.2 MOH	3.1.4.4.4. 2.2 % hospitals start submitti ng process and outcome indicator s for SSI.
		3.1.4.4.4. 2.3	3.1.4.4.4. 2.3	3.1.4.4.4. 2.3	3.1.4.4.4. 2.3 5 years	3.1.4.4.4. 2.3 Hospitals	3.1.4.4.4. 2.3	3.1.4.4.4. 2.3 None	3.1.4.4.4. 2.3 MOH	3.1.4.4.4. 2.3

				Data collectio n	Data collectio n	1			MOH/IPC directora te /Hospital s			% hospitals start submitti ng process and outcome indicator s for SSI.
3.1.5 IPC programs /office in LTCF working in close cooperati on with IPC directora te in MOH	3.1.5.1 Mandate from MOH that IPC program/ with employe e with TOR and specified budget are available in LTCF the central and the "branche s".	3.1.5.1.1 Communication between MOH and Public Authorit y of Disability Affairs regardin g IPC in LTCF and being overseen by the IPC directora te	3.1.5.1.1. 1 Letter from MOH to Public Authority of Disability Affairs asking to establish IPC in each LTCF and to have the personne I work closely with IPC directora te delegate to LTCF.		3.1.5.1.1. 1 Letter	3.1.5.1.1. 1	3.1.5.1.1. 1 2 months	3.1.5.1.1. 1 MOH/ Public Authority of Disability Affairs	3.1.5.1.1. 1 MOH/ Main LTCF	3.1.5.1.1. 1 No	3.1.5.1.1. 1 No	3.1.5.1.1. 1 Present

	3.1.5.1.1. 2 Letter from Public Authority of Disability Affairs to LTCF that recomme nds IPC in each LTCF and organizes the communi cation with MOH		3.1.5.1.1. 2 Letter	3.1.5.1.1. 2 1	3.1.5.1.1. 2 2 months	3.1.5.1.1. 2 Public Authority of Disability Affairs	3.1.5.1.1. 2 Public Authority of Disability Affairs	3.1.5.1.1. 2 No	3.1.5.1.1. 2 No	3.1.5.1.1. 2 Letter sent
3.1.5.1.2 Establish ed Guidelin es for IPC in LTCF	3.1.5.1.2. 1 Preparati on of the LTCF guideline s and their endorse ment by Public Authority of Disability Affairs	-	3.1.5.1.2. 1 Guideline s Policies Procedur es	3.1.5.1.2. 1	3.1.5.1.2. 1 1 year	3.1.5.1.2. 1 IPC Directora te/ Public Authority of Disability Affairs	3.1.5.1.2. 1 MOH/IPC Directora te	3.1.5.1.2. 1 None	3.1.5.1.2. 1 None	3.1.5.1.2. 1 Availabili ty of the endorsed guideline s and set of policies and procedur es

Strategic C	Objective 3.2	To have the	3.1.5.1.2. 2 Dissemin ation of LTCF Policies and Procedur es	3.1.5.1.2. 2.1 TOT of LTCF IPC focal persons	3.1.5.1.2. 2 Worksho p once/y	3.1.5.1.2. 2 1/year	3.1.5.1.2. 2 1.5 years	3.1.5.1.2. 2 Public Authority of Disability Affairs	3.1.5.1.2. 2 MOH/ Public Authority of Disability Affairs	3.1.5.1.2. 2 20,000 KD	3.1.5.1.2. 2 MOH/ Public Authority of Disability Affairs	3.1.5.1.2. 2 % focal IPC persons in LTCF trained on the guideline s/total number of LTCF IPC focal persons.
3.2.1 Software for hospital data entry, analysis and generati ng both hospital and national reports	3.2.1.1 Subcom mittee from IPC directora te sends a letter to Informati on system departm ent at MOH explainin g what is required and asking				3.2.1.1 Letter	3.2.1.1	3.2.1.1 2 months	3.2.1.1 MOH	3.2.1.1 MOH/ IPC directora te	3.2.1.1 None	3.2.1.1 None	3.2.1 Software develope d and working

for the software									
3.2.1.2 Informati on system dep. presents projects (either in-house, or ready to buy) with budget providing		3.2.1.2 Project	3.2.1.2	3.2.1.2 12 months	3.2.1.2 MOH	3.2.1.2 MOH/ Informati on system departm ent	3.2.1.2 To be determin ed	3.2.1.2 MOH	See above
3.2.1.3 Agreeme nt between IPC directora te and informati on system dep. to choose one system		3.2.1.3 Agreeme nt letter	3.2.1.3	3.2.1.3 1 year 3 months	3.2.1.3 MOH	3.2.1.3 MOH/ IPC directora te/Healt h Informati on system departm ent	3.2.1.3 None	3.2.1.3 None	See above
3.2.1.4 Apply the system in the hospitals		3.2.1.4 System applicati on	3.2.1.4	3.2.1.4 1 year 9 months	3.2.1.4 Hospitals	3.2.1.4 MOH/ IPC directora te/ Informati	3.2.1.4 To be determin ed by the MOH	3.2.1.4 MOH	See above

									on system departm ent/Hosp itals			
	3.2.1.5 Apply the Digital HAI/AMR System for National data collectio n from individua I hospitals to central IPC directora te				3.2.1.5 System applicati on	3.2.1.5	3.2.1.5 2 years	3.2.1.5 Hospitals	3.2.1.5 MOH/ IPC directora te/Infor mation system departm ent/Hosp itals	3.2.1.5 To be determin ed by the MOH	3.2.1.5 MOH	See above
3.2.2 To Shift all hospital informati on system from paper to digital mode.	3.2.2.1 MOH recomme nds from hospitals to have an electroni c patient and data informati on system.	3.2.2.1.1 Letter from MOH to hospitals	-	-	3.2.2.1.1 Letter	3.2.2.1.1 1 to each hospital	3.2.2.1.1 2 months	3.2.2.1.1 MOH/Ho spitals	3.2.2.1.1 MOH/ Informati on system dep.	3.2.2.1.1 None	3.2.2.1.1 None	3.2.2.1.1 Letter is sent

	3.2.2.2 Increase the number of hospitals that have digitalize d their medical informati on system	-	-	-	3.2.2.2 Digital informati on system in each hospital	3.2.2.2 1/hospit al	3.2.2.2 5 years	3.2.2.2 Hospitals	3.2.2.2 Hospitals /MOH/ Informati on system dep.	3.2.2.2 to be determin ed by MOH	3.2.2.2 Private hospitals: the hospitals themselv es Public hospitals: MOH	3.2.2.2 % hospitals with digital informati on system
3.2.3 Connect Infection Control Departm ents to electroni c medical records	3.2.3.1 Connect Infection Control departm ents in all hospitals with the HIS and LIS at the hospital level	3.2.3.1.1 Letter from MOH to ask all hospitals to connect IC dep. with HIS and LIS at the hospital level	-	-	3.2.3.1.1 Letter	3.2.3.1.1 all private hospitals all governm ental hospitals	3.2.3.1.1 2 months	3.2.3.1.1 MOH	3.2.3.1.1 Hospitals MOH/Inf ormation system dep.	3.2.3.1.1 None	3.2.3.1.1 None	3.2.3.1.1 % IC dep connecte d with HIS/LIS in hospitals
		3.2.3.1.2 Decree from MOH to all hospitals who have an electroni	-	-	3.2.3.1.2 Letter	3.2.3.1.2 all private hospitals / all governm ental hospitals	3.2.3.1.2 4 months	3.2.3.1.2 MOH	3.2.3.1.2 MOH	3.2.3.1.2 No	3.2.3.1.2 Hospitals	3.2.3.1.2 % IC dep connecte d with electroni c AMR data in hospitals

	c Health informati on system to link their Lab AMR data to the Hospital system									
3.2.3.2 Develop electroni c outbreak reporting system for data entry, analysis and generati ng reports.			3.2.3.2 electroni c outbreak reporting system	3.2.3.2 1 central	3.2.3.2 3 years	3.2.3.2 MOH	3.2.3.2 MOH /Informa tion system departm ent	3.2.3.2 to be determin ed by MOH	3.2.3.2 MOH	3.2.3.2 electroni c outbreak reporting system available
3.2.3.3 Develop electroni c reporting system for data entry, analysis and			3.2.3.3 Electroni c system for data collectio n analysis and generati ng	3.2.3.3	3.2.3.3 3 years	3.2.3.3 MOH/Ho spitals	3.2.3.3 MOH /Informa tion system departm ent	3.2.3.3 to be determin ed by MOH	3.2.3.3 MOH	3.2.3.3 Bundles data collectio n, analysis and generati on of hospital /

	generati ng hospital/ national reports for bundles of care.				hospitals /national reports for bundles process and outcome indicator s							national reports is done through. an electroni c advance d system Yes/No
	3.2.3.4 Develop electroni c infection preventi on and control system capable of self-identifyin g HAI and detecting outbreak s	3.2.3.4.1 Electroni c system has a function for self diagnosis of HAIs and automati cally detects outbreak			3.2.3.4.1 Function in the electroni c system	3.2.3.4.1	3.2.3.4.1 5 years	3.2.3.4.1 MOH/Ho spitals	3.2.3.4.1 MOH/ Informati on system departm ent	3.2.3.4.1 to be determin ed by MOH	3.2.3.4.1 MOH	3.2.3.4.1 % of HAIs and outbreak s in hospitals are automati cally detected by the IT system
		Enhance IP	C education	in all health-	related spe	cialties and	among HCW	l				
3.3.1 Include IPC as a core element in educatio n and	3.3.1.1 Include infection control in undergra duate curricula for	3.3.1.1.1 Prepare module about IPC and AMR to be included	-	-	3.3.1.1.1 Educatio nal IPC module	3.3.1.1.1	3.3.1.1.1 2 years	3.2.1.1.1 MOH	3.3.1.1.1 MOH/IPC directora te	3.3.1.1.1 None	3.3.1.1.1 MOH	3.3.1.1.1 Module is available

training of healthcar e professio nals	health care students (medical, nursing, dental and pharmac eutical).	in all health specialtie s in universiti es/PAAE T.										
		3.3.1.1.2 MOH letter to MOHE and PAAET to ask to include this modules into the undergra ds and graduate health students' curricula	-	-	3.3.1.1.2 1 letter to MOHE and Letter to PAAET	3.3.1.1.2	3.3.1.1.2 2 years	3.2.1.1.2 MOH	3.3.1.1.2 MOH/IPC Directora te	3.3.1.1.2 None	3.3.1.1.2 No	3.3.1.1.2 Letters sent
		3.3.1.1.3 Modules sent to Kuwait universit y/ PAAET to include it in curricula			3.3.1.1.3 Module sent to Kuwait Universit y and PAAET	3.3.1.1.3	3.3.1.1.3 2.5 years	3.3.1.1.3 MOHE / PAAET	3.3.1.1.3 MOHE / PAAET	3.3.1.1.3 None	3.3.1.1.3 No	3.3.1.1.3 % health specialtie s that include IPC module

3.3.2 Enhance IPC educatio n among undergra duates and higher educatio n health students	3.3.2.1 Include infection control in continuin g educatio n of healthcar e workers and mandato ry prerequis ite for promotio n	3.3.2.1.1 Provide IPC courses with CE credits in universiti es, hospitals , MOH, PAAET PAAF Include CME Credit 10% IPC		3.3.2.1.1 IPC courses	3.3.2.1.1 At least once/yea r in universiti es /IPC directora te and PAAF	3.3.2.1.1 2 years	3.3.2.1.1 Universiti es/ Hospitals / IPC Directora te/ PAAET PAAF	3.3.2.1.1 IPC directora te/MOHE / PAAET / PAAF	3.3.2.1.1 No extra	3.3.2.1.1 Universiti es/ Hospitals /-PAAET / PAAF	3.3.2.1.1 Number of IPC courses/ year
		3.3.2.1.2 Request that when CE credits for physician s, nurses, or pharmaci		3.3.2.1.2 Decree	3.3.2.1.2 -1 for physician s -1 for Nurses -1 for pharmaci sts.	3.3.2.1.2 3 months	3.3.2.1.2 MOH	3.3.2.1.2 MOH/IPC Directora te	3.3.2.1.2 None	3.3.2.1.2 None	3.3.2.1.2 % professio nals have 10% of their CE credits are related to IPC

	sts is needed, 10% of these credits should be about IPC.										
	3.3.2.1.3 In hospitals , yearly and upon employm ent, mandato ry attendan ce of IPC lectures	-	-	3.3.2.1.3 Letter from MOH to all hospitals	3.3.2.1.3 1 to each hospital	3.3.2.1.3 3 months	3.3.2.1.3 MOH/Ho spitals	3.3.2.1.3 MOH/Ho spitals	3.3.2.1.3 None	3.3.2.1.3 None	3.3.2.1.3 % hospitals that mandate d that all employe e get IPC training upon employm ent and yearly IPC lecture.
3.3.2.2 Hygiene and IPC educatio n provided outside formal teaching programs	3.3.2.2.1 Require IPC educatio n by drug compani es, hospitals , LTCFs, FAO, and medical/ veterinar y	3.3.2.2.1. 1 Request to include few topics about IPC in conferen ces agendas	3.3.2.2.1. 11 Letter to be sent	3.3.2.2.1. 11 letter	3.3.2.2.1. 1 1 letter /associati on or entity	3.3.2.2.1. 1 1 6 months	3.3.2.2.1. 1 1 MOH	3.3.2.2.1. 1 1 MOH/Dir ectorate IPC	3.3.2.2.1. 1 1 None	3.3.2.2.1. 1 1 None	3.3.2.2.1. 1 1 Yes/No

		associati ons										
Strategic C	Objective 3.4	Prevent oc	cupational in	fections in I	ICWs			•				•
3.4.1 Develop a compreh ensive program of vaccinati on to protect HCWs from vaccine preventa ble diseases	-	-	-	-	3.4.1 HCW vaccinati on program	3.4.1 1 all inclusive	3.4.1 Available And applied	3.4.1 MOH	3.4.1 MOH/Pu blic Health directora te	3.4.1 Price of vaccines	3.4.1 MOH	3.4.1 % of HCWs professio nals that are vaccinate d
3.4.2 Develop policies and procedur es to protect healthcar e workers from occupati onal exposure to	-	-	-		3.4.2 policies and procedur es to protect healthcar e workers from occupati onal exposure to infection s	3.4.2	3.4.2 Available And applied	3.4.2 MOH	3.4.2 MOH	3.4.2 None	3.4.2 None	3.4.2 Yes/No

infection												
S.												
3.4.3 Availabili ty of Airborne Isolation Rooms in all hospitals	3.4.3.1 screenin g mapping and updating the recomme ndations accordin g to the expected need accordin g to need	-	-	-	3.4.3.1 Mapping and Updated recomme ndations	3.4.3.1 1/1	3.4.3.1 5 years	3.4.3.1 MOH/Ho spitals	3.4.3.1 MOH/En gineering Directora te	3.4.3.1 None	3.4.3.1 MOH for Public hospitals private hospitals: the hospitals themselv es.	3.4.3.1 Yes/No
3.4.4 Mandate to hospitals to provide airborne isolation rooms (AII) available as per the updated recomme ndations	-	-	-	-	3.4.4 Mandate	3.4.4 1/hospit al	3.4.4 5 years	3.4.4 MOH	3.4.4 MOH	3.4.4 none	3.4.4 none	3.4.4 Percent hospitals have enough AII rooms as per MOH recomm endation
	bjective 3.5	Adequate v	vaste manag	ement in ho	spitals							
3.5.1	3.5.1.1	3.5.1.1.1	3.5.1.1.1. 1	-	3.5.1.1.1. 1	3.5.1.1.1. 1	3.5.1.1.1. 1	3.5.1.1.1. 1	3.5.1.1.1. 1	3.5.1.1.1. 1	3.5.1.1.1. 1	3.5.1.1.1. 1

Put National Guideline s	National policies, strategie s and plans for healthcar e waste manage ment is in place	Unify recomm endation s between KEPA and MOH	Create a committe e from MOH and KEPA that unifies the recomme ndation and the road map of all types of waste that are generate d in healthcar e		Committ	1	6 months	MOH/KE PA	МОН	None	None	Committ ee members are nominat ed
			3.5.1.1.1. 2 The committe e reviews waste policies that are related to HCF and unifies recomme ndations and provides	-	3.5.1.1.1. 2 Revision	3.5.1.1.1. 2 1	3.5.1.1.1. 2 1 year	3.5.1.1.1. 2 KEPA	3.5.1.1.1. 2 KEPA	3.5.1.1.1. 2 None	3.5.1.1.1. 2 None	3.5.1.1.1. 2 Percent chapters that represen t conflict between recomm endation s of the 2 ministrie s that have been reviewed

			the proper ways of supervisi on									and unified
3.5.2 Supervisi on of waste disposal from public hospitals	3.5.2.1 Create a committ ee for the supervisi on of waste disposal from public hospitals	-	-	-			3.5.2.1 Available and ongoing					3.5.2.1 Available and ongoing
3.5.3 Supervisi on of waste disposal in private hospitals	3.5.3.1 Create a committ ee for the supervisi on of waste disposal in private hospitals	-	-	-	3.5.3.1 Committ ee	3.5.3.1	3.5.3.1 1 year	3.5.3.1 KEPA	3.5.3.1 MOH/KE PA	3.5.3.1 None	3.5.3.1 None	3.5.3.1 Yes/No
3.5.4 Waste policies are being audited by KEPA	3.5.4.1 Organiza tion of supervisi on of waste disposal	-	-	-	3.5.4.1 Audit plan	3.5.4.1 1	3.5.4.1 1.5 years	3.5.4.1 KEPA	3.5.4.1 KEPA	3.5.4.1 1 employe e KEPA	3.5.4.1 KEPA	3.5.4.1 % Public hospitals are being audited for waste disposal.

	inside and from private hospitals 3.5.4.2 Designat e 1 unit at KEPA to oversee the waste in hospitals		-	-	3.5.4.2 1 Unit At KEPA	3.5.4.2	3.5.4.2 1 year	3.5.4.2 KEPA	3.5.4.2 MOH/KE PA	3.5.4.2 to be determin ed	3.5.4.2 KEPA/Pri vate hospitals /MOH	3.5.4.2 % Private hospitals where waste manage ment is being supervise
Strategic C	biective 3.6	To enhance	infection pr	evention an	d control pr	ograms in ve	eterinary and	d agriculture	settings			d
3.6.1 Availability of IPC guidelines in veterinary practices	3.6.1.1 Impleme ntation of OIE guideline s regarding IPC in veterinar y field	3.6.1.1.1 Committ ee in PAAF that regularly checks updates in OIE guideline s regardin g IPC in veterinar y field, and updates the guideline s	-	-	3.6.1.1.1 IPC Committ ee in PAAF	3.6.1.1.1	3.6.1.1.1 6 months	3.6.1.1.1 PAAF	3.6.1.1.1 PAAF	3.6.1.1.1 None	3.6.1.1.1 None	3.6.1.1.1 Committ ee is formed

		accordin gly										
		3.6.1.1.2 Overseei ng the impleme ntation of national IPC guideline s in Vet field and OIE good practices	3.6.1.1.2 Regular audits to farms	3.6.1.1.2 Put a yearly audit plan	3.6.1.1.2 Plan	3.6.1.1.2	3.6.1.1.2 1 year	3.6.1.1.2 PAAF	3.6.1.1.2 PAAF	3.6.1.1.2 none	3.6.1.1.2 none	3.6.1.1.2 % Farms that are being audited yearly for IPC measure s.
3.6.2 Include IPC in licensing of Vets	3.6.2.1 Include question s about OIE IPC guideline s and vaccinati ons in all vets licensing exams or interview s.	3.6.2.1.1 Prepare a set of question s related to IPC in Vets to be included in licensing exams and interview s.	-	-	3.6.2.1.1 Set of question s	3.6.2.1.1 1	3.6.2.1.1 6 months	3.6.2.1.1 PAAF	3.6.2.1.1 IPC Committ ee in PAAF	3.6.2.1.1 none	3.6.2.1.1 none	3.6.2.1.1 Availabili ty of IPC question s in licensing exams (Yes/No)
3.6.3 Include IPC in Continuo us educatio	3.6.3.1 Prepare a slide bank related to IPC in	-	-	-	3.6.3.1 IPC slide bank	3.6.3.1 1	3.6.3.1 6 months	3.6.3.1 PAAF	3.6.3.1 PAAF	3.6.3.1 none	3.6.3.1 none	3.6.3.1 Yes/No

n of Vets and professio nals in Agricultu re	vet and agricultu re, to be given to drug compani es that usually do the continuo us educatio n activities.											
	3.6.3.2 Mandate that all drug company events or seminars include a specific number of slides related to IPC and AMR.	-	-		3.6.3.2 Mandate	3.6.3.2	3.6.3.2 9 months	3.6.3.2 PAAF	3.6.3.2 PAAF	3.6.3.2 None	3.6.3.2 None	3.6.3.2 Mandate available or not
3.6.4 Include hygiene and infection preventi on and	3.6.4.1 Close communi cation between. FAO and PAAF to	-	-	-	3.6.4.1 Yearly educatio n plan that targets vaccinati	3.6.4.1	3.6.4.1 1 year	3.6.4.1 PAAF	3.6.4.1 PAAF/FA O	3.6.4.1 None	3.6.4.1 None	3.6.4.1 Joint educatio n plan between FAO and PAAF

control as core content in training of veterinar y professio nals.	put a joint yearly plan of educatio n activities related to IPC.				on and IPC							that includes IPC- related training
	3.6.4.2 Include hygiene and infection preventi on and control as mandato ry prerequis ite for promotio n of for the veterinar y workers.	3.6.4.2.1 IPC credits mandato ry for promotio n in Vet. Associati ons	-	-	3.6.4.2.1 Mandate	3.6.4.2.1	3.6.4.2.1 6 months	3.6.4.2.1 Veterinar y associati on	3.6.4.2.1 PAAF/Ve terinary associati on	3.6.4.2.1 None	3.6.4.2.1 None	3.6.4.2.1 Mandate available or not

N.B. Refer to Annex 4 for IPC Axis Strategic and Monitory plans

Antimicrobial Use Operational and Budget Plan

Global action plan strategic objective 4: Optimize the use of antimicrobial medicines in human and animal health.

Activity	Sub- activity	Sub-sub- activity	Unit	Quantity	Date	Location	Responsibl e entity	Cost	Source Of funding	Indicator
Strategic Obj	ective 4.1 Ens	ure governance	e of antibiotic	axis pillar and	follow up of the	ne activities of	this pillar			
4.1.1 Identify who will be the ABX arm pillar focal point in each ministry /authority and assign the TOR					4.1.1 6 months		MOH -PAAF -KEPA -PAFN	4.1.1 Already available	4.1.1 None	4.1.1 available
4.1.2 Prepare clear TOR of the focal person for ABX in each ministry/au thority			4.1.2 TOR	4.1.2	4.1.2 2 months	4.1.2 MOH	4.1.2 AMR committee that reports to secretary	4.1.2 None	4.1.2 None	4.1.2 Yes/No
4.1.3 Assign the job to a defined person/pos ition in each ministry/au thority,			4.1.3 Letter of assignment of duties	4.1.3 1/each ministry	4.1.3 3 months	4.1.3 -MOH -PAAF -KEPA PAFN	4.1.3 AMR Inter- ministerial/ authorities executive committee	4.1.3 None	4.1.3 None	4.1.3 Proportion of ministries/ authorities where ABX arm focal person is assigned

and nomination by each ministry/au thority										
4.1.4 Put TOR And reimburse ment plan for the different tasks.			4.1.4 TOT/ Reimburse ment plan	4.1.4	4.1.4 3 months	4.1.4 MOH	4.1.4 NMCG/ Focal person	4.1.4 None	4.1.4 None	4.1.4 Yes/No
4.1.5 Nominate the stakeholde rs for the task force of the pillar			4.1.5 Decree with names of stakeholde rs	4.1.5 1	4.1.5 4 months	4.1.5 MOH PAAF KEPA PAFN	4.1.5 MOH PAAF KEPA PAFN	4.1.5 None	4.1.5 None	4.1.5 ABX use task force group is formed Yes/No
Strategic Obj	ective 4.2 Ens	ure uninterrup	ted access to I	nigh-quality an	timicrobial me	edicines				
			4.2 Already available		4.2 System is in place	(Pharmace utical and Herbal Medicines Registratio n and Control Administrat ion/Drug Inspection Control				4.2 Already available
4.2.1 Put clear criteria for					4.2.1 System is in place					4.2.1 System is in place

standards of quality safety and efficiency of ABX 4.2.2 Put a system of controlling that all available ABX in the country should meet the internation al standards listed above otherwise they are not allowed to be in the market.			4.2.2 Already available, Only ABX that are licensed by WHO, EMEA or FDA, are allowed in the market			4.2.2 Already available
4.2.3 All ABX should be tested by a system that identifies counterfeit prior to licensing. 4.2.4 Put			4.2.3 Available 4.2.4			4.2.3 System already in place
regulations			Available			System

that prohibit the sale of										already in place
counterfeit										
products										
4.2.5 The										
system that										
controls										
quality										
assurance										
of ABX has the										4.2.5
tne authority					4.2.5					System
to stop					available					already
counterfeit										In place
products										
from being										
licensed										
and										
marketed										
	ective 4.3 Dev	elop and enfo	rce legislation	and regulation	s on prescripti	on and dispen	sing of medici	ne including a	ntibiotics	
4.3.1										
Develop and								4.3.1.1 No		
implement	4.3.1.1		4.3.1.1					need		
national	Finalize the		National				4.3.1.1	Then		
essential	essential		list issued			4.3.1.1	Central	accordingly		4.3.1.1 List
medicine	medicine		and	4.3.1.1	4.3.1.1	MOH	Medical	might need	4.3.1.1 No	is issued
list guided	list that is		endorsed	1	6 months		Stores	new need	need	Yes/No
by the	being prepared at		by MOH director				Directorate	antibiotics that may		
WHO	MOH		general					need a		
Model List	10.011		Belleral					budget		
of Essential										
Medicines			422	422	422	422	4 2 2 5 4 0 1 1	422		422
4.3.2 Mandate			4.3.2	4.3.2	4.3.2	4.3.2 MOH	4.3.2 MOH	4.3.2	4.3.2 None	4.3.2
ivianuate			Mandate	1	6 months	IVIUTI	Undersecre	None		Yes/No

that every institution has an essential medicine list							tary general/ ABX focal person			
4.3.3 Each institution has an essential medicine list and this is checked in accreditati on standards include checking the availability of the essential medicine list	4.3.3.1 Mandate from MOH that once the national essential medicine list is available, all hospitals including public and private hospitals should formulate their own list		4.3.3.1 Mandate	4.3.3.1	4.3.3.1 8 months	4.3.3.1 MOH	4.3.3.1 MOH Undersecre tary general / Abx focal person	4.3.3.1 None	4.3.3.1 None	4.3.3.1 Yes/No
	4.3.3.2 Include the availability of essential medicine list into the license renewal	4.3.3.2.1 Private hospital renewal license checklist should include the availability of essential	4.3.3.2.1 Checklist	4.3.3.2.1	4.3.3.2.1 8 months	4.3.3.2.1 MOH	4.3.3.2.1 Medical Licensing Dep/ MOH	4.3.3.2.1 None	4.3.3.2.1 None	4.3.3.2.1 Licensing checklist includes the item of essential medicine list or not

4.3.4 Re imburseme nt plan is based on essential medicine list in public and private hospitals	conditions of private hospitals	medicine list	4.3.4 After issuing the essential medicine list, Mandate from Undersecre tary General	4.3.4 1	4.3.4 8 months	4.3.4 MOH	4.3.4 MOH/ Undersecre tary General / NMCG	4.3.4 None	4.3.4 None	4.3.4 % of reimbursed ABX that are listed in the essential medicine list
4.4.1 Create a technical group that puts national ABX use guidelines for outpatient care based on Internation al guidelines and National AMR profiles:	4.4.1.1 Put TOR and renewal conditions of the focal group of stakeholde rs	t AMS program	4.4.1.1 TOR	4.4.1.1 1	4.4.1.1 3 months	4.4.1.1 MOH	4.4.1.1 MOH/ National Committee of proper ABX use.	4.4.1.1 None	4.4.1.1 None	4.4.1.1 TOR put Yes or no

can be a subcommit tee from the national task force for the ABX pillar for proper use of ABX									
	4.4.1.2 Assign the members of this committee	4.4.1.2 Committee members	4.4.1.2 1 committee	4.4.1.2 3 months	4.4.1.2 MOH	4.4.1.2 MOH/ National Committee of proper ABX use.	4.4.1.2 None	4.4.1.2 None	4.4.1.2 TOR put Yes or no
	4.4.1.3 Put TOR and official renewal time for the guidelines committee	4.4.1.3 TOR	4.4.1.3	4.4.1.3 3 months	4.4.1.3 MOH	4.4.1.3 MOH/ National Committee of proper ABX use.	4.4.1.3 None	4.4.1.3 None	4.4.1.3 TOR put Yes or no
4.4.2 Guidelines in primary healthcare are available and are regularly updated		4.4.2 Guidelines	4.4.2	4.4.2 6 months	4.4.2 MOH	4.4.2 Primary Healthcare Directorate / technical committee / National Committee for ABX use	4.4.2 None	4.4.2 MOH	4.4.2 Guidelines available Yes or no

	4.4.2.1 Training workshop about ABX guidelines in primary healthcare		4.4.2.1 Workshop	4.4.2.1 Once/year	4.4.2.1 5 years	4.4.2.1 MOH	4.4.2.1 Primary Health care Directorate / technical committee	4.4.2.1 None	4.4.2.1 MOH	4.4.2.1 Proportion of primary healthcare physician who attend every year
/4.4.3 Apply AMS in primary healthcare: Monitoring ABX consumpti on and Link prescriptio n to diagnoses	4.4.3.1 Project ABX consumpti on documenta tion and surveillanc e in primary care	4.4.3.1.1 Put a proposal for this project	4.4.3.1.1 Proposal	4.4.3.1.1	4.4.3.1.1 8 months	4.4.3.1.1 Primary Health care	4.4.3.1.1 AMR committee /Primary Health care Directorate	4.4.3.1.1 COST: No need for extra manpower	4.4.3.1.1 None	4.4.3.1.1 Project is put
	4.4.3.2 Start collecting data and link it to diagnosis		4.4.3.2 Data collection	4.4.3.2 Same number as primary healthcare centers	4.4.3.2 10 months	4.4.3.2 Primary healthcare centers	4.4.3.2 Primary Healthcare Directorate /Informatio n system dep.	4.4.3.2 No	4.4.3.2 No	4.4.3.2 Proportion of primary Healthcare centers that are feeding data into the project
	4.4.3.3 Data analysis and feedback to prescribers		4.4.3.3 Report of analyzed data	4.4.3.3 Once/year	4.4.3.3 1 year	4.4.3.3 Central primary Heath care Directorate	4.4.3.3 Clinical Pharmacist in primary Healthcare.	4.4.3.3 Employ 1 Clinical pharmacist in charge of the project	4.4.3.3 MOH	4.4.3.3 Proportion of primary Healthcare centers

							in primary healthcare		that receive annual report with feedback about ABX prescriptio n
4.4.3.4 Employ more pharmacist s in primary healthcare		4.4.3.4 New employed Pharmacist s	4.4.3.4 According to the needed clinical pharmacist s	4.4.3.4 1 year	4.4.3.4 Primary Health care/ MOH	4.4.3.4 Pharmaceu tical Services Directorate	4.4.3.4 Salaries for new pharmacist s	4.4.3.4 MOH	4.4.3.4 Number of pharmacist s that are employed
4.4.3.5 Improve manpower with professiona Is for the AMS project in primary healthcare	4.4.3.5.1 Train available pharmacist s on AMS and clinical pharmacy principles	4.4.3.5.1 Workshops /Courses	4.4.3.5.1 To be assessed	4.4.3.5.1 18 months	4.4.3.5.1 Primary Health care directorate	4.4.3.5.1 ABX use taskforce group	4.4.3.5.1 30,000 KD/ workshop	4.4.3.5.1 MOH	4.4.3.5.1 Number of pharmacist s in primary Health care who have a certificate in AMS training
	4.4.3.5.2 Employ new clinical pharmacist specialized in AMS and 1 full time clinical microbiolo gist in	4.4.3.5.2 Clinical pharmacist specialized in AMS	4.4.3.5.2 2	4.4.3.5.2 1 8 months	4.4.3.5.2 Primary Healthcare directorate	4.4.3.5.2 Pharmaceu tical Services Directorate	4.4.3.5.2 Salary x2	4.4.3.5.2 MOH	4.4.3.5.2 Clinical pharmacist s are employed, Yes/No

		primary healthcare directorate								
4.4.4 Update the digital system that is available in primary healthcare to integrate the ABX guidelines into the electronic system in the form of pop-ups			4.4.4 ABX guidelines integrated in primary Heath care digital system	4.4.4 1	4.4.4 2 years	4.4.4 Primary Healthcare directorate	4.4.4 MOH/Prim ary Healthcare / technical committee /Informatio n System Dep.	4.4.4 To be determined	4.4.4 MO	4.4.4 updated electronic system available
Strategic Obj	ective 4.5 App	ly AMS In Pub	lic and Private	hospitals	1	I	T .	T .		
4.5.1 Improve and measure appropriat e use of antimicrobi al agents in healthcare	4.5.1.1 Nominate who will be the AMS focal person (Could be the ABX pillar focal person)		4.5.1.1 Decree	4.5.1.1 1	4.5.1.1 3 months	4.5.1.1 MOH	4.5.1.1 MOH Undersecre tary r general /NMCG	4.5.1.1 Already employed in MOH No cost (The Abx focal person)	4.5.1.1 None	4.5.1.1 AMS focal person is nominated
4.5.2 Identify the National	4.5.2.1 AMS technical		4.5.2.1 Group of specialists	4.5.2.1 1	4.5.2.1 3 months	4.5.2.1 MOH	4.5.2.1 MOH/	4.5.2.1 Cost/Activit y/	4.5.2.1 MOH	4.5.2.1 AMS

AMS Task force (Could be a subcommit tee of the ABX pillar Task force) the National AMS Technical team	group is assigned (subgroup from the ABX technical committee. +other specialists like IT, etc)					ABX taskforce group / AMS Focal person	professiona I		Technical Task force members are nominated Yes/No
	4.5.2.2 TOR of the technical groups are well defined They include: -guidelines for inpatient and outpatient careput or endorse diagnostic pathways, treatment and national AMS targets	4.5.2.2 TOR	4.5.2.2	4.5.2.2 3 months	4.5.2.2 MOH	4.5.2.2 ABX Pillar Task force / ABX focal person/ MOH Undersecre tary general.	4.5.2.2 None	4.5.2.2 None	4.5.2.2 TOR put

4.5.3 Put national guidelines either de novo or Endorse existing internation al guidelines with regular updates	4.5.3.1 MOH puts national guidelines for common community acquired infections (CAI)		4.5.3.1 National guidelines for common CAI	4.5.3.1	4.5.3.1 6 months	4.5.3.1 MOH	4.5.3.1 AMS Task force Group	4.5.3.1 none	4.5.3.1 MOH	4.5.3.1 CAI guidelines developed Yes/No
	4.5.3.2 Hospitals put institutiona I guidelines for treatment of Healthcare associated infections (HAI) according to their local epidemiolo gy, with the help of taskforce members	4.5.3.2.1 Mandate for hospitals to issue HAI treatment guidelines based on local epidemiolo gy and with the help of taskforce members	4.5.3.2.1 Mandate	4.5.3.2.1	4.5.3.2.1 6 months	4.5.3.2.1 MOH	4.5.3.2.1 MOH Undersecre tary General/Ta sk Force	4.5.3.2.1 None	4.5.3.2.1 None	4.5.3.2.1 Mandate Yes/No
		4.5.3.2.2	4.5.3.2.2	4.5.3.2.2 1	4.5.3.2.2 8 months	4.5.3.2.2	4.5.3.2.2 Task force	4.5.3.2.2	4.5.3.2.2 Hospitals	4.5.3.2.2

	Organize the work of taskforce members with the different hospitals to assist in putting their own HAI guidelines	Organizatio nal plan and distribution of the hospitals to the different taskforce members			MOH/ Hospitals		To be determined		Proportion of hospitals that have HAI guidelines based on their own epidemiolo gy.
	4.5.3.2.3 HAI treatment guidelines should be reviewed every year	4.5.3.2.3 Revision	4.5.3.2.3	4.5.3.2.3 once / year	4.5.3.2.3 Hospitals/ MOH	4.5.3.2.3 Hospitals/ MOH/ Taskforce	4.5.3.2.3 None	4.5.3.2.3 None	4.5.3.2.3 Proportion of hospitals that have guidelines updated
4.5.3.3 Dissemina on and implement ation of these treatment guidelines in hospita practice.	and endorseme nt of these guidelines with practicing	4.5.3.3.1 Workshops	4.5.3.3.1 Number of hospitals	4.5.3.3.1 1 year	4.5.3.3.1 MOH/Hosp itals	4.5.3.3.1 MOH/ Taskforce/ ABX Focal person	4.5.3.3.1 Number of workshops X 5000 KD	4.5.3.3.1 MOH	4.5.3.3.1 Number of hospitals where guidelines workshops have been performed
	4.5.3.3.2 Distributio n of	4.5.3.3.2 Material	4.5.3.3.2	4.5.3.3.2 1 year	4.5.3.3.2 MOH/Hosp itals	4.5.3.3.2 MOH/Hosp itals	4.5.3.3.2 To be determined	4.5.3.3.2 MOH/ Hospitals	4.5.3.3.2 Proportion of hospitals

	educational material in the form of booklets, brochures, posters, &pocket cards, or equivalent softcopies available on mobile phones		1 for national CAI guidelines 1 for HAI guidelines in each hospital						that have the softcopies available on the mobile phones of staff
	4.5.3.3.3 Post the guidelines on the MOH website and integrate them into the intranet of each hospital	4.5.3.3.3 IT project	4.5.3.3.3 1/Hospital	4.5.3.3.3 1yr, 3 months	4.5.3.3.3 Hospitals	4.5.3.3.3 Hospitals/I nformation System Dep./Hospi tal IT	4.5.3.3.3 To be determined	4.5.3.3.3 Hospitals	4.5.3.3.3 Proportion of hospitals that have the guidelines posted on their intranet
	4.5.3.3.4 Integrate the guidelines into the electronic medical record, in the form of pop-ups	4.5.3.3.4 IT Project	4.5.3.3.4 1/hospital	4.5.3.3.4 2 years	4.5.3.3.4 Hospitals	4.5.3.3.4 Hospitals/I nformation System Dep/ Hospital IT	4.5.3.3.4 To be determined	4.5.3.3.4 Hospitals	4.5.3.3.4 Proportion of hospitals that have guidelines recommen dations integrated in the electronic

4.5.4										medical records system
Make AMS programs mandatory in each hospital (presence of AMS team/ committee)	4.5.4.1 Mandate by MOH for this issue		4.5.4.1 Mandate	4.5.4.1 1	4.5.4.1 1 y 6 months	4.5.4.1 MOH	4.5.4.1 MOH Undersecre tary General/ AMS Task force.	4.5.4.1 None	4.5.4.1 None	4.5.4.1 Mandate issued Yes/No
	4.5.4.2 Build AMS manpower	4.5.4.2.1 Train microbiolo gists, clinical pharmacist s and Infectious Disease specialists on AMS through national workshops	4.5.4.2.1 Workshops	4.5.4.2.1 Depends on the REGIONS	4.5.4.2.1 1 y 6 months	4.5.4.2.1 Hospitals/ Universities / MOH	4.5.4.2.1 Task force	4.5.4.2.1 30,000 KD x number of workshops	4.5.4.2.1 MOH	4.5.4.2.1 Percent microbiolog ists and clinical pharmacist s that have attended at least 1 AMS workshop
Strategic Obj	ective 4.6 Qua	lity control of	antimicrobials	used in anima	als and plants					
4.6 ABX used in the agriculture/ veterinary are under the same control as					4.6 Already available					4.6 Already available

those used in humans Pharmaceu tical and Herbal Medicines Registratio n and Control Administrat ion/The Dug Inspection Control Strategic Obj	ective 4.7 Res	trict the use of	critically impo	ortant ABX to h	numan health i	in the veterina	ry field			
4.7.1 Establish the list of critically important antibiotics for humans in the animal sector.			4.7.1 List	4.7.1 1	4.7.1 3 months	4.7.1 MOH	4.7.1 ABX Task force	4.7.1 None	4.7.1 None	4.7.1 List put Yes/No
4.7.2 Determine from this list which antibiotics should be restricted for use to humans			4.7.2 List	4.7.2 1	4.7.2 3 months	4.7.2 PAAF/MOH	4.7.2 ABX Task force	4.7.2 None	4.7.2 None	4.7.2 List put Yes/No
4.7.3			4.7.3 Decree	4.7.3 1	4.7.3 6 months	4.7.3 PAAF	4.7.3	4.7.3 None	4.7.3 None	4.7.3

Issue a decree that the critically important ABX to human health are not to be used in animals whether in individual treatment or in herd therapy							ABX Task force/ ABX Focal person/ PAAF			Decree issued Yes /No
4.7.4 Prohibit the importatio n of these critically important ABX for vet use	4.7.4.1 Decree that prohibits the importatio n and sale of these ABX in the vet field		4.7.4.1 Decree	4.7.4.1 1	4.7.4.1 6 months	4.7.4.1 PAAF	4.7.4.1 PAAF	4.7.4.1 None	4.7.4.1 None	4.7.4.1 Decree is issued Yes/No
Strategic Obj	jective 4.8 Orga	anize the use o	of antimicrobia	ıl agents in ter	restrial and aq	uatic animals	and agriculture	e.		
4.8.1 Put (or endorse) guidelines regarding the use of ABX in animals			4.8.1 Guidelines	4.8.1	4.8.1 8 months	4.8.1 PAAF/MOH	4.8.1 ABX Task force /PAAF Focal person	4.8.1 Bonus for the members	4.8.1 PAAF/ MOH	4.8.1 Guidelines are established Yes/No

and									
4.8.2 Disseminat e these guidelines among veterinaria ns	4.8.2.1 Booklets/ Brochures/ app that summarize the key points in these guidelines.	4.8.1.1 -Booklets -Brochures - Application	4.8.1.1	4.8.1.1 18 months	4.8.1.1 PAAF	4.8.1.1 PAAF focal person/PA AF	4.8.1.1 To be determined	4.8.1.1 PAAF	4.8.1.1 Proportion of Vet clinics that have easy access to these updated guidelines
	4.8.2.1 Regular workshops for veterinary specialists regarding the application of these guidelines	4.8.2.1 Workshop	4.8.2.1 2*/year	4.8.2.1 1 year	4.8.2.1 PAAF	4.8.2.1 PAAF	4.8.2.1 7000 to 10000 KD per workshop	4.8.2.1 PAAF	4.8.2.1 proportion of veterinary specialists attending these workshops
4.8.3 Restrict selling ABX for animal and agriculture health to exclusively prescriptio ns by vets	4.8.3.1 Decree that vet pharmacies should dispense ABX only according to vet prescriptio ns, and should submit these	4.8.3.1 Decree	4.8.3.1	4.8.3.1 2 years	4.8.3.1 PAAF	4.8.3.1 ABX focal person / PAAF/ MOH	4.8.3.1 None	4.8.3.1 None	4.8.3.1 Decree is issued Yes/No

4.8.4 Control the use of ABX in animals	prescriptio ns with the evidence of selling to PAAF. 4.8.4.1. Detection of the ABX residues in milk, meat, poultry and eggs		4.8.4.1. Already available		4.8.4.1. system in place	PAAF/MOH public Health Lab				4.8.4.1. Already available
	ective 4.9 Cap	acity building a	among vets in	antimicrobial	use and AMR	I	T	T	T	
4.9.1 Provide training courses for vets for ABX use according to guidelines and provide certificate.			4.9.1 Workshop	4.9.1 2/year	4.9.1 2 years	4.9.1 PAAF	4.9.1 PAAF	4.9.1 5 ,000 KD X 2 /YR	4.9.1 PAAF	4.9.1 Proportion of vets that have a yearly certificate
4.9.2 Study baseline consumpti on imported antibiotics in this field	4.9.2.1 Pharmaceu tical Dep in PAAF Committee that will study and document this consumpti on and		4.9.2.1 Report	4.9.2.1 1/year	4.9.2.1 2 years	4.9.2.1 PAAF	4.9.2.1 PAAF	4.9.2.1 None	4.9.2.1 PAAF	4.9.2.1 Report issued Yes/No

compare it					
to					
internation					
al					
benchmark					

N.B. Refer to Annex 5 for Antimicrobial Use Axis Strategic and Monitory plans

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Annexes

Annex 1

A. Governance Strategic Plan

Activity	Sub-activity	Sub-sub- activity	Sub-sub-sub-activity	Milestone
Strategic Objective G1. Nation	nal coordination between focal	points in the different sectors a	nd different pillars	
G1.1 Identify the needed AMR focal points in human health, animal health, welfare, and production, food safety and security, Plants and agriculture, and environment. The TOR should include coordination among the different ministries and authorities along with a yearly meeting of the different focal persons.	G1.1.1 Define the TOR of focal points in each of these sectors			G1. 6 months
	G1.1.2 Employ and assign the functions of a focal person for each pillar and for AMR in general	G1.1.2.1 Assign AMR focal points for each pillar in human health	G1.1.2.1.1 one focal point for each of the following pillars: IPC, Surveillance, ABX, Awareness	
		G1.1.2.2 Assign the general focal Person in MOH	G1.1.2.2.1 Dr. Abeer Aly pending assignment/ New employee	
		G1.1.2.3 Assign 1 focal person in animal health and agriculture fields for all pillars including IPC, Surveillance, ABX, Awareness	G1.1.2.3.1 Representative in NMCG is the focal person or authorities appoint a person	
		G1.1.2.4	G1.1.2.4.1	

		Assign 1 focal person in KEPA for all pillars including IPC, Surveillance, ABX, Awareness	Representative in NMCG is the focal person or authorities appoint a person	
		G1.1.2.5 Assign 1 Focal person in food safety regarding IPC, Surveillance, ABX, Awareness	G1.1.2.5.1 leave the decision to ministries/authorities	
	G1.1.2 Include inter-ministerial communication and coordination with the general AMR focal person in the TOR of the focal points			
Strategic Objective G2. Provide		for the realization of the activiti	ies listed in the plan	
G2.1 Form inter-ministerial NMCG committee	G2.1.1 Nominate the members and have a ministerial decree about their appointment			G2. 3 months
	G2.1.2 Write the TOR of the NMCG that include: to facilitate and oversee implementation, monitoring and evaluation of the AMR action plan			
	G2.1.3 The NMCG has dedicated funds for administrative costs.	G2.1.3.1 NMCG puts a budget for administrative costs		
	an is based on evidence-based	medicine and put by profession	als	
G3.1 Technical groups for specific activities are created	G3.1.1			G3. 3 months

when needed from all	Mapping of professionals			
sectors	throughout the country from			
	public and private sectors			
	that could be part of			
	technical groups when			
	needed.			
Strategic Objective G4. Provid	le guidance and tools to form a	basis for preparation of a nation	nal action plan on AMR	
	G4.1.1			
	Each technical group for			
G4.1 Formulate or endorse	each pillar will agree on			
	existing guidelines or			G4.
guidelines for the different	producing new guidelines or			1 year
topics, along with tools	endorsing international			
	guidelines and make them			
	national			

B. Governance Monitoring Plan

Activity	Sub- activity	Sub-sub- activity	Sub-sub- sub- activity	Indicator	Purpose	Calculation	Frequency	Data source	Method	Baseline
Strategic Ob	jective G1. Nat	tional coordina	tion between	focal points in	the different s	ectors and dif	ferent pillars			
G1.1 Identify the needed AMR focal points in human health, animal health, welfare, and production, food safety and security, Plants and agriculture, and environme nt. The TOR should include coordinatio n among the different ministries	G1.1.1 Define the TOR of focal points in each of these sectors			G1.1.1 TOR are put						

and authorities along with a yearly meeting of the different focal persons.										
	G1.1.2 Employ and assign the functions of a focal person for each pillar and for AMR in general	Assign AMR focal points for each pillar in human health	G1.1.2.1.1 one focal point for each of the following pillars: IPC, Surveillanc e, ABX, Awareness	Focal persons for each pillar at MOH are assigned. Focal points in PAAF, and KEPA, and food safety are assigned	G1.1.2 Oversee the implement ation of different activities in the NAP and ensure communica tion between different authorities	G1.1.2 Yes/No	G1.1.2 Once	G1.1.2 MOH	G1.1.2 Checking	G1.1.2 Partially available
		G1.1.2.2 Assign the general focal person in MOH	G1.1.2.2.1 Dr. Abeer Aly pending assignment / New employee							
		G1.1.2.3 Assign 1 focal person in animal	G1.1.2.3.1 Representa tive in NMCG							

	health and agriculture fields for all pillars including IPC, Surveillanc e, ABX, Awareness	is the focal person or authorities appoint a person				
	G1.1.2.4 Assign 1 focal person in KEPA for all pillars including IPC, Surveillanc e, ABX, Awareness	G1.1.2.4.1 Representa tive in NMCG is the focal person or authorities appoint a person				
	G1.1.2.5 Assign 1 Focal person in food safety IPC: Surveillanc e: ABX Awareness	G1.1.2.5.1 leave the decision to ministries/ authorities				
G1.1.2 Include inter- ministerial			G1.1.2 None			

Strategic Ob	communica tion and coordinatio n with the general AMR focal person in the TOR of the focal points jective G2. Pro	vide legal and	ministerial sup	pport for the re	ealization of th	e activities list	ed in the plan		
G2.1 Form inter- ministerial NMCG committee	G2.1.1 Nominate the members and have a ministerial decree about their appointme nt			G2.1.1 Already available					
	G2.1.2 Write the TOR of the NMCG that include: to facilitate and oversee implement ation, monitoring and evaluation of the AMR action plan			G2.1.2 Already available					

	G2.1.3 The NMCG has dedicated funds for administrat ive costs.	G2.1.3.1 NMCG puts a budget for administrat ive costs		G2.1.3.1 A budget is put and provided		G2.1.3.1 Yes/No	G2.1.3.1 Once	G2.1.3.1 NMCG	G2.1.3.1 Checking	G2.1.3.1 Partially available
Strategic Ob	1	plan is based	on evidence-b	ased medicine	and put by pr	ofessionals		T	ı	
G3.1 Technical groups for specific activities are created when needed from all sectors	G3.1.1 Mapping of professiona Is throughout the country from public and private sectors that could be part of technical groups when needed.			G3.1.1 Technical groups are formed for each pillar, including representat ive of One Health approach	G3.1.1 Ensure proper implement ation and monitoring of the activities of the NAP	G3.1.1 Yes/No	G3.1.1 Once	G3.1.1 NMCG/ MOH Undersecre tary general /Director General in PAAF/KEPA /PAFN	G3.1.1 Checking	G3.1.1 Partially available in MOH
Strategic Obj	jective G4. Pro	vide guidance	and tools to fo	rm a basis for	preparation o	f a national ac	tion plan on A	MR		
G4.1 Formulate or endorse guidelines for the different topics, along with tools	G4.1.1 Each technical group for each pillar will agree on existing guidelines or producing new guidelines			G4.1.1 Guidelines are available when needed	G4.1.1 Standardize practice related to AMR in different sectors	G4.1.1 Yes/No	G4.1.1 Once/year	G4.1.1 Technical groups/ Focal persons of each pillar	G4.1.1 Checking	G4.1.1 Partially available

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Annex 2

A. Awareness Strategic Plan

Activity	Sub-activity	Sub-sub- activity	Sub-sub-sub-activity	Date	Milestone
Strategic Objective 1.1 O	rganize the activities of th	is axis and ensure a prope	r follow up		
1.1.1 To designate a specific employee or post that is responsible of realizing the plan, do the proper communications and follow up on the progress of the plan	1.1.1 Assign one employee/post to overlook these activities in MOH, KEPA, PAAF, food safety			1.1.1 3 months	1.1 8 months
	1.1.2 Put TOR of employment			1.1.2 2 months	
	1.1.3 Letter of employment or assignment of Responsibilities of focal persons in the 4 ministries/authorities			1.1.3 3 months	
Strategic Objective 1.2 Ir		areness of AMR as per One	e health approach through	communication program	s
1.2.1 Establish an evidence-based public communications program targeting the general public in the community	1.2.1.1 Estimate the knowledge, attitude and practice of general public regarding AMR in humans, animals, agriculture and environment			1.2.1.1 Done	1.2 5 years
	1.2.1.2 Conduct national public awareness campaigns based on the One health approach	1.2.1.2.1 Preparation of material regarding the awareness campaign based on the One health approach	1.2.1.2.1.1 Assign a team to formulate and design the public communication	1.2.1.2.1.1 Done	

	material regarding AMR		
	1.2.1.2.1.2 Produce the public communication material regarding AMR	1.2.1.2.1.1 Done	
	1.2.1.2.1.3 Update the communication material to cover all aspects of the One health approach: Animal Health, Agriculture, food safety and environment	1.2.1.2.1.3 6 months	
1.2.1.2.2 Put a yearly plan on broadcasting and posting the public communication material on AMR	1.2.1.2.2.1 Broadcasting is all year long, not only during the AMR awareness week (specific timing, place)	1.2.1.2.2.1 6 months	
	1.2.1.2.2.2 Prepare costing of the plan	1.2.1.2.2.2 8 months	
1.2.1.2.3 Launch the broadcasting plan		1.2.1.2.3 1 year	
1.2.1.2.4 Engage different types of media including social media in AMR awareness such as putting material/ads periodically on ministry/governmental websites, social media		1.2.1.2.4 1 year	

		accounts like Facebook, Twitter, Instagram, etc. 1.2.1.2.5 Involve a public figure like Dr. Abdallah Al-Sanad (spokesperson MOH) in delivering AMR messages, like tweets on twitter or messages in Talk Shows or		1.2.1.2.5 3 months	
		through TV ads/News bar 1.2.1.2.6 Discuss AMR in TV talk shows	1.2.1.2.6.1 MOH sends a letter to Ministry of Information asking to host professionals in AMR to discuss this issue on TV and thus raise public awareness	1.2.1.2.6.1 2 months	
		1.2.1.2.7 Involve Non- Governmental Organizations (NGOs) in AMR awareness campaign	1.2.1.2.7.1 Contact FAO and OIE and ask about their communications program regarding AMR and communicate with them the NAP on AMR	1.2.1.2.7.1 2 months	
Strategic Objective 1.3 In	clude AMR education as p	er One health approach a	t an early age in schools		
1.3.1 Include basic AMR education in school curricula	1.3.1.1 MOH sends a letter with key facts about AMR, antibiotics, hygiene and IPC basics			1.3.1.1 2 months	1.3 2 years

	through the School Health Administration asking to include material related to these topics in school curricula				
	1.3.1.2 Prepare a module/Add material related to basic AMR knowledge, antibiotics, hygiene and IPC in the curricula	1.3.1.2.1 Create a committee of comprising of AMR professional and education professionals to create a module about basic AMR knowledge, antibiotics, hygiene and IPC to be included in school curricula 1.3.1.2.2 Include this		1.3.1.2.1 2 months	
		module in secondary school education		1.3.1.2.2 8 months	
		1.3.1.2.3 Yearly AMR module workshops involving teachers		1.3.1.2.3 1 year	
Strategic Objective 1.4 R	aise awareness of healthca	are workers about AMR ba	sed on the One Health app	proach	
1.4.1 Establish an evidence-based communication program targeting the healthcare providers for human health, animal health, agriculture and environment sector	1.4.1.1 Estimate knowledge, attitudes and practices related to AMR among healthcare workers	1.4.1.1.1 Prepare a survey to estimate this knowledge among healthcare workers and identify the gaps		1.4.1.1.1 Done	1.4 3 years
	1.4.1.2 Conduct awareness program	1.4.1.2.1 MOH mandates to all		1.4.1.2.1 2 months	

within hospitals, primary healthcare, among physicians, nurses, pharmacists, dentists, physiotherapists, and other HC providers	hospitals and primary health care centers to provide mandatory sessions about AMR to all employees who are subject to contract/license renewal		
1.4.1.3 AMR mandatory courses for pharmacists in private practice who are subject to contract/ license renewal	1.4.1.3.1 Letter from MOH to Drug Inspection Control to request evidence of AMR course attendance about AMR upon license renewal	1.4.1.3.1 6 months	
1.4.1.4 Link of AMR course attendance to hospital pharmacists' and primary Health care pharmacists' promotion.	1.4.1.4.1 MOH recommends yearly attendance for promotion	1.4.1.4.1 1 year	
1.4.1.5 Preparation of a bank of presentations, films, articles, etc. diffused to these syndicates and hospitals to be used as material in their AMR communication programs.	1.4.1.5.1 Toolkit for AMR communication program to be used in hospital and syndicates	1.4.1.5.1 1 year	

Strategic Objective 1.5 R approach	1.4.1.6 Syndicates and hospitals send yearly report to MOH about this activity aise awareness of veterina	arians, agriculture professi	onals and environmental v	1.4.1.6 Yearly starting 1 year from starting the plan workers about AMR based	on the One Health
1.5.1 Conduct an awareness program among veterinarians, agriculture professionals, environmental workers, in addition to professionals in PAAF, KEPA, and food safety.	1.5.1.1 Prepare AMR awareness toolkit including slide sets, short films, lectures, posters for these professionals			1.5.1.1 6 months	1.5 3 years
	1.5.1.2 Integrate AMR awareness in continuous education programs of veterinarians, agriculture professionals, environmental workers, in addition to professionals in PAAF, KEPA, and food safety	1.5.1.2.1 The focal person in each ministry/authority checks educational activities in their facilities and Includes AMR awareness as a topic of continuous education		1.5.1.2.1 1 year	
	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	1.5.1.2.2 Mandate that all drug companies' educational activities should include AMR awareness		1.5.1.2.2 6 months	

1.5.2 Country participates the regional/international AMR awareness campaign	1.5.2.1 Participate actively in WHO regional AMR campaign and FAO and OIE activities related to AMR	1.5.2.1.1 Register Kuwait ministries/authorities (KEPA, PAAF, MOH) to participate in the yearly schedule of AMR activities in WHO, FAO, OIE And encourage individual healthcare professionals, veterinarians, agriculture professionals, environmental workers to participate		1.5.2.1.1 1 year	
	1.5.2.2 Participate in GCC activities related to AMR	1.5.2.2.1 Kuwait is part of GCC AMR Group		1.5.2.2.1 Done	
		1.5.2.2.2 Kuwait participates in all the AMR activities GCC		1.5.2.2.2 Done	
		ther education/post-gradu	ate studies in all human ho	ealth specialties, agricultu	re, veterinary food
safety, and environment	1.6.1.1 Include AMR				
1.6.1 Include AMR and related topics based on the One Health approach as a core component of professional education, training, certification	and related topics based on the One Health approach in undergraduate curricula for human health (medical, nursing, dental and pharmaceutical), agriculture, veterinary food safety, and	1.6.1.1.1 preparation of modules for these specialties		1.6.1.1.1 6 months	1.6 1.5 years

	environment related specialties.	1.6.1.1.2			
		Letter from MOH, PAAF, PAFN and KEPA to Ministry of Higher Education to impose on universities/authorities to include these modules in the corresponding education programs.		1.6.1.1.2 6 months	
Strategic Objective Irely	1.6.1.2 AMR and related topics based on the One Health approach included in the licensing procedure for nurses, dentists, physicians, pharmacists, veterinarians and other professionals for which licensing is required.	1.6.1.2.1 Letter from MOH, PAAF, PAFN and KEPA to licensing bodies of these professionals to include AMR and its related topics in licensing exams and interviews or in the evaluation procedure for license renewal		1.6.1.2.1 3 months	
chain and environment		pics in quality assurance a	nd accreditation programs	in numan neaith, animai	neaith, agriculture, food
1.7.1 IPC, AMR surveillance and AMS are included in the accreditation standards of healthcare facilities	1.7.1.1 Letter from MOH to accreditation bodies to include AMR surveillance and implementation of antimicrobial stewardship programs as part of accreditation standards in all			1.7.1.1 3 months	1.7 9 months

	healthcare facilities as well as in the licensing renewal procedures of private healthcare facilities IPC is already part of these standards			
1.7.2 IPC and rational use of antimicrobials is checked in the procedure of licensing farms, animal husbandry, agriculture, aquatic and marine fields	1.7.2.1 Review of large animal/poultry farms licensing policies and procedures, to include IPC and proper ABX use and prevention of ABX use as growth promoters	1.7.2.1.1 Mandate that ABX are not to be used as growth promoters	1.7.2.1.1 3 months	
		1.7.2.1.2 Letter from PAAF Director General to licensing office to include these points in the standards	1.7.2.1.2 3 months	

B. Awareness Monitoring Plan

Activity	Sub- activity	Sub-sub- activity	Sub-sub- sub- activity	Indicator	Purpose	Calculation	Frequency	Data source	Method	Baseline
Strategic Obj	ective 1.1 Org	anize the acti	vities of this ax	is and ensure a	proper follow	/ up				
1.1.1 To designate a specific employee or post that is responsible of realizing the plan, do the proper communica tions and follow up on the progress of the plan	1.1.1 Assign one employee/ post to overlook these activities in MOH, KEPA, PAAF, food safety			1.1.1 Awareness Focal person assigned	1.1.1 Oversee activities of this pillar	1.1.1 Yes/No	1.1.1 Once	1.1.1 MOH KEPA, PAAF, PAFN	1.1.1 Checking	1.1.1 Not available
	1.1.2 Put TOR of employme nt			1.1.2 TOR is put for all 4 focal persons.	1.1.2 Organize work in this pillar	1.1.2 Yes/No	1.1.2 Once	1.1.2 MOH	1.1.2 Checking	1.1.2 Not available
	1.1.3 Letter of employme nt or assignment of Responsibil ities of			1.1.3 Letter of employme nt or assignment of duties to specific focal	1.1.3 Oversee activities in this pillar	1.1.3 Yes/No	1.1.3 Once	1.1.3 MOH, KEPA, PAAF, PAFN	1.1.3 Checking	1.1.3 Not available

Strategic Obj	focal persons in the 4 ministries/ authorities ective 1.2 Incr	ease national	public awarenc	persons in the 4 ministries/ authorities is done	per One health	approach thr	ough commun	ication progra	ms	
1.2.1 Establish an evidence- based public communica tions program targeting the general public in the community	1.2.1.1 Estimate the knowledge, attitude and practice of general public regarding AMR in humans, animals, agriculture and environme nt			1.2.1.1 Done						1.2.1.1 Done
	1.2.1.2 Conduct national public awareness campaigns based on the One health approach	1.2.1.2.1 Preparation of material regarding the awareness campaign based on the One health approach	1.2.1.2.1.1 Assign a team to formulate and design the public communication material regarding AMR	1.2.1.2.1.1 Done						1.2.1.2.1.1 Done

1.2.1.2.1. Produce the public communition material regarding AMR	121211						1.2.1.2.1.1 Done
1.2.1.2.1. Update the communition material acover all aspects of the One health approach Animal Health, Agricultute food safe and environment	1.2.1.2.1.3 The communication material covers all fields of One Health approach (Yes/No)	1.2.1.2.1.3 Highlight the effect of AMR on all fields not just the human health	1.2.1.2.1.3 Yes/No	1.2.1.2.1.3 Once	1.2.1.2.1.3 The awareness technical task force from MOH/PAAF /KEPA/PAF N	1.2.1.2.1.3 Checking	1.2.1.2.1.3 Partially available
1.2.1.2.2 Put a yearly plan on g is all ye long, not only duri posting the public communica tion 1.2.1.2.2 Broadcas g is all ye long, not only duri the AMR awarenes week (specific	1.2.1.2.2.1 Schedule is put	1.2.1.2.2.1 Organize the diffusion of AMR related messages all yearlong	1.2.1.2.2.1 Yes/No	1.2.1.2.2.1 Once	1.2.1.2.2.1 MOH/ Media Office	1.2.1.2.2.1 Checking	1.2.1.2.2.1 Partially available

material o	n timing, place)							
	1.2.1.2.2.2 Prepare costing of the plan	1.2.1.2.2.2 Costing plan is finalized	1.2.1.2.2.2 Organize the diffusion of AMR related messages all yearlong	1.2.1.2.2.2 Yes/No	1.2.1.2.2.2 Once	1.2.1.2.2.2 MOH/ Media Office	1.2.1.2.2.2 Checking	1.2.1.2.2.2 Not available
1.2.1.2.3 Launch th broadcast g plan		1.2.1.2.3 Proportion of the plan that was put on schedule	1.2.1.2.3 Diffusion of AMR messages all year long	1.2.1.2.3 Number of broadcasts related to AMR per month	1.2.1.2.3 Once per year	1.2.1.2.3 MOH	1.2.1.2.3 Counting	1.2.1.2.3 Partially available
1.2.1.2.4 Engage different types of media including social media in AMR awarenes such as putting material/s s periodical on ministry/s vernment websites, social	d y o	1.2.1.2.4 Types of media or social media that mention AMR/mont h	1.2.1.2.4 Relevant informatio n on AMR and related topics reaches to the public as much as possible	1.2.1.2.4 Yes/No	1.2.1.2.4 Once	1.2.1.2.4 MOH/ Media Office	1.2.1.2.4 Checking	1.2.1.2.4 Not available

media accounts like Facebook, Twitter, Instagram, etc.								
1.2.1.2.5 Involve a public figure like Dr. Abdallah Al Sanad (spokesper son MOH) in delivering AMR messages, like tweets on twitter or messages in Talk Shows or through TV ads/News bar		1.2.1.2.5 Public figure (Dr. Abdulla Al- Sanad) sending message regarding AMR	1.2.1.2.5 The public figure influences the public awareness on AMR	1.2.1.2.5 Yes/No	1.2.1.2.5 Once	1.2.1.2.5 MOH/Medi a Office/ Awareness focal person	1.2.1.2.5 Checking	1.2.1.2.5 Partially available
1.2.1.2.6 Discuss AMR in TV talk shows	1.2.1.2.6.1 MOH sends a letter to Ministry of Informatio n asking to host	1.2.1.2.6.1 Proportion of talk shows episodes that discuss AMR	1.2.1.2.6.1 enhance public knowledge about the dangers of AMR	1.2.1.2.6.1 Number of talk show episodes that target AMR/total number of	1.2.1.2.6.1 Once/year	1.2.1.2.6.1 MOH	1.2.1.2.6.1 Data collection	1.2.1.2.6.1 Not available

		1.2.1.2.7 Involve Non- Governmen tal Organizatio ns (NGOs) in AMR awareness campaign	professiona Is in AMR to discuss this issue on TV and thus raise public awareness 1.2.1.2.7.1 Contact FAO and OIE and ask about their communica tions program regarding AMR and communica te with	1.2.1.2.7.1 None		episodes*1 00				
6			them the NAP on AMR							
Strategic Obj	ective 1.3 Incl 1.3.1.1	ude AMR educ	ation as per O	ne health appr	oach at an ear	rly age in schoo	ols			
1.3.1 Include basic AMR education in school curricula	MOH sends a letter with key facts about AMR, antibiotics, hygiene and IPC basics and to School			1.3.1.1 Letter sent Yes or No	1.3.1.1 Improve AMR awareness starting in young age	1.3.1.1 Yes/No	1.3.1.1 Once	1.3.1.1 MOH	1.3.1.1 Checking	1.3.1.1 Not available

Health Administrat ion asking to include material related to these topics in school curricula								
1.3.1.2 Prepare a module/Ad d material related to basic AMR knowledge, antibiotics, hygiene and IPC in the curricula	1.3.1.2.1 Create a committee of comprising of AMR professiona I and education professiona Is to create a module about basic AMR knowledge, antibiotics, hygiene and IPC to be included in school curricula	1.3.1.2.1 Module prepared Yes/No	1.3.1.2.1 Standardize AMR education at the school level	1.3.1.2.1 Yes/No	1.3.1.2.1 Once	1.3.1.2.1 MOH/ Awareness Focal person/ awareness technical task force	1.3.1.2.1 Checking	1.3.1.2.1 Not available
	1.3.1.2.2 Include this module in	1.3.1.2.2 Percent of schools	1.3.1.2.2 Enhance the	1.3.1.2.2 Number of schools	1.3.1.2.2 Once/year	1.3.1.2.2 MOH/ School	1.3.1.2.2 Data collection	1.3.1.2.2 Not available

		secondary school education		that teach the AMR module in their secondary school curricula	education and knowledge about the dangers of AMR starting from a young age	that teach AMR module/To tal number of schools*10 0		Health Administrat ion/ Ministry of Education		
		1.3.1.2.3 Yearly AMR module workshops involving teachers		1.3.1.2.3 Proportion of teachers who attend these workshops	1.3.1.2.3 Standardize AMR teaching in schools	1.3.1.2.3 Number of teachers who attend these modules/T otal number of teachers*1	1.3.1.2.3 Once/year	1.3.1.2.3 MOH/ School Health Administrat ion/Health promotion Departmen t/ Ministry of Education	1.3.1.2.3 Data collection	1.3.1.2.3 ` Not available
1.4.1 Establish an evidence- based communica tion program targeting the healthcare providers for human health, animal	1.4.1.1 Estimate knowledge, attitudes and practices related to AMR among healthcare workers	1.4.1.1.1 Prepare a survey to estimate this knowledge among healthcare workers and identify the gaps	f healthcare w	1.4.1.1.1 Done	AMK based on	tne Une Healt	n approach			1.4.1.1.1 Done

health, agriculture and environme nt sector									
	1.4.1.2 Conduct awareness program within hospitals, primary Health care, among physicians, nurses, pharmacist s dentists, physiother apists, and other HC providers	1.4.1.2.1 MOH mandates to all hospitals and primary Healthcare centers to provide mandatory sessions about AMR to all employees who are subject to contract/lic ense renewal	1.4.1.2.1 Proportion of hospitals and primary Health care centers that provide and request yearly attendance of AMR conference s /sessions	1.4.1.2.1 AMR and related topics are an important elements education and practice	1.4.1.2.1 Number of healthcare facilities that provide and request yearly attendance of AMR conference s/Total number of healthcare facilities*1 00	1.4.1.2.1 Once/year	1.4.1.2.1 MOH	1.4.1.2.1 Data collection	1.4.1.2.1 Not available
	1.4.1.3 AMR mandatory courses for pharmacist s in private practice who are subject to contract/	1.4.1.3.1 Letter from MOH to Drug Inspection Control to request evidence of AMR	1.4.1.3.1 Letter is issued	1.4.1.3.1 Make sure that AMR education/ CE is mandatory for license renewal in the human health sector	1.4.1.3.1 Yes/No	1.4.1.3.1 Once	1.4.1.3.1 MOH	1.4.1.3.1 Checking	1.4.1.3.1 Not available

r	license renewal 1.4.1.4	course attendance about AMR upon license renewal							
	Link of AMR course attendance to hospital pharmacist s' and primary care pharmacist s' promotion.	1.4.1.4.1 MOH recommen ds yearly attendance for promotion	1.4.1.4.1 Mandate is issued	1.4.1.4.1 Emphasize the importance of AMR education and practice in the human health sector	1.4.1.4.1 Yes/No	1.4.1.4.1 Once	1.4.1.4.1 MOH Undersecre tary general / AMR Focal person	1.4.1.4.1 Checking	1.4.1.4.1 Not available
	1.4.1.5 Preparation of a bank of presentatio ns, films, articles, etc. diffused to these syndicates and hospitals to be used as material in their AMR communica	1.4.1.5.1 Toolkit for AMR communica tion program to be used in hospital and syndicates	1.4.1.5.1 Toolkit available	1.4.1.5.1 Standardize AMR CE and knowledge in the human health sector	1.4.1.5.1 Yes/No	1.4.1.5.1 Once	1.4.1.5.1 NMCG/ Awareness Technical task force	1.4.1.5.1 Checking	1.4.1.5.1 Not available

	tion									
	1.4.1.6 Syndicates and hospitals send yearly report to MOH about this activity			1.4.1.6 Proportion of hospitals that send report about AMR Awareness activities to MOH	1.4.1.6 Keep track of AMR activities in hospitals	1.4.1.6 Number of hospitals that send report about AMR Awareness activities to MOH/Total number of hospitals*1 00	1.4.1.6 Once/year	1.4.1.6 MOH	1.4.1.6 Data collection	1.4.1.6 Not available
Strategic Obj approach	ective 1.5 Rais	e awareness o	of veterinarians	, agriculture p	rofessionals a	nd environme	ntal workers a	bout AMR bas	ed on the One	Health
1.5.1 Conduct an awareness program among veterinaria ns, agriculture professionals, environme ntal workers, in addition to professionals in PAAF, KEPA, and food safety.	1.5.1.1 Prepare AMR awareness toolkit including slide sets, short films, lectures, posters for these professiona ls			1.5.1.1 AMR toolkit is available	1.5.1.1 Standardize AMR CE and knowledge in the non- human health sector	1.5.1.1 Yes/No	1.5.1.1 Once	1.5.1.1 Awareness Technical Task force PAAF KEPA PAFN	1.5.1.1 Checking	1.5.1.1 Not available

1.5.1.2 Integrate AMR awareness in continuous education programs of veterinaria ns, agriculture professiona Is, environme ntal workers, in addition to professiona Is in PAAF, KEPA, and food safety	1.5.1.2.1 The focal person in each ministry/au thority checks educational activities in their facilities and Includes AMR awareness as a topic of continuous education	1.5.1.2.1 Number of AMR Awareness sessions per year in each Ministry/au thority	1.5.1.2.1 Enhance AMR knowledge and facing its dangers on the official/gov ernmental level	1.5.1.2.1 Number of sessions	1.5.1.2.1 Once/year	1.5.1.2.1 PAAF KEPA-PAFN	1.5.1.2.1 Counting	1.5.1.2.1 Not available
	1.5.1.2.2 Mandate that all drug companies' educational activities should include AMR awareness	1.5.1.2.2 Proportion of promotion al activities by drug companies in these fields that include messages about AMR	1.5.1.2.2 Engage drug companies in improving AMR awareness among veterinaria ns, agriculture professiona ls	1.5.1.2.2 Number of activities held by drug companies that include messages on AMR/Total number of activities held by	1.5.1.2.2 Once/year	1.5.1.2.2 PAAF Awareness focal person / Director General of PAAF	1.5.1.2.2 Data collection	1.5.1.2.2 Not available

1.5.2 V Country participates the regional/int ernational AMR awareness campaign	1.5.2.2 Km M /a (k P N 1.5.2.1 Participate actively in WHO regional AMR campaign and FAO and OIE en activities in related to AMR Is ve ns ag pr Is en m w participate ns ag pr Is en ns ag pr Is en m w participate ns ag pr Is en m w pr Is Is Is Is Is Is Is Is Is I	preterinaria ns, agriculture professiona ss, environme ntal workers to participate 1.5.2.2.1	1.5.2.1.1 Proportion of events related to AMR in which the country participates	1.5.2.1.1 Engage the country as a whole in the regional and internation al fighting of AMR	1.5.2.1.1 Number of events related to AMR attended by the country/To tal number of events attended by the country*10 0	1.5.2.1.1 Once/year	1.5.2.1.1 MOH PAAF KEPA Awareness Technical Task force	1.5.2.1.1 Data collection	1.5.2.1.1 Not available
P	Participate K	Cuwait is	Done						Done

Strategic Objects safety, and env	vironment re		 1.5.2.2.2 Done education/pos	t-graduate stu	dies in all hum	an health spec	cialties, agricul	lture, veterina	1.5.2.2.2 Done
1.6.1 Include AMR and related topics based on the One Health approach as a core component of professiona I education, training, certificatio n	1.6.1.1 Include AMR and related topics based on the One Health approach in undergradu ate curricula for human health (medical, nursing, dental and pharmaceu tical), food safety, and environme nt related specialties.	1.6.1.1.1 preparatio n of modules for these specialties	1.6.1.1.1 Modules prepared yes/no	1.6.1.1.1 Standardize AMR education and knowledge based on the One Health Approach	1.6.1.1.1 Yes/No	1.6.1.1.1 Once	1.6.1.1.1 Awareness focal persons in all ministries/authorities/Awarenes s technical task force	1.6.1.1.1 Checking	1.6.1.1.1 Not available

	1.6.1.1.2 Letter from MOH, PAFN and KEPA to Ministry of Higher Education to impose on universities /authorities to include these modules in the correspond ing education programs.	1.6.1.1.2 Letter sent Yes/No	1.6.1.1.2 Standardize AMR knowledge based on the One Health Approach	1.6.1.1.2 Yes/No	1.6.1.1.2 Once	1.6.1.1.2 Awareness focal persons in all ministries/ authorities /awareness technical Task force	1.6.1.1.2 Checking	1.6.1.1.2 Not available
1.6.1.2 AMR and related topics based on the One Health approach included in the licensing procedure for nurses, dentists, physicians, pharmacist	1.6.1.2.1 Letter from MOH, PAAF, PAFN and KEPA to licensing bodies of these professiona Is to include AMR and its related topics in licensing exams and	1.6.1.2.1 Letters sent Yes/No	1.6.1.2.1 Standardize AMR knowledge based on the One Health Approach	1.6.1.2.1 Yes/No	1.6.1.2.1 Once	1.6.1.2.1 MOH, PAAF,PAFN and KEPA /AMR focal persons in all authorities	1.6.1.2.1 Checking	1.6.1.2.1 Not available

	s, veterinaria ns and other professiona Is for which licensing is required.	interviews or in the evaluation procedure for license renewal								
		ude AMR and	related topics i	in quality assu	rance and accr	editation prog	grams in huma	n health, anim	al health, agric	culture, food
chain and en		<u> </u>		ı	I	I		ı	1	
	1.7.1.1 Letter from									
	MOH to									
	accreditati									
	on bodies									
	to include			1.7.1.1						
1.7.1 IPC,	AMR			AMR	1.7.1.1					
AMR	surveillanc			surveillanc	Standardize					
surveillanc	e and			e and AMS	quality					
e and AMS	implement ation of			programs	assurance practices					
are	antimicrobi			are	related to					4744
included in	al			included in the	AMR	1.7.1.1	1.7.1.1	1.7.1.1	1.7.1.1	1.7.1.1
the	stewardshi			accreditati	including	Yes/No	Once	мон	Checking	Not
accreditati	p programs			on and	surveillanc					available
on standards	as part of			license	e and					
of	accreditati			renewal	stewardshi					
healthcare	on			standards	p in					
facilities	standards in all			for	healthcare facilities					
	healthcare			healthcare	lacilities					
	facilities as			facilities						
	well as in									
	the									
	licensing									
	renewal									

	procedures of private healthcare facilities IPC is already part of these standards								
1.7.2 IPC and rational use of antimicrobi als is checked in the procedure of licensing farms, animal husbandry, agriculture, aquatic and marine fields	1.7.2.1 Review of large animal/poultry farms licensing policies and procedures, to include IPC and proper ABX use and prevention of ABX use as growth promoters	1.7.2.1.1 Mandate that ABX are not to be used as growth promoters	1.7.2.1.1 Mandate issued	1.7.2.1.1 Put a legal frame for the control of ABX use in the non- human sector	1.7.2.1.1 Yes/No	1.7.2.1.1 Once	1.7.2.1.1 PAAF/ PAAF Director General /PAAF awareness focal person	1.7.2.1.1 Checking	1.7.2.1.1 Not available
		1.7.2.1.2 Letter from PAAF Director General to licensing office to include	1.7.2.1.2 AMR- related topics are included in licensing conditions.	1.7.2.1.2 Emphasize on the importance of AMR and related topics in the non-	1.7.2.1.2 Yes/No	1.7.2.1.2 Once	1.7.2.1.2 PAAF/ Director General of PAAF	1.7.2.1.2 Checking	1.7.2.1.2 Not available

these		human			
points in		sector			
the					
standards					

Annex 3

A. Surveillance Strategic Plan

Strategic interventions	Activity	Sub-activity	Sub-sub- activity	Sub-sub-sub- activity	Date	Milestone for strategic objective
Strategic Objective 2		urveillance system for	r antimicrobial resistance	based on One Health	Approach	
2.1.1 Establish a national coordination structure for AMR surveillance in humans, veterinary agriculture and environment	2.1.1.1 Write and approve terms of reference for a national coordinating committee for AMR surveillance with the mandate to oversee the AMR surveillance program, including AMR surveillance in Humans, Veterinary, agriculture, environment and Food safety.				2.1.1 6 months	2.1 3 years
	2.1.1.2. Write and approve terms of reference for a central office that executes the National surveillance in humans and receives data from other AMR offices in PAAF, KEPA and food safety	2.1.1.2.1 Write terms of reference of the AMR office at MOH			2.1.1.2.1 3 months	

t d	2.1.1.2.2 Create the AMR coordinating office at MOH.	2.1.1.2.2.1 Include the already existing Glass team into the MOH National AMR office for Human sector at MOH.	2.1.1.2.2.1.1 Add 2 employees with a support from IT at MOH to be able to handle nationwide data (to do the functions of the central office).	2.1.1.2.2.1.1 6 months	
		2.1.1.2.2.2 Mandate that each MOH hospital has an AMR surveillance focal person that coordinates with Central AMR surveillance office.		2.1.1.2.2.2 3 months	
	2.1.1.2.3 Inform PAAF, PAFN and KEPA about the need for AMR surveillance in their respective fields.			2.1.1.2.3 1 month	
1	2.1.1.2.4 Nominate AMR focal persons in PAAF, KEPA, and Food safety			2.1.1.2.4 3 months	
	2.1.1.2.5 Write the job description of these focal persons. The job description would be to collect AMR data and supply it			2.1.1.2.5 3 months	

	to the Central AMR office and full coordination with the Central AMR office at MOH.			
2.1.1.3 Identify priority organisms for the purpose of national AMR surveillance in human sector, animal sector, plants and Environment.			2.1.1.3 3 months	
2.1.1.4 Systematically collect, analyze and report data on AMR in humans in order to inform decision-making at national and international levels (Choose sentinel group of labs)	2.1.1.4.1 Choose a list of labs (dealing with specimen from humans) that can provide reliable AMR data will form a sentinel for starting AMR data collection that will be submitted to AMR central office.		2.1.1.4.1 3 months	
	2.1.1.4.2 Increase the number of labs that are included in the list on yearly basis until it becomes representative of the national data.	2.1.1.4.2.1 Establish close communication between central AMR OFFICE that also collects data for glass with the lab council to have a list of labs that are accredited by the council, and to	2.1.1.4.2.1 1 year	

			include them in the list of labs eligible to submit data to GLASS and National AMR surveillance		
	2.1.1.5 Establish an antimicrobial resistance surveillance network for human health	2.1.1.5.1 integrate all MOH laboratories in network with the national reference Lab.	2.1.1.5.1.1 Training of hospital AMR Surveillance focal persons for data cleaning and data entry into the MOH surveillance system	2.1.1.5.1.1 1 year	
		2.1.1.5.2 Generate yearly report about AMR in humans in general and mainly priority organisms in humans by the MOH central office.		2.1.1.5.2 2 years	
2.1.2 Participate in Global Antimicrobial Resistance Surveillance System (GLASS- AMR)	2.1.2.1 Nominate a hospital ad a team for GLASS reporting with TOR			2.1.2.1 Already done	
	2.1.2.2 Give the GLASS team the authority to request and collect data from labs nationwide			2.1.2.2 Already done	
		2.1.2.2.1 The team initiates data		2.1.2.2.1 Started 2021	

	2.1.2.3 Send yearly report to glass with increasing number of contributing labs	collection from a sentinel of labs 2.1.2.3.1 Central GLASS focal group (Part of AMR central office), chooses from the list of eligible accredited labs submitted by central entity of medical labs to AMR office,	2.1.2.3.1 1 Yearly training of GLASS focal persons on GLASS data collection, management and entry	2.1.2.3.1.1.1 Central workshops yearly	2.1.2.3.1.1.1 2 years	
Stratogic Objective 2	2. Ruild laboratory o	2.1.2.3.2 Yearly lab report is sent to GLASS with increasing number of contributing labs every year.	h-quality microbiologica	data for nations man	2.1.2.3.2 Started already	survaillance
activities in human,	Bullu laboratory C	apacity to produce hig	ni-quality inicrobiological	i uata ioi patient man	agement and support	Sui veillance
2.2.1 Lab in Hospitals become CAP certified (College of American pathologists)	2.2.1.1 Pilot with 2 labs				2.2.1.1 1 year	2.2 5 years
	2.2.1.2 Increase the number of labs that are CAP members	2.2.1.2.1 Lab council puts a plan for sequential recruitment of labs into the CAP			2.2.1.2.1 1 year	
		2.2.1.2.2 Sequential recruitment of labs into CAP			2.2.1.2.2 5 years	

2.2.2 Improve private labs capacity to be eligible for submitting data to GLASS and National AMR	2.2.2.1 Reinforce the accreditation activity of the lab council that audits labs according to availability of qualified microbiologist, results of External QC and abidance to guidelines	2.2.2.1.1 Mandate from MOH that each private lab should pass the accreditation by the Lab council		2.2.2.1.1 12-24 months	
		2.2.2.1.2 Accreditation of the lab council includes the availability of qualified microbiologist, results of External QC and abidance to guidelines		2.2.2.1.2 6 months	
Strategic objective 2	.3: Collect data from V	et, Agriculture and En	vironment		
2.3.1 Collect AMR data from Vets	2.3.1.1 Map available data On AMR in vet and agriculture in the country.	2.3.1.1.1 Agriculture and Environment focal persons collect available info about AMR in the country and issue a report	2.3.1.1.1.1 Map PAAF labs and the type of AMR data that is generated.	2.3.1.1.1.1 6 months	2.3 5 years
	2.3.1.2	2.3.1.2.1	2.3.1.1.1.2 Collect available data about AMR in PAAF/PAFN	2.3.1.1.1.2 6 months	
	2.3.1.2	2.3.1.2.1	2.3.1.2.1.1	2.3.1.2.1.1	

Improve vet lab capacity in a way they generate data about AMR vets and agriculture	Establish quality control of Vet Microbiology labs	Assign expert to establish a system for quality control and put a plan	1 year	
		2.3.1.2.1.2 Execute the plan of quality control in Vet labs	2.3.1.2.1.2 2 year	
	2.3.1.2.2 Extend Vet Micro Labs to identify priority organisms antibiogram	2.3.1.2.2 1 Increase man power	2.3.1.2.2 1 6 months	
		2.3.1.2.2.2 Road map and full project	2.3.1.2.2.2 1 year	
		2.3.1.2.2.3 Workshops for training about antibiogram	2.3.1.2.2.3 1 year	
	2.3.1.2.3 Implement national surveillance for antibiogram priority animal pathogens, zoonotic and commensal bacterial isolates.	2.3.1.2.3.1 Choose a number of Vet labs that could submit AMR data to PAAF for AMR surveillance	2.3.1.2.3.1 1 year	
		2.3.1.2.3.2 Collect data from a Sentinel of vet hospitals or labs that submit data to the	2.3.1.2.3.2 2 years	

		2.3.1.2.4 The focal AMR entity at PAAF submits AMR data to the Central AMR surveillance lab.	department of surveillance. (They belong to the list).	2.3.1.2.4 1 year	
2.3.2 Collect AMR data from Environment.	2.3.2.1 Map available data from KEPA	2.3.2.1 Check KEPA Labs and publications about any AMR data in environment		2.3.2.1 3 months	
	2.3.2.2 Initiate AMR data collection from Environment By research projects.	2.3.2.2.1 The director of the office of research and studies puts AMR surveillance as one of the priority subjects in research	2.3.2.2.1.1 The plan of prioritizing AMR surveillance in Environment and marine environment re and is being discussed with universities/institutes.	2.3.2.2.1.1 6 months	
			2.3.2.2.1.2 Call for proposal from universities/institutes	2.3.2.2.1.2 1 year	
			2.3.2.2.1.3 The director of the office of research and studies applies for funds from KEPA for projects related to AMR surveillance in Vet, agriculture, and Environment.	2.3.2.2.1.3 6 months	

2.3.2.2.2 Increase lab capacity of KEPA Lab that deals with the quality of coastal water to include AMR identification and surveillance in water and sediment	2.3.2.2.1 Letter from KEPA Director General to Technical affairs sector of micro labs asking to prepare a project with a budget that aims at expanding their work from only identification to antibiogram testing		2.3.2.2.2.1 3 months	
	2.3.2.2.2 KEPA analytical central lab KEPA prepare road map for the labs to be able to do antibiogram testing	2.3.2.2.2.1 Identify the needed machinery and their budget	2.3.2.2.2.1 6 months	
		2.3.2.2.2.2 Put a plan of needed extra staffing	2.3.2.2.2.2 6 months	
	2.3.2.2.3 KEPA provides the related budget	2.3.2.2.3 Communication between KEPA and ministry of finance	2.3.2.2.2.3 2 years	
	2.3.2.2.4 KEPA microbiology labs start generating antibiogram results for priority organisms		2.3.2.2.2.4 3 years	
2.3.2.2.3 Generate report about AMR			2.3.2.2.3 4 years	

		surveillance in			
		Environment			
2.3.3 The national AMR Surveillance Office at MOH gathers data from KEPA, PAAF and PAFN compiles it with Human AMR data and generates yearly ONE Health AMR surveillance report.	2.3.3.1 KEPA, PAAF and PAFN submit available data to Central AMR surveillance office at MOH on yearly basis	2.3.3.1.1 Create a digital network for data sharing with MOH		2.3.3.1.1 5 years	
		2.3.3.1.2 Add this function to the TOR of the focal person at PAAF and KEPA and food safety.		2.3.3.1.2 3 years	
	.4: Create a National F	Reference Lab for AMR			
2.4.1 To have A Kuwaiti Center of Disease Control that will be overseeing bacterial ,viral and AMR reference labs	2.4.1.1 Project of reference lab with its budget and having it endorsed by MOH			2.4.1.1 Already available	2.4 5 years
	2.4.1.2 Kuwait CDC becomes functional			2.4.1.2 5 years	
	2.4.1.3 Write and approve terms of reference for a national			2.4.1.3 3 months	

	A N A D f				
	AMR reference				
	laboratory with				
	expertise in				
	method				
	s for confirming				
	and characterizing				
	specific pathogens,				
	putting				
	guidelines, SOP for				
	identification and				
	antibiogram testing				
	of priority				
	organisms for				
	surveillance				
	training according				
	to guidelines, and				
	organizing quality				
	assurance schemes				
	2.4.1.4				
	Prepare a full				
	project and			2.4.1.4	
	proposal for the			5 years	
	reference lab				
	2.4.1.5				
	Prepare a budget				
	for the building,			2.4.1.5	
	equipment and			5 years	
	man power of the			5 , 5 4 5	
	reference Lab				
	2.4.1.6				
	Execution of the			2.4.1.6	
	Reference Lab			5 years	
2.4.2	2.4.2.1	2.4.2.1.1			
Delegate the	Identify institution	A letter from MOH			
function of	affiliated to MOH	to all labs that asks		2.4.2.1.1	
				Already available	
surveillance of the	that will host the	them to coordinate		•	

Reference Lab pending its readiness	manpower and activities of the AMR Surveillance office for the actual work on the National and GLASS AMR reports pending the finalization of the reference lab.	with the GLASS lab by delivering AMR data for the purpose of AMR Surveillance				
2.5.1 Lab audit by lab accreditation body according to microbiologist consultant in charge, external quality control and use of CLSI guidelines	2.5.1.1 Accreditation of the medical Laboratories by Medical Laboratories Council	2.5.1.1.1 Involve all laboratories in external quality assurance programs	2.5.1.1.1.1 Mandate that all labs including all private should be accredited and all perform external QC.	2.5.1.1.1.1 List of accreditation bodies accepted in Kuwait, all labs should be accredited by one of them	2.5.1.1.1.1.1 6 months	2.5 5 years

B. Surveillance Monitoring Plan

Strategic interventions	Activity	Sub- activity	Sub-sub- activity	Sub-sub- sub- activity	Indicator	Purpose	Calculatio n	Frequenc y	Data source	Method	Baseline
Strategic O	bjective 2.1:	Set up a natio	onal surveilla	nce system fo	r antimicrob	ial resistance	based on On	e Health App	roach		
2.1.1 Establish a national coordinati on structure for AMR surveillan ce in humans, veterinary agricultur e and environm ent	2.1.1.1 Write and approve terms of reference for a national coordinating committe e for AMR surveillance with the mandate to oversee the AMR surveillance program, including AMR surveillance in Humans, Veterinary, agriculture, environm				2.1.1 Committe e is formed yes/No	2.1.1 To oversee and coordinat e surveillan ce activities in Humans, Vets, Agricultur e and Environm ent.	2.1.1 Yes/No	2.1.1 Once	2.1.1 MOH	2.1.1 Checking	2.1.1 Not formed

ent and Food safety. 2.1.1.2. Write and approve terms of reference for a central office that executes the National surveillan ce in humans and receives data from other AMR offices in PAAF, KEPA and	2.1.1.2.1 Write terms of reference of the AMR office at MOH			2.1.1.2.1 TOR written	2.1.1.2.1 To organize and frame the work of this committe e	2.1.1.2.1 Yes/No	2.1.1.2.1 Once	2.1.1.2.1 MOH	2.1.1.2.1 Checking	2.1.1.2.1 Not written yet
KEPA and food safety										
	2.1.1.2.2 Create the AMR coordinati ng office at MOH.	2.1.1.2.2.1 Include the already existing Glass team into the MOH National	2.1.1.2.2.1 .1 Add 2 employee s with a support from IT at MOH to be able to	2.1.1.2.2.1 .1 2 extra employee s are employed. Yes/No	2.1.1.2.2.1 .1 To execute all the Human AMR surveillan ce and to	2.1.1.2.2.1 .1 Office with employed personnel is formed.	2.1.1.2.2.1 .1 Once	2.1.1.2.2.1 .1 MOH/ Informati on system administr ation	2.1.1.2.2.1 .1 Checking	2.1.1.2.2. 1.1 Not available

	AMR office for Human sector at MOH.	handle nationwid e data (to do the functions of the central office).		collect data from the Vet and Agricultur e and KEPA Surveillan ce.					
	2.1.1.2.2.2 Mandate that each MOH hospital has an AMR surveillan ce focal person that coordinat es with Central AMR surveillan ce office.		2.1.1.2.2.2 Mandate issued Yes/No	2.1.1.2.2.2 Mandate is issued	2.1.1.2.2.2 Yes/No	2.1.1.2.2.2 Once	2.1.1.2.2.2 MOH	2.1.1.2.2.2 Checking	2.1.1.2.2. 2 Not available
2.1.1.2.3 Inform PAAF,PAF N and KEPA about the need for AMR surveillan ce in their			2.1.1.2.3 Letter sent						

respective fields.								
2.1.1.2.4 Nominate AMR focal persons in PAAF, KEPA, and Food safety		2.1.1.2.4 AMR surveillan ce focal persons appointed in all 3 authoritie s	2.1.1.2.4 To start the activities that lead to data collection and ultimately do the surveillan ce data collection in their respective authoritie s	2.1.1.2.4 Yes/No	2.1.1.2.4 ONCE in each authority	2.1.1.2.4 MOH PAAF KEPA PAFN	2.1.1.2.4 Checking	2.1.1.2.4 Only GLASS focal person is employed at MOH. No national surveillan ce job descriptio n. None in PAAF, KEPA, PAFN
2.1.1.2.5 Write the job descriptio n of these focal persons. The job descriptio n would be to collect AMR data and supply it to the Central		2.1.1.2.5 TOR are set	2.1.1.2.5 To organize the job and reach the goal of surveillan ce	2.1.1.2.5 Yes/No	2.1.1.2.5 Once in each authority	2.1.1.2.5 MOH PAAF KEPA PAFN	2.1.1.2.5 Checking	2.1.1.2.5 Not available

	AMR office and full coordinati on with the Central AMR office at MOH.								
2.1.1.3 Identify priority organisms for the purpose of national AMR surveillan ce in human sector, animal sector, plants and Environm ent.			2.1.1.3 List is available and includes all listed sectors.	2.1.1.3 To focus surveillan ce on the most important AMR trends that are of interest in all sectors	2.1.1.3 List is put Yes/No	2.1.1.3 Once	2.1.1.3 MOH	2.1.1.3 Checking	Not available.
2.1.1.4 Systematically collect, analyze and report data on	2.1.1.4.1 Choose a list of labs (dealing with specimen from humans)		2.1.1.4.1 List is chosen(8 Labs submitted GLASS data for 2021	2.1.1.4.1 To include only data that is microbiol ogically brought according	2.1.1.4.1 Yes/No	2.1.1.4.1 Once/year	2.1.1.4.1 MOH	2.1.1.4.1 Checking	2.1.1.4.1 List for GLASS data has been put for 2022

AMR in humans in order to inform decision-making at national and internatio nal levels (Choose sentinel group of labs)	that can provide reliable AMR data will form a sentinel for starting AMR data collection that will be submitted to AMR central office.	-Amiri -Adan -Mubarak Alkhabeer -Jaber -Farwania -Jahra -Chest -Ibn Sina)	to recogniza ble microbiol ogy guidelines like CLSI or EUCAST					
	2.1.1.4.2.1 2.1.1.4.2 Increase the number of labs that are included in the list on yearly basis until it becomes represent ative of the national data. 2.1.1.4.2.1 Establish close communic ation between central AMR OFFICE that also collects data for glass with the lab council to have a list of labs that are accredited by the	2.1.1.4.2.1 Proportio n of labs in the country that are included in the list	2.1.1.4.2.1 To have surveillan ce data that is represent ative of all the country	2.1.1.4.2.1 Number of labs in the list/total number of labs	2.1.1.4.2.1 Once/year	2.1.1.4.2.1 MOH/ Surveillan ce office	2.1.1.4.2.1 Checking	2.1.1.4.2. 1 10 labs will be included in 2022 in GLASS data

		council, and to include them in the list of labs eligible to submit data to GLASS and National AMR surveillan ce							
2.1.1. Estab an antim bial resist surve ce netwood for huma healt	olish integrate all MOH laboratori es in network with the national reference Lab.	2.1.1.5.1.1 Training of hospital AMR Surveillan ce focal persons for data cleaning and data entry into the MOH surveillan ce system	2.1.1.5.1.1 Yearly workshop performe d Yes/No	2.1.1.5.1.1 To provide adequate training about data managem ent in a unanimou s way throughou t all the included labs	2.1.1.5.1.1 Workshop done Yes /No	2.1.1.5.1.1 Once/year	2.1.1.5.1.1 MOH	2.1.1.5.1.1 Checking	2.1.1.5.1. 1 Only GLASS focal is being trained
	2.1.1.5.2 Generate yearly report about AMR in humans in		2.1.1.5.2 Yearly human AMR report is issued	2.1.1.5.2 To direct empiric therapy and strategic plan in	2.1.1.5.2 Yes/No	2.1.1.5.2 Once/Year	2.1.1.5.2 MOH Surveillan ce office	2.1.1.5.2 Checking	2.1.1.5.2 GLASS report started 2021

		general and mainly priority organisms in humans by the MOH central office.			treatment and preventio n and antibiotic use in humans.			
2.1.2 Participat e in Global Antimicro bial Resistance Surveillan ce System (GLASS- AMR)	2.1.2.1 Nominate a hospital ad a team for GLASS reporting with TOR			2.1.2.1 Already done				Already available.
,	2.1.2.2 Give the GLASS team the authority to request and collect data from labs nationwid e			2.1.2.2 Already done				2.1.2.2 Already done
		2.1.2.2.1 The team initiates		2.1.2.2.1 Started 2021				2.1.2.2.1 Started 2021

	data collection from a sentinel of labs									
2.1.2.3 Send yearly report to glass with increasing number of contributi ng labs	2.1.2.3.1 Central GLASS focal group (Part of AMR central office), chooses from the list of	2.1.2.3.1 1 Yearly training of GLASS focal persons on GLASS data collection, managem ent and entry	2.1.2.3.1.1 .1 Central workshop s yearly	2.1.2.3.1.1 .1 Workshop done, yes/No have 10- 20 % increase in number of participati ng labs in GLASS annually	2.1.2.3.1.1 .1 To do proper data entry into GLASS	2.1.2.3.1.1 .1 Yes/No	2.1.2.3.1.1 .1 Once/Year (and for new employee)	2.1.2.3.1.1 .1 Surveillan ce/GLASS office at MOH	2.1.2.3.1.1 .1 Checking	2.1.2.3.1. 1.1 Not being done on yearly basis systematically
	2.1.2.3.2 Yearly lab report is sent to GLASS with increasing number of contributi			2.1.2.3.2 Data is available in the yearly GLASS report	2.1.2.3.2 To be included in the global AMR map and to evaluate benchmar	2.1.2.3.2 Yes/No	2.1.2.3.2 Once/Year	2.1.2.3.2 GLASS Report	2.1.2.3.2 Checking	2.1.2.3.2 Started 2021

		ng labs every year.				k with other countries					
Strategic Ol activities in	=	Build laborate	ory capacity t	o produce hig	gh-quality mi	crobiological	data for pation	ent managem	ent and supp	ort surveillar	ice
		2.2.1 Lab in Hospitals become CAP certified (College of American pathologis ts)	2.2.1.1 Pilot with 2 labs		2.2.1.1 Started already	2.2.1.1 To check feasibility					2.2.1.1 Started already
			2.2.1.2 Increase the number of labs that are CAP members	2.2.1.2.1 Lab council puts a plan for sequential recruitme nt of labs into the CAP	2.2.1.2.1 List is put already						2.2.1.2.1 List is put already
				2.2.1.2.2 Sequential recruitme nt of labs into CAP	2.2.1.1 Proportio n of Labs that are CAP affiliated	2.2.1.1 To ensure high standard lab work and	2.2.1.1 Number of labs that are CAP Members/ Total labs	2.2.1.1 Once/Year	2.2.1.1 MOH Central laboratori es	2.2.1.1 Checking	2.2.1.1 2 started already as pilot

					get internatio nal scientific support					
	2.2.2 Improve private labs capacity to be eligible for submittin g data to GLASS and National AMR	2.2.2.1 Reinforce the accreditati on activity of the lab council that audits labs according to availabilit y of qualified microbiol ogist, results of External QC and abidance to guidelines	2.2.2.1.1 Mandate from MOH that each private lab should pass the accreditati on by the Lab council	2.2.2.1 Proportio n of labs that are functional and that have passed the accreditati on testing	2.2.2.1 To have quality AMR data from all the labs	2.2.2.1 Number of of labs that are functional and that have passed the accreditati on testing/to tal number of labs	2.2.2.1 Once/Year	2.2.2.1 Central Medical laboratory council/ Medical Licensing Dep.	2.2.2.1 Checking	2.2.2.1 All governme ntal labs are accredite d Not all private labs are accredite d and many do not pass the accreditat ion check yearly.
			2.2.2.1.2 Accreditat ion of the lab includes the availabilit y of	2.2.2.1.2 Updated accreditati on standards of Labs prepared or not	To improve private lab capacity for detecting AMR	Update done yes/No	once	Central Medical laboratory council	checking	available in MOH Labs

				qualified microbiol ogist, results of External QC and abidance to guidelines							
Strategic ob	jective 2.3: C		om Vet, Agric	ulture and Er	ivironment						
2.3.1 Collect AMR data from Vets	2.3.1.1 Map available data On AMR in vet and agricultur e in the country.	2.3.1.1.1 Agricultur e and Environm ent focal persons collect available info about AMR in the country and issue a report	2.3.1.1.1.1 Map PAAF labs and the type of AMR data that is generated		2.3.1.1.1.1 Report is Submitted to central AMR surveillan ce office	2.3.1.1.1.1 To check the available Situation regarding AMR identificat ion and reporting in Vet labs.	2.3.1.1.1.1 Report done Yes/No	2.3.1.1.1.1 Once	2.3.1.1.1.1 Vet Labs PAAF	2.3.1.1.1.1 Data collection according to a checklist	2.3.1.1.1 1 Unknown
		3.00010	2.3.1.1.1.2 Collect available data about AMR in PAAF/PAF N		2.3.1.1.1.2 AMR Vets/food reports are submitted to central AMR Surveillan ce office	2.3.1.1.1.2 To collect available AMR data	2.3.1.1.1.2 AMR report from PAAF/PAF N	2.3.1.1.1.2 Once/year	2.3.1.1.1.2 Vet Labs/PAF N	2.3.1.1.1.2 Data collection	2.3.1.1.1. 2 Not available
	2.3.1.2	2.3.1.2.1	2.3.1.2.1.1		2.3.1.1.1.1	2.3.1.1.1.1	2.3.1.1.1.1 Yes/No	2.3.1.1.1.1 Once	2.3.1.1.1.1 PAAF	2.3.1.1.1.1 Checking	2.3.1.1.1. 1

Improve vet lab capacity in a way they generate data about AMR vets and agricultur e	Establish quality control of Vet Microbiol ogy labs	Assign expert to establish a system for quality control and put a plan	Profession al is assigned	To put a plan with a defined budget for Vet labs to start generatin g adequate AMR data					The Vet labs generate only microorga nism identificat ion data, no AMR data is generated
		2.3.1.2.1.2 Execute the plan of quality control in Vet labs	2.3.1.2.1.2 Proportio n of VET Labs that Participat e in external quality control activities	2.3.1.2.1.2 To provide periodic check up on the accuracy of AMR detection in vet labs.	2.3.1.2.1.2 Number of of VET Labs that participat e in external quality control activities/ Total number of vet labs.	2.3.1.2.1.2 once/year	2.3.1.2.1.2 Vet labs/PAAF	2.3.1.2.1.2 Checking	2.3.1.2.1. 2 Not available
	2.3.1.2.2 Extend Vet Micro Labs to identify priority organisms antibiogra m	2.3.1.2.2 1 Increase man power	2.3.1.2.2 1 None						

	2.3.1.2.2.2 Road map and full project	2.3.1.2.2.2 The project is put with specified budget and ready for execution	2.3.1.2.2.2 To upgrade Vet labs to become able to generate AMR data	2.3.1.2.2.2 Yes/No	2.3.1.2.2.2 Once	2.3.1.2.2.2 PAAF	2.3.1.2.2.2 Checking	2.3.1.2.2. 2 Not available
	2.3.1.2.2.3 Workshop s for training about antibiogra m	2.3.1.2.2.3 Workshop is performe d.	2.3.1.2.2.3 To train microbiol ogists and technician in Vet labs about AMR IDENTIFIC ATION	2.3.1.2.2.3 Proportio n of vet labs that are represent ed in these workshop s	2.3.1.2.2.3 Once/year	2.3.1.2.2.3 PAAF	2.3.1.2.2.3 Checking	2.3.1.2.2. 3 Not available
2.3.1.2.3 Implement national surveillance for antibiogram priority animal pathogens, zoonotic and commens al bacterial isolates.	2.3.1.2.3.1 Choose a number of Vet labs that could submit AMR data to PAAF for AMR surveillan ce	2.3.1.2.3.1 List of labs is available yearly	2.3.1.2.3.1 To collect Microbiol ogically sound AMR data from Vet Labs.	2.3.1.2.3.1 Yes/No	2.3.1.2.3.1 Once/Year	2.3.1.2.3.1 PAAF	2.3.1.2.3.1 Checking	2.3.1.2.3. 1 Not available

			2.3.1.2.3.2 Collect data from a Sentinel of vet hospitals or labs that submit data to the departme nt of surveillan ce. (They belong to the list).	2.3.1.2.3.2 Data is yearly collected	2.3.1.2.3.2 To generate AMR data from Vets and Agricultur e and correlate it with Human AMR and decide upon ABX use in agricultur e and vets.	2.3.1.2.3.2 Yes/No	2.3.1.2.3.2 Once/Year	2.3.1.2.3.2 PAAF Vets Lab	2.3.1.2.3.2 Checking	2.3.1.2.3. 2 Not available.
2.3.2	2.3.2.1	2.3.1.2.4 The focal AMR entity at PAAF submits AMR data to the Central AMR surveillan ce lab		2.3.1.2.4 Data submitted to MOH Central Office.	2.3.1.2.4 To pool data into the central national One Health AMR office, to generate decisions related to ABX in all sectors of One Health. 2.3.2.1	2.3.1.2.4 Yes/No	2.3.1.2.4 Once/Year	2.3.1.2.4 MOH Central AMR One Health office	2.3.1.2.4 Checking	2.3.1.2.4 Not available

Collect AMR data from Environm ent.	Map available data from KEPA	Check KEPA Labs and publicatio ns about any AMR data in environm ent		Report is issued	To collect available baseline informatio n about AMR in the environm ent.	Yes/No	Once	Publicatio ns from Kuwait	Checking	Not available.
	2.3.2.2 Initiate AMR data collection from Environm ent By research projects.	2.3.2.2.1 The director of the office of research and studies puts AMR surveillan ce as one of the priority subjects in research	2.3.2.2.1.1 The plan of prioritizin g AMR surveillan ce in Environm ent and marine environm ent re and is being discussed with universitie s/institute s.	2.3.2.2.1.1 % of research meetings with universitie s/institute s where AMR is requested to be apriority in research projects.	2.3.2.2.1.1 To prioritize AMR in environm ent as research target, and attract related projects and funds	2.3.2.2.1.1 Number of research meetings with universitie s where AMR is requested to be a priority in research projects/ total number of research meetings.	2.3.2.2.1.1 Twice /year	2.3.2.2.1.1 KEPA/ Universiti es/KISR	2.3.2.2.1.1 Checking	2.3.2.2.1. 1 Not available
			2.3.2.2.1.2 Call for proposal from universitie s/institute s	2.3.2.2.1.2 Proportio n of universitie s that send projects related to AMR in	2.3.2.2.1.2 To increase the number of research projects about	2.3.2.2.1.2 Number of universitie s that send projects related to AMR in	2.3.2.2.1.2 Twice/yea r	2.3.2.2.1.2 KEPA Research office	2.3.2.2.1.2 Checking	2.3.2.2.1. 2 Not available

		environm ent to KEPA research office	AMR in the environm ent.	environm ent to KEPA research office/Tot al number of universitie s.				
	2.3.2.2.1.3 The director of the office of research and studies applies for funds from KEPA for projects related to AMR surveillan ce in Vet, agricultur e, and Environm ent.	2.3.2.2.1.3 % of proposed projects related to AMR in environm ent that have been funded	2.3.2.2.1.3 To secure funds for the projects in order for them to be realized	2.3.2.2.1.3 Number of proposed projects related to AMR in environm ent that have been funded/to tal number of projects related to AMR that were proposed.	2.3.2.2.1.3 Once/6 months	2.3.2.2.1.3 KEPA Research office	2.3.2.2.1.3 Checking	2.3.2.2.1. 3 Almost none
2.3.2.2.2 Increase lab capacity of KEPA Lab that	2.3.2.2.1 Letter from KEPA Director General to Technical	2.3.2.2.2.1 KEPA Lab starts reporting AMR data	2.3.2.2.2.1 To expand the capacity of the lab from	2.3.2.2.2.1 Yes/No	2.3.2.2.2.1 Once/year	2.3.2.2.2.1 KEPA lab	2.3.2.2.2.1 Checking	2.3.2.2. 1 Not available

deals with the quality of coastal water to include AMR identificat ion and surveillan ce in water and sediment	affairs sector of micro labs asking to prepare a project with a budget that aims at expanding their work from only identificat ion to antibiogra			microorga nism identificat ion to AMR detection					
	2.3.2.2.2.2 KEPA analytical central lab KEPA prepare road map for the labs to be able to do antibiogra m testing	2.3.2.2.2.2 .1 Identify the needed machinery and their budget	2.3.2.2.2.2 Road map project with budget is put to extend KEPA microbiol ogy lab from bacterial identificat ion to include also antibiogra m testing	2.3.2.2.2.2 To put a plan for how to reach the target of making KEPA lab (marine) detect AMR.	2.3.2.2.2.2 Yes /No	2.3.2.2.2 Once/6 months	2.3.2.2.2.2 KEPA Analytical Lab center	2.3.2.2.2 Checking	2.3.2.2.2. 2 KEPA marine lab identifies microorga nisms, does not detect AMR.

			2.3.2.2.2 .2 Put a plan of needed extra staffing	See above						
		2.3.2.2.3 KEPA provides the related budget	2.3.2.2.2.3 Communi cation between KEPA and ministry of finance	2.3.2.2.2.3 None						
		2.3.2.2.4 KEPA microbiol ogy labs start generatin g antibiogra m results for priority organisms		2.3.2.2.2.4 KEPA Marine lab actually generates AMR data	2.3.2.2.4 To have AMR data from marine environm ent	2.3.2.2.2.4 Yes/No	2.3.2.2.2.4 Once/Year	2.3.2.2.2.4 KEPA marine lab	2.3.2.2.2.4 Checking	2.3.2.2.2. 4 Not available
	2.3.2.2.3 Generate report about AMR surveillan ce in Environm ent	Ü		2.3.2.2.3 Yes/No						
2.3.3	2.3.3.1.1			2.3.3.1.1	2.3.3.1.1	2.3.3.1.1 Yes/No	2.3.3.1.1	2.3.3.1.1 MOH	2.3.3.1.1 Checking	2.3.3.1.1

The	KEPA,	Create a			Digital	То		Once/6			Not
national	PAAF,	digital			System is	facilitate		months			available
AMR	and PAFN	network			put	AMR data					
Surveillan	submit	for data				collection					
ce Office	available	sharing				in health					
at MOH	data to	with MOH				approach					
gathers	Central					by making					
data from	AMR					it digital					
KEPA,	surveillan										
PAAF, and	ce office										
PAFN	at MOH										
compiles	on yearly										
it with	basis										
Human											
AMR data											
and											
generates											
yearly											
ONE											
Health											
AMR											
surveillan											
ce report.											
-		2.3.3.1.2									
		Add this									
		function			2.3.3.1.2						
		to the			TOR						
		TOR of			includes						
		the focal			surveillan						
		person at			ce						
		PAAF and			function						
		KEPA and			detail.						
		food									
		safety.									
Strategic ob		reate a Natio	nal Reference	e Lab for AMF	2						
2.4.1	2.4.1.1				2.4.1.1	2.4.1.1	2.4.1.1	2.4.1.1	2.4.1.1	2.4.1.1	2.4.1.1

To have A Kuwaiti Center of Disease Control that will be overseein g bacterial, viral and AMR reference labs	Project of reference lab with its budget and having it endorsed by MOH		CDC will oversee the needs and functions of all reference laboratori al, including viral, bacterial and AMR.	To centralize all the reference labs	Yes/No	Once/year	МОН	Checking	Project is already under the umbrella of Minister of Health and under serious considera tion.
	2.4.1.2 Kuwait CDC becomes functional		2.4.1.2 Kuwait CDC is establishe d	2.4.1.2 To put the project of Kuwait CDC into action	2.4.1.2 Yes/No	2.4.1.2 Once/Year	2.4.1.2 MOH	2.4.1.2 Checking	2.4.1.2 The Minister of Health has put the project among his priorities and has provided the budget for it.
	2.4.1.3 Write and approve terms of reference for a national		2.4.1.3 TOR of AMR reference Lab are put.	2.4.1.3 To define the functions and fulfill them	2.4.1.3 Yes/No	2.4.1.3 Once	2.4.1.3 MOH/ Kuwait CDC/Medi cal Lab.Counc il	2.4.1.3 Checking	2.4.1.3 Not done yet

AMR			They						
reference			include:						
laboratory			expertise						
with			in method						
expertise			s for						
in method			confirmin						
s for			g and						
confirmin			characteri						
g and			zing						
characteri			specific						
zing			pathogens						
specific			, putting						
pathogens			guidelines						
, putting			, SOP for						
guidelines			identificat						
, SOP for			ion and						
identificat			antibiogra						
ion and			m testing						
antibiogra			of priority						
m testing			organisms						
of priority			for						
organisms			surveillan						
for			ce training						
surveillan			according						
ce training			to						
according	•		guidelines						
to			, and						
guidelines			organizing						
_									
, and			quality						
organizing			assurance						
quality			schemes						
assurance									
schemes									
2.4.1.4			2.4.1.4		2.4.1.4	2.4.1.4	2.4.1.4	2.4.1.4	2.4.1.4
Prepare a			Yes/No	2.4.1.4	Yes/No	Once	Kuwait	checking	Not yet
full			103/110		. 03/110	01100	CDC/MOH	C.ICCINIII B	

	project and proposal for the reference lab				To put the full road map and define the budget and submit it for funding					
	2.4.1.5 Prepare a budget for the building, equipmen t and man power of the reference Lab			2.4.1.5 Budget for AMR lab is approved within the budget of Kuwait CDC	2.4.1.5 To provide the needed budget for Manpowe r and equipmen t	2.4.1.5 Yes/No	2.4.1.5 Once/year	2.4.1.5 Kuwait CDC/ MOH	Checking	Not yet
	2.4.1.6 Execution of the Reference Lab			2.4.1.6 Reference lab is set and functionin g	2.4.1.6 To fulfill the functions listed in the TOR	2.4.1.6 Yes/no	2.4.1.6 Once/year	2.4.1.6 Kuwait CDC	2.4.1.6 Checking	2.4.1.6 Not yet
2.4.2 Delegate the function of surveillan ce of the reference Lab pending	2.4.2.1 Identify institution affiliated to MOH that will host the manpowe r and activities	2.4.2.1.1 A letter from MOH to all labs that asks them to coordinat e with the GLASS lab		2.4.2.1.1 Already available	2.4.2.1.1 To start surveillan ce while waiting for the reference lab, which might					2.4.2.1.1 AMR surveillan ce is being submitted to GLASS National AMR not

its readiness	of the AMR Surveillan ce office for the actual work on the National and GLASS AMR reports pending the finalizatio n of the reference lab.	by delivering AMR data for the purpose of AMR Surveillan ce				take few years					finalized yet A plan is put above
2.5.1 Lab audit by lab accreditation body according to microbiol ogist consultant in charge, external quality control and use of CLSI guidelines	2.5.1.1 Accreditation of the medical Laboratories by Medical Laboratories Council	2.5.1.1.1 Involve all laboratori es in external quality assurance programs	2.5.1.1.1.1 Mandate that all labs including all private should be accredited and all perform external QC.	2.5.1.1.1.1 .1 List of accreditati on bodies accepted in Kuwait, all labs should be accredited by one of them	2.5.1.1.1.1 .1 Number of internatio nal accreditati on bodies that are recomme nded in Kuwait for the labs to get their accreditati on	2.5.1.1.1.1 .1 To identify internatio nal accreditati on bodies to perform accreditati on of the labs in Kuwait and not to be limited	2.5.1.1.1.1 .1 Number	2.5.1.1.1.1 .1 Once/Year	2.5.1.1.1.1 .1 MOH/ Reference lab/ Medical Lab.counc il	2.5.1.1.1.1 .1 Checking	2.5.1.1.1 1.1 CAP

			to the			
			local			
			accreditati			
			on activity			

Annex 4

A. IPC Strategic Plan

Strategic interventions	Activity	Sub-activity	Sub-sub- activity	Sub-sub-sub- activity	Date	Milestone
	1 Emphasize the natio	nal infection prevention	n and control progran	n in healthcare at the N	lational and Health ca	re facility levels.
3.1.1 Create a formal organizational structure to ensure proper development and use of infection prevention and control policies and strategies	3.1.1.1 Infection Control Organizational structure is already established and functioning.	-	-	-	3.1.1.1 Done 1980	3.1 5 years
	3.1.1.2 To make official that IPC is part of the organogram of each hospital with Terms of Reference and allocated staff	3.1.2.1 Decree from MOH about IPC department in each hospital with TOR, and staffing organization			3.1.2.1 2 Months	
	3.1.1.3 IPC departments in hospitals and they work closely with the central IPC Directorate	3.1.1.3.1 To impose IPC programs in all hospitals	3.1.1.3.1.1 IPC program availability is checked by accreditation standards		3.1.1.3.1.1 Already available	
		3.1.1.3.2 IPC programs are available in all hospitals and they work under the	-	-	3.1.1.3.2 Already available	

		umbrella of the Central IPC directorate.	3.1.2.1.1.1 IPC			
3.1.2 Implement laws for mandatory compliance of the facilities with IPC standards	3.1.2.1 Decree from MOH	3.1.2.1.1 Decree that recommends that all HCF have to abide by IPC Recommendations.	Directorate puts the detailed IPC measures that should be required from each HCF, and sent to MOH to be included in ministry	-	3.1.2.1.1.1 2 months	
			3.1.2.1.1.2 Decree is issued by MOH	-	3.1.2.1.1.2 3 months	
		3.1.2.1.2 Medical responsibilities. Add details IPC standard compliance in hospitals	3.1.2.1.2.1 Add compliance to IPC measures and recommendations to the list of medical responsibilities of hospitals as listed by MOH	-	3.1.2.1.2.1 3 months	
3.1.3 WHO IPC Core component based guidelines are established in all HCF	3.1.3.1 Availability of endorsed IPC based on WHO IPC core components	-	-	-	3.1.3.1 Done	
	3.1.3.2 IPC policies and procedures are included in hospital accreditation standards	-	-	-	3.1.3.2 Done	
	3.1.3.3 IPC policies and procedures are being audited	-	-	-	3.1.3.3 Done	

regularly by accreditation bodies				
3.1.3.4 Natio infection con policies and guidelines and distributed a periodically updated	3.1.3.4.1 Regular update of national	3.1.3.4.1.1 On going, but needs to be speeded up by increasing manpower in IPC nationwide in central office and in hospitals	-	3.1.3.4.1.1 2 years
	3.1.3.4.2 A direct channel with WHO consultant regarding few queries and new recommendations	3.1.3.4.2.1 Communicate this need to WHO office in Kuwait	3.1.3.4.2.1.1 Letter from MOH/IPC directorate to WHO office in Kuwait asking for direct channel with a consultant for queries whenever they occur	3.1.3.4.2.1.1 1 month
		3.1.3.4.2.2 Communicate this need to Regional WHO office	3.1.3.4.2.2 Request from WCO to Regional office.	3.1.3.4.2.2 2 months
	3.1.3.4.3 Regular update of National IPC policies and procedures	3.1.3.4.3.1 Need updating + speeding up the process		3.1.3.4.3.1 ongoing
3.1.3.5 Natio Guidelines ar policies are b taught	periodically	-	-	3.1.3.5.1 Already being performed

		3.1.3.5.2 Continuous training of HCW about IPC policies in hospitals is available	3.1.3.5.2.1 Mandate from MOH that it is mandatory to all HCW to attend at least 1 yearly and 1 upon employment		3.1.3.5.2.1 2 months
			3.1.3.5.2.2 Improve attendance to these general HCW IPC training sessions	3.1.3.5.2.2.1 Attendance is reported to hospital administrators and action taken to improve attendance	3.1.3.5.2.2.1 1 year
				3.1.3.5.2.2.2 Attendance evidence is available in yearly Hospital IPC report and monitored by IPC Directorate	3.1.3.5.2.2.2 1 year
3.1.4 Monitor and evaluate compliance with the IPC guidelines	3.1.4.1 Checklists and audit tools are prepared and applied	-	-	-	3.1.4.1 Started years ago, and ongoing for new policies.
	3.1.4.2 Regular updates of these audit tools	3.1.4.2.1 Check new policies if they have their audit tools prepared and disseminated and check old tools if they need update	-	-	3.1.4.2.1 Ongoing
	3.1.4.3 Audit plan available in hospitals	3.1.4.3.1 Audit plan is checked in standards of the	-	-	3.1.4.3.1 done ongoing

	National Accreditation Program applied by the Quality and Accreditation Directorate 3.1.4.3.2 Central audit plan is being			3.1.4.3.2
	prepared annually and distributed to all hospitals	-	-	Done and ongoing yearly
	3.1.4.3.3 Audit report is sent to central IPC directorate with corrective actions.	-	-	3.1.4.3.3 Done and ongoing
3.1.4.4 Establish national process indicators	3.1.4.4.1 1st set of process indicators: -CLABSI -CAUTI -VAP	3.1.4.4.1.1Put the basic training material	-	3.1.4.4.1.1Already available
		3.1.4.4.1.2Update of the already available bundle of care in each of the mentioned bundles, by including new evidence- based items to the bundle	3.1.4.4.1.2 Prepare the updated bundles	3.1.4.4.1.2 2 years
		3.1.4.4.1.3Train the IPC professionals nationwide on the updated bundles	3.1.4.4.1.3.1 TOT workshops	3.1.4.4.1.3.1 2 years
		3.1.4.4.1.4 Engage all HCW with the	3.1.4.4.1.4.1	3.1.4.4.1.4.1 CLABSI months

	bundles by training,	Prepare the ground	Each bundle 1 y
	audit and feedback.	in the hospitals: Letter to hospital	
		directors about the	
		bundles project its	
		importance and its	
		processes	
		3.1.4.4.1.4.2	
		Update the	
		material to be	3.1.4.4.1.4.2
		taught to HCW by	2 years
		IPC professionals,	,
		according to the	
		updated bundles. 3.1.4.4.1.4.3	
		In-hospital training	
		sessions by IPC	
		professionals to all	
		concerned HCW.	244442
		Include Bundle	3.1.4.4.1.4.3
		training in the	4 years
		yearly IPC training	
		and in peri-	
		employment	
		training.	
		3.1.4.4.1.4.4	
		Auditing compliance to	
		these bundles on	3.1.4.4.1.4.4
		monthly basis and	
		report to IPC	5 years
		directorate	
		monthly	
3.1.4.4.2	3.1.4.4.2.1		3.1.4.4.2.1
Update outcome	Review updated	-	6 months
indicators for the 3	CDC outcome		o mondis

		bundles above (CLABSI, CAUTI, VAP) according to new CDC/ guidelines	indicators and update already measured bundles outcome indicators accordingly		
		3.1.4.4.3 Integrate process and outcome indicators for each bundle, and use in the analysis and corrective actions	3.1.4.4.3.1 Put a road map for integrating analysis of outcome and process indicators related to the bundles	-	3.1.4.4.3.1 1 year
		3.1.4.4.4 Introduce SSI bundle	3.1.4.4.4.1 preparation of material	-	3.1.4.4.4.1 4 years
			3.1.4.4.4.2 Implementation of SSI bundle	3.1.4.4.4.2.1 Teaching of IPC professionals by central IPC directorate	3.1.4.4.4.2.1 5 years
				3.1.4.4.4.2.2 Teaching of healthcare providers inside the hospitals by IPC professionals	3.1.4.4.4.2.2 5 years
				3.1.4.4.4.2.3 Data collection	3.1.4.4.4.2.3 5 years
3.1.5	3.1.5.1	3.1.5.1.1	3.1.5.1.1.1		
IPC programs/office in LTCF working in close cooperation	Mandate from MOH that IPC program/with	Communication between MOH and Public Authority of	Letter from MOH to Public Authority of Disability Affairs		3.1.5.1.1.1 2 months
with IPC directorate in MOH	employee with TOR and specified	Disability Affairs regarding IPC in	asking to establish IPC in each LTCF		

	budget are available in LTCF the central and the "branches".	LTCF and being overseen by the IPC directorate	and to have the personnel work closely with IPC directorate delegate to LTCF.			
			3.1.5.1.1.2Letter from Public Authority of Disability Affairs to LTCF that recommends IPC in each LTCF and organizes the communication with MOH		3.1.5.1.1.2 2 months	
		3.1.5.1.2 Established Guidelines for IPC in LTCF	3.1.5.1.2.1 Preparation of the LTCF guidelines and their endorsement by Public Authority of Disability Affairs	-	3.1.5.1.2.1 1 year	
			3.1.5.1.2.2 Dissemination of LTCF Policies and Procedures	3.1.5.1.2.2.1 TOT of LTCF IPC focal persons	3.1.5.1.2.2 1.5 years	
Strategic objective 3.		AMR information digit	talized and sent to IPC	directorate as a digital	report	
3.2.1 Software for hospital data entry, analysis and generating both hospital and national reports	3.2.1.1 Subcommittee from IPC directorate sends a letter to Information system department at MOH explaining what is required				3.2.1.1 2 months	3.2 5 years

	and asking for the				
	software				
	3.2.1.2				
	Health Information				
	system dep.			2242	
	presents projects			3.2.1.2	
	(either in-house, or			12 months	
	ready to buy) with				
	budget providing				
	3.2.1.3				
	Agreement				
	between IPC			3.2.1.3	
	directorate and			1 year 3 months	
	information system			1 year 3 months	
	dep. to choose one				
	system				
	3.2.1.4			3.2.1.4	
	Apply the system in			1 year 9 months	
	the hospitals			1 year 5 monens	
	3.2.1.5				
	Apply the Digital				
	HAI/AMR				
	System for National			3.2.1.5	
	data collection			2 years	
	from individual				
	hospitals to central				
	IPC directorate				
2.2.2.To Chift all	3.2.2.1 MOH				
3.2.2 To Shift all	recommends from	3.2.2.1.1 Letter			
hospital information system	hospitals to have an electronic patient	from MOH to		3.2.2.1.1	
from paper to	and data	hospitals		2 months	
digital mode.	information	Hospitals			
uigitai iiioue.	system.				
	system.				

	3.2.2.2 Increase the number of hospitals that have digitalized their medical information system	-	-	-	3.2.2.2 5 years	
3.2.3 Connect Infection Control Departments to electronic medical records	3.2.3.1 Connect Infection Control departments in all hospitals with the Heath information system and Laboratory information system at the hospital level	3.2.3.1.1 Letter from MOH to ask all hospitals to connect IC dep. With HIS system and LIS at the hospital level	-	-	3.2.3.1.1 2 months	
		3.2.3.1.2 Decree from MOH to all hospitals who have an electronic Health information system to link their Lab AMR data to the Hospital system	-	-	3.2.3.1.2 4 months	
	3.2.3.2 Develop electronic outbreak reporting system for data entry, analysis and generating reports.				3.2.3.2 3 years	
	3.2.3.3 Develop electronic reporting system for data entry, analysis and generating hospital/national				3.2.3.3 3 years	

Strategic objective 3.	reports for bundles of care. 3.2.3.4 Develop electronic infection prevention and control system capable of selfidentifying HAI and detecting outbreaks 3 Enhance IPC educations	3.2.3.4.1 Electronic system has a function for self-diagnosis of HAIs and-automatically detects outbreak	d specialties and amon	g HCW.	3.2.3.4.1 5 years	
3.3.1 Include IPC as a core element in education and training of healthcare professionals	3.3.1.1 Include infection control in undergraduate curricula for health care students (medical, nursing, dental and pharmaceutical).	3.3.1.1.1 Prepare module about IPC and AMR to be included in all health specialties in universities/PAAET.	-	-	3.3.1.1.1 2 years	3.3 5 years
		3.3.1.1.2 MOH letter to Ministry of higher education and The Public Authority for Applied Education and Training to ask to include these modules into the undergrads and graduate health students' curricula	-	-	3.3.1.1.2 2 years	

		3.3.1.1.3 Modules sent to Kuwait university/ The Public Authority for Applied Education and Training to include it in curricula			3.3.1.1.3 2.5 years	
3.3.2 Enhance IPC education among undergraduates and higher education health students	3.3.2.1 Include infection control in continuing education of healthcare workers and mandatory prerequisite for promotion	3.3.2.1.1 Provide IPC courses with CE credits in universities, hospitals, MOH, PAAET, PAAF Include CME Credit 10% IPC			3.3.2.1.1 2 years	
		3.3.2.1.2 Request that when CE credits for physicians, nurses, or pharmacists is needed, 10% of these credits should be about IPC.			3.3.2.1.2 3 months	
		3.3.2.1.3 In hospitals, yearly and upon employment, mandatory attendance of IPC lectures	-	-	3.3.2.1.3 3 months	

	3.3.2.2 Hygiene and IPC education provided outside formal teaching programs	3.3.2.2.1 Require IPC education by drug companies, hospitals, LTCFs, FAO, and medical/veterinary associations	3.3.2.2.1.1 Request to include few topics about IPC in conferences agendas	3.3.2.2.1.1 Letter to be sent	3.3.2.2.1.1 6 months	
Strategic objective 3.	4 Prevent occupationa	l infections in HCWs				
3.4.1 Develop a comprehensive program of vaccination to protect HCWs from vaccine preventable diseases	-		-	-	3.4.1 Available And applied	3.4 5 years
3.4.2 Develop policies and procedures to protect healthcare workers from occupational exposure to infections.	-	-	-	-	3.4.2 Available And applied	
3.4.3 Availability of Airborne Isolation Rooms in all hospitals	3.4.3.1 screening mapping and updating the recommendations according to the expected need according to need	-	-	-	3.4.3.1 5 years	
3.4.4 Mandate to hospitals to provide airborne isolation rooms (AII)	-	-	-	-	3.4.4 5 years	

available as per the						
updated						
recommendations						
Strategic objective 3.	5 Adequate waste mai	nagement in hospitals				
3.5.1 Put National Guidelines	3.5.1.1 National policies, strategies and plans for healthcare waste management is in place	3.5.1.1.1 Unify recommendations between KEPA and MOH	3.5.1.1.1.1 Create a committee from MOH and KEPA that unifies the recommendation and the road map of all types of waste that are generated in healthcare		3.5.1.1.1.1 6 months	3.5 5 years
			3.5.1.1.1.2 The committee reviews waste policies that are related to HCF and unifies recommendations and provides the proper ways of supervision	-	3.5.1.1.1.2 1 year	
3.5.2 Supervision of waste disposal from public hospitals	3.5.2.1 Create a committee for the supervision of waste disposal from public hospitals	-	-	-	3.5.2.1 Available and ongoing	
3.5.3 Supervision of waste disposal in private hospitals	3.5.3.1 Create a committee for the supervision of waste disposal in private hospitals	-	-	-	3.5.3.1 1 year	

3.5.4 Waste policies are being audited by KEPA	3.5.4.1 Organization of supervision of waste disposal inside and from private hospitals	-	-	-	3.5.4.1 1.5 years	
	3.5.4.2 Designate 1 unit at KEPA to oversee the waste in hospitals		-	-	3.5.4.2 1 year	
Strategic objective 3.	6 To enhance infection		ol programs in veterin	ary and agriculture set	tings	I
3.6.1 Availability of IPC guidelines in veterinary practices	3.6.1.1 Implementation of OIE guidelines regarding IPC in veterinary field	3.6.1.1.1 Committee in PAAF that regularly checks updates in OIE guidelines regarding IPC in veterinary field, and updates the guidelines accordingly	-	-	3.6.1.1.1 6 months	3.6 5 years
		3.6.1.1.2 Overseeing the implementation of national IPC guidelines in Vet field and OIE good practices	3.6.1.1.2 Regular audits to farms	3.6.1.1.2 Put a yearly audit plan	3.6.1.1.2 1 year	
3.6.2 Include IPC in licensing of Vets	3.6.2.1 Include questions about OIE IPC guidelines and	3.6.2.1.1 Prepare a set of questions related to IPC in Vets to be	-	-	3.6.2.1.1 6 months	

	vaccinations in all vets licensing exams or interviews.	included in licensing exams and interviews.				
3.6.3 Include IPC in Continuous education of Vets and professionals in Agriculture	3.6.3.1 Prepare a slide bank related to IPC in vet and agriculture, to be given to drug companies that usually do the continuous education activities.	-	-	-	3.6.3.1 6 months	
	3.6.3.2 Mandate that all drug company events or seminars include a specific number of slides related to IPC and AMR.	-	-	-	3.6.3.2 9 months	
3.6.4 Include hygiene and infection prevention and control as core content in training of veterinary professionals.	3.6.4.1 Close communication between. FAO and PAAF to put a joint yearly plan of education activities related to IPC.	-	-		3.6.4.1 1 year	
	3.6.4.2 Include hygiene and infection prevention and control as	3.6.4.2.1 IPC credits mandatory for promotion in Vet. Associations	-	-	3.6.4.2.1 6 months	

mandatory prerequisite for promotion of for			
the veterinary workers.			

B. <u>IPC Monitoring Plan</u>

Strategic interventions Strategic Obs	Activity	Sub- activity mphasize the	Sub-sub- activity	Sub-sub- sub- activity	Indicator	Purpose rol program i	Calculatio n n healthcare	Frequenc Y at the Nation	Data source nal and Health	Method	Baseline levels.
3.1.1 Create a formal organizati onal structure to ensure proper developm ent and use of infection preventio n and control policies and strategies	3.1.1.1 Infection Control Organizati onal structure is already establishe d and functionin g.	-	-	-	3.1.1.1 Already available						3.1.1.1 Already available
	3.1.1.2 To make official that IPC is part of	3.1.2.1 Decree from MOH about IPC			3.1.2.1 The decree was issued.						3.1.2.1 Already available

the organog m of each hospital with Terms of Reference and allocated staff	h hospital with TOR, and staffing organizati on						
3.1.1.3 IPC departments in hospitals and they work closely with the central IPC Director e	3.1.1.3.1 To impose IPC programs in all hospitals	3.1.1.3.1.1 IPC program availabilit y is checked by accreditati on standards		3.1.1.3.1.1 already exists			3.1.1.3.1. 1 Already establishe d
	3.1.1.3.2 IPC programs are available in all hospitals and they work under the umbrella of the	-	-	3.1.1.3.2 % of Hospitals with IPC programs			3.1.1.3.1. 2 Already establishe d

		Central IPC directorat e.									
3.1.2 Implemen t laws for mandator y complianc e of the facilities with IPC standards	3.1.2.1 Decree from MOH	3.1.2.1.1 Decree that recomme nds that all HCF have to abide by IPC Recomme ndations.	3.1.2.1.1.1 IPC Directorat e puts the detailed IPC measures that should be required from each HCF, and sent to MOH to be included in ministry	-	3.1.2.1.1.1 List is sent from IPC Directorat e to MOH Director		3.1.2.1.1.1 Yes/No	3.1.2.1.1.1 Once	3.1.2.1.1.1 IPC Directorat e/ MOH	3.1.2.1.1.1 Checking	3.1.2.1.1. 1 Not available
			3.1.2.1.1.2 Decree is issued by MOH	-	3.1.2.1.1.2 Decree availabilit y and dissemina tion	3.1.2.1.1.2 Abiding by IPC standards is a must in hospitals	3.1.2.1.1.2 Yes/No	3.1.2.1.1.2 once	3.1.2.1.1.2 IPC Directorat e/ MOH	3.1.2.1.1.2 Checking	3.1.2.1.1. 2 Not available
		3.1.2.1.2 Medical responsibi lities. Add details IPC standard	3.1.2.1.2.1 Add complianc e to IPC measures and recomme ndations	-	3.1.2.1.2.1 The Medical responsibi lity list is updated and it includes	3.1.2.1.2.1 To make it official through medical Decisions that obedience	3.1.2.1.2.1 Yes/No	3.1.2.1.2.1 Once /year until they appear in the law	3.1.2.1.2.1 IPC Directorat e/ MOH	3.1.2.1.2.1 Checking	3.1.2.1.2. 1 IPC exists in hospitals and centrally but there

		complianc e in hospitals	to the list of medical responsibi lities of hospitals as listed by MOH		the statement about IPC measures.	to IPC measures is a must in Hospitals			is no law, so far, that enforces it in the hospitals.
3.1.3 WHO IPC Core compone nt based guidelines are establishe d in all HCF	3.1.3.1 Availabilit y of endorsed IPC based on WHO IPC core compone nts	-	-	-	3.1.3.1 Available				3.1.3.1 Available
	3.1.3.2 IPC policies and procedure s are included in hospital accreditati on standards	-	-	-	3.1.3.2 Available				3.1.3.2 Available
	3.1.3.3 IPC policies and procedure s are being audited regularly by	-	-	-	3.1.3.3 Available				3.1.3.3 Available

	creditati bodies										
Natinfe con poli and guid are dist dan per	olicies Rud unidelines e III gstribute g	3.1.3.4.1 Regular update of national PC guidelines	3.1.3.4.1.1 On going, but needs to be speeded up by increasing manpowe r in IPC nationwid e in central office and in hospitals	-	3.1.3.4.1.1 Proportio n of staff that is needed that has been actually employed	3.1.3.4.1.1 speed up the process of updating policies and guidelines	3.1.3.4.1.1 Number of newly employed in IPC. /Number needed X 100	3.1.3.4.1.1 Once /6 months until all are employed.	3.1.3.4.1.1 IPC Directorat e / MOH	3.1.3.4.1.1 Checking	3.1.3.4.1. 1 The request has been sent to MOH and approval was given, employm ent still in process.
	A c w c ri fi q a	3.1.3.4.2 A direct channel with WHO consultant regarding few queries and new recomme andations	3.1.3.4.2.1 Communi cate this need to WHO office in Kuwait	3.1.3.4.2.1 .1 Letter from MOH/IPC directorat e to WHO office in Kuwait asking for direct channel with a consultant for queries whenever they occur	3.1.3.4.2.1 .1 Percent of queries to WHO that have been timely answered.	3.1.3.4.2.1 To have a direct channel with a WHO consultant that helps giving answers to questions based on WHO guidelines and recomme ndations	3.1.3.4.2.1 Number of timely answered queries /Number of needed consultati ons X 100	3.1.3.4.2.1 Once/year	3.1.3.4.2.1 MOH/WH O	3.1.3.4.2.1 Checking	3.1.3.4.2. 1 Not available

					that might arise					
		3.1.3.4.2.2 Communi cate this need to Regional WHO office	3.1.3.4.2.2 Request from WCO to Regional office.							
	3.1.3.4.3 Regular update of National IPC policies and procedure s	3.1.3.4.3.1 Need updating + speeding up the process		3.1.3.4.3.1 % of hospitals that have polices updated in the last 2 years or less.		3.1.3.4.3.1 Number of hospitals with updated policies in the last 2 years or less/Total number of hospitals X 100	3.1.3.4.3.1 Once/Year	3.1.3.4.3.1 Hospitals / Central IPC Directorat e	3.1.3.4.3.1 Checking	3.1.3.4.3. 1 Partially available
3.1.3.5 National IPC Guidelines and policies are being taught	3.1.3.5.1 TOT periodicall y performe d nationally	-	-	3.1.3.5.1 % IPC profession als that are trained annually	3.1.3.5.1 To have a unified centralize d training of IPC profession als.	3.1.3.5.1 Number of IPC profession als trained annually /total number of IPC profession als X100	3.1.3.5.1 Once/year	3.1.3.5.1 MOH/ Central IPC Directorat e	3.1.3.5.1 Checking	3.1.3.5.1 Ongoing, Exact numbers are unknown.
	3.1.3.5.2 Continuou s	3.1.3.5.2.1 Mandate		3.1.3.5.2.1 Decree is done and						3.1.3.5.2. 1 Yes/no

training of HCW about IPC policies in hospitals is available	from MOH that it is mandator y to all HCW to attend at least 1 yearly and 1 upon employm ent		sent to hospitals.					
	3.1.3.5.2.2 Improve attendanc e to these general HCW IPC training sessions	3.1.3.5.2.2 .1 Attendanc e is reported to hospital administr ators and action taken to improve attendanc e	3.1.3.5.2.2 .1 % of HCW that attend the training sessions yearly	3.1.3.5.2.2 .1 Number of HCW attendees /total number of HCW /year X100	3.1.3.5.2.2 .1 Once/year	3.1.3.5.2.2 .1 Hospitals/ Central IPC Directorat e	3.1.3.5.2.2 .1 Data collection	3.1.3.5.2. 2.1 Training is recomme nded and provided, attendanc e is suboptim al
		3.1.3.5.2.2 .2 Attendanc e evidence is available in yearly Hospital IPC report						

				and monitored by IPC Directorat e							
3.1.4 Monitor and evaluate complianc e with the IPC guidelines	3.1.4.1 Checklists and audit tools are prepared and applied	_	_	_	3.1.4.1 and 3.1.4.2 Percent of policies that have correspon ding audit tools.	3.1.4.1 and 3.1.4.2 To make audit tools unified nationally and readily available and to facilitate National data collection if needed	3.1.4.1 and 3.1.4.2 Number of policies with audit tools/Tota I number of policies X100	3.1.4.1 and 3.1.4.2 Once/year	3.1.4.1 and 3.1.4.2 IPC directorat e	3.1.4.1 and 3.1.4.2 Checking	3.1.4.1 and 3.1.4.2 Partially available
	3.1.4.2 Regular updates of these audit tools	3.1.4.2.1 Check new policies if they have their audit tools prepared and dissemina ted and check old tools if	-	-							

	they need update									
3.1.4.3 Audit plan available in hospitals	3.1.4.3.1 Audit plan is checked in standards of the National Accreditat ion Program applied by the Quality and Accreditat ion Directorat e	-	-	3.1.4.3.1 Done and ongoing						3.1.4.3.1 Done and ongoing
	3.1.4.3.2 Central audit plan is being prepared annually and distribute d to all hospitals	-	-	3.1.4.3.2 Done and ongoing	3.1.4.3.2 % of hospitals that perform audit according to the centralize d audit plan and audit. tools.	3.1.4.3.2 Number of hospitals that perform audit according to the centralize d audit plan and audit tools/total number of	3.1.4.3.2 Once/year	3.1.4.3.2 Hospitals/ Digital central IPC program	3.1.4.3.2 Data collection	3.1.4.3.2 Partially performe d

	3.1.4.3.3 Audit report is sent to central IPC directorat e with corrective actions.	-	-	3.1.4.3.3 Done and ongoing	3.1.4.3.3 % of hospitals that report the results of their audits to IPC	hospitals X100 3.1.4.3.3 Number of hospitals that report the results of their audits to IPC/Total number of hospitals X100	3.1.4.3.3 Once/year	3.1.4.3.3 Hospitals/ Digital central IPC program	3.1.4.3.3 Data collection	3.1.4.3.3 Partially performe d
3.1.4.4 Establish national process indicators	3.1.4.4.1 1 st set of process indicators: -CLABSI -CAUTI -VAP	3.1.4.4.1.1 Put the basic training material	-	3.1.4.4.1.1 Each one of the 4 procedure s has its own bundle establishe d	3.1.4.4.1.1 To improve quality of care by using the bundle approach in all HCF	3.1.4.4.1.1 Yes/No	3.1.4.4.1.1 Checking	3.1.4.4.1.1 MOH/ IPC directorat e	3.1.4.4.1.1 Checking	3.1.4.4.1. 1 Not available
		3.1.4.4.1.2 Update of the already available bundle of care in each of the mentione d bundles, by	3.1.4.4.1.2 Prepare the updated bundles	3.1.4.4.1.2 Availabilit y of 3 updated bundles	3.1.4.4.1.2 To make quality of care up- to-date regarding the 3 procedure s and to improve outcome	3.1.4.4.1.2 Yes or no x 3 (For each bundle)	3.1.4.4.1.2 Once /6 months until availabilit y	3.1.4.4.1.2 IPC directorat e	3.1.4.4.1.2 Checking	3.1.4.4.1. 2 Old version of bundles available

	including new evidence- based items to the bundle								
	3.1.4.4.1.3 Train the IPC profession als nationwid e on the updated bundles	3.1.4.4.1.3 .1 TOT workshop s	3.1.4.4.1.3 % IPC profession als trained about the new bundles	3.1.4.4.1.3 To dissemina te the updated bundles to all IPC profession als nationally	3.1.4.4.1.3 Number of IPC profession als trained about the new bundles /Total number of IPC profession als.	3.1.4.4.1.3 Once/year	3.1.4.4.1.3 IPC directorat e/Hospital s	3.1.4.4.1.3 Data collection	3.1.4.4.1. 3 IPC profession als have been trained on the older version of the bundles.
	3.1.4.4.1.4 Engage all HCW with the bundles by training, audit and feedback.	3.1.4.4.1.4 .1 Prepare the ground in the hospitals: Letter to hospital directors about the bundles project its importanc	3.1.4.4.1.4 .1 Letter sent or not		3.1.4.4.1.4 .1 Yes /No	3.1.4.4.1.4 .1 Once	3.1.4.4.1.4 .1 IPC directorat e/ MOH	3.1.4.4.1.4 .1 Checking	3.1.4.4.1. 4.1 Not available

	e and its processes 3.1.4.4.1.4 .2 Update the material to be taught to HCW by IPC profession als, according to the updated bundles.	3.1.4.4.1.4 .2 % of material directed to HCW that has been updated	3.1.4.4.1.4 .2 To synchroni ze the teaching of the IPC bundle material with the training of the HCW other than the IPC profession als.	3.1.4.4.1.4 .2 Updated bundle material directed to HCW/Nu mber of bundles x100	3.1.4.4.1.4 .2 Once/Year	3.1.4.4.1.4 .2 IPC directorat e/Hospital training material	3.1.4.4.1.4 .2 Checking	3.1.4.4.1. 4.2 The material to be taught to HCW is based on. the classical bundle training
	3.1.4.4.1.4 .3 In- hospital training sessions by IPC profession als to all concerned HCW. Include Bundle training in the yearly IPC training	3.1.4.4.1.4 .3 Complianc e with performa nce indicator for each bundle	3.1.4.4.1.4 .3 To increase the complianc e to bundles to prevent HAI and improve the quality of care		3.1.4.4.1.4 .3 Once/Year	3.1.4.4.1.4 .3 Hospitals	3.1.4.4.1.4 .3 Checking	3.1.4.4.1. 4.3 The material to be taught to HCW is based on. the classical bundle training

		and in peri- employm ent training. 3.1.4.4.1.4 .4 Auditing complianc e to these bundles on monthly basis and report to IPC directorat e monthly	3.1.4.4.1.4 .4 Process Indicator for each of the 3 bundles: For each bundle: % Complianc e by the specific bundle /opportun ities of audit for each bundle	3.1.4.4.1.4 .4 To increase the complianc e with the updated bundles and improve the related quality of care	3.1.4.4.1.4 .4 % of auditing opportuni ties that are compliant with the bundle checklist/t otal number of audited complianc e with updated bundle X 100	3.1.4.4.1.4 .4 twice/yea r	3.1.4.4.1.4 .4 Hospitals Central Digital IPC system	3.1.4.4.1.4 .4 Data collection	3.1.4.4.1. 4.4 There are already process indicators related to the old bundles
3.1.4.4.2 Update outcome indicators for the 3 bundles above (CLABSI, CAUTI, VAP)	3.1.4.4.2.1 Review updated CDC outcome indicators and update already measured	-	3.1.5.4.2.1 Number of outcome indicator tools and benchmar k is updated.	3.1.5.4.2.1 To be in line with internatio nal data benchmar k.	3.1.5.4.2.1 Number of updates outcome bundles	3.1.5.4.2.1 Once until all 3 indicators are updates	3.1.5.4.2.1 IPC Directorat e	3.1.5.4.2.1 Checking	3.1.5.4.2. 1 Outcome bundles methodol ogy need revision and update

according to new CDC/ guidelines	bundles outcome indicators accordingl y								
3.1.4.4.3 Integrate process and outcome indicators for each bundle, and use in the analysis and corrective actions	3.1.4.4.3.1 Put a road map for integratin g analysis of outcome and process indicators related to the bundles	-	3.1.5.4.3.1 Proportio n of bundles whose analysis integrates process and outcome data	3.1.5.4.3.1 To use all the available data and means to improve outcome results	3.1.5.4.3.1 Number of bundles whose analysis integrates process and outcome data/3	3.1.5.4.3.1 Once/6 months	3.1.5.4.3.1 IPC Directorat e	3.1.5.4.3.1 Data collection	3.1.5.4.3. 1 Process and outcome bundles are not being interprete d together yet
3.1.4.4.4 Introduce SSI bundle	3.1.4.4.4.1 preparatio n of material	-	3.1.5.4.4.1 material that is prepared	3.1.5.4.4.1 To track and minimize SSI	3.1.5.4.4.1 Yes/no	3.1.5.4.4.1 Once	3.1.5.4.4.1 IPC Directorat e	3.1.5.4.4.1 Literature search WHO/Inte rnational literature	3.1.5.4.4. 1 Not done
	3.1.4.4.4.2 Implemen tation of SSI bundle	3.1.4.4.4.2 .1 Teaching of IPC profession als by central IPC directorat e	3.1.5.4.4.2 % hospitals start submittin g process and outcome indicators for SSI.	3.1.5.4.4.2 To measure SSI complianc e to bundles elements and to track SSI incidence	3.1.5.4.4.2 Number of hospitals start submittin g process and outcome indicators for SSI/	3.1.5.4.4.2 Once/Year	3.1.5.4.4.2 Hospitals/ IPC directorat e	3.1.5.4.4.2 Checking	3.1.5.4.4. 2 Not available

						Total number of hospitals x100		
				3.1.4.4.4.2 .2 Teaching of healthcar e providers inside the hospitals by IPC profession als				
				3.1.4.4.4.2 .3 Data collection				
3.1.5 IPC programs/ office in LTCF working in close cooperati on with IPC directorat e in MOH	3.1.5.1 Mandate from MOH that IPC program/ with employee with TOR and specified budget are available in LTCF	3.1.5.1.1 Communi cation between MOH and Public Authority of Disability Affairs regarding IPC in LTCF and being overseen	3.1.5.1.1.1 Letter from MOH to Public Authority of Disability Affairs asking to establish IPC in each LTCF and to have the		3.1.5.1.1.1 Present			

the central and the "branches ".	by the IPC directorat e	personnel work closely with IPC directorat e delegate to LTCF.								
		3.1.5.1.1.2 Letter from Public Authority of Disability Affairs to LTCF that recomme nds IPC in each LTCF and organizes the communic ation with MOH		3.1.5.1.1.2 Letter sent		3.1.5.1.1.2 Yes /No	3.1.5.1.1.2 Once	3.1.5.1.1.2 MOH	3.1.5.1.1.2 Checking	3.1.5.1.1. 2 Partially available
	3.1.5.1.2 Establishe d Guidelines for IPC in LTCF	3.1.5.1.2.1 Preparatio n of the LTCF guidelines and their endorsem ent by Public Authority	-	3.1.5.1.2.1 Availabilit y of the endorsed guidelines and set of policies and procedure s	3.1.5.1.2.1 Proper unified IPC practices at LTCF	3.1.5.1.2.1 Yes /No	3.1.5.1.2.1 Once	3.1.5.1.2.1 MOH	3.1.5.1.2.1 Checking	3.1.5.1.2. 1 Not available

Strategic of	ojective 3.2 To	have the HA	of Disability Affairs 3.1.5.1.2.2 Dissemina tion of LTCF Policies and Procedure s	3.1.5.1.2.2 .1 TOT of LTCF IPC focal persons	3.1.5.1.2.2 % focal IPC persons in LTCF trained on the guidelines /total number of LTCF IPC focal persons.	3.1.5.1.2.1 Proper unified IPC practices at LTCF	3.1.5.1.2.1 Number of focal persons trained /total number of focal persons *100	3.1.5.1.2.1 Once/year	мон	3.1.5.1.2.1 Checking	3.1.5.1.2. 1 Not available
3.2.1 Software for hospital data entry, analysis and generatin g both hospital and national reports	3.2.1.1 Subcommittee from IPC directorate sends a letter to Information system department at MOH explaining what is required and asking for the software 3.2.1.2				3.2.1 Software developed and working	3.2.1 Simplify the follow-up on tracking AMR/HAI data by MOH IPC directorat e	3.2.1 Yes/No	3.2.1 Once	3.2.1 MOH/ IPC directorat e	3.2.1 Checking	3.2.1 Not available

Informati on system dep.	
dep.	
Droconts	
presents projects	
(either in-	
house, or	
ready to	
buy) with	
budget	
providing	
3.2.1.3	
Agreemen	
t between 100 100 100 100 100 100 100 100 100 10	
IPC III III III III III III III III III	
directorat	
e and See above	
Informatio	
n system	
dep. to	
choose choose	
one one	
system	
3.2.1.4	
Apply the Apply the	
system in See above	
the land and the l	
hospitals	
3.2.1.5	
Apply the Apply the	
Digital Digi	
HAI/AMR See above	
System See above	
for	
National National	
data	

	collection from individual hospitals to central IPC directorat e										
3.2.2 To Shift all hospital informatio n system from paper to digital mode.	3.2.2.1 MOH recomme nds from hospitals to have an electronic patient and data informatio n system.	3.2.2.1.1 Letter from MOH to hospitals	-	-	3.2.2.1.1 Letter is sent		3.2.2.1.1 Yes /No	3.2.2.1.1 Once	3.2.2.1.1 MOH	3.2.2.1.1 Checking	3.2.2.1.1 Partially available in some hospitals
	3.2.2.2 Increase the number of hospitals that have digitalized their medical informatio n system	-	-	-	3.2.2.2 % hospitals with digital informatio n system	3.2.2.2 improving all aspects of patient care	3.2.2.2 Number of hospitals with electronic medical records/T otal number of hospitals *100	3.2.2.2 Once/year	3.2.2.2 Hospitals	3.2.2.2 Checking	3.2.2.2 Partially available in some hospitals
3.2.3 Connect Infection Control	3.2.3.1 Connect Infection Control departme	3.2.3.1.1 Letter from MOH to ask all	-	-	3.2.3.1.1 % IC dep connected with	3.2.3.1.1 proper HAI surveillan ce and	3.2.3.1.1 Number of-IC dep connected with	3.2.3.1.1 Once/year	3.2.3.1.1 Hospitals/ IPC directorat e / MOH	3.2.3.1.1 Checking	3.2.3.1.1 partially available in some hospitals

Departme nts to electronic medical records	nts in all hospitals with the Heath informatio n system and Laborator y informatio n system at the hospital level	hospitals to connect IC dep. with HIS and LIS at the hospital level		HIS/LIS in hospitals	preventio n thus improving patient care	HIS/LIS in hospitals /Total number of hospitals *100				
		3.2.3.1.2 Decree from MOH to all hospitals who have an electronic Health informatio n system to link their Lab AMR data to the Hospital system	-	3.2.3.1.2 % of IC dep connected with electronic AMR data in hospitals	3.2.3.1.2 Better HAI and AMR surveillan ce on the national level	3.2.3.1.2 number of IC dep connected with electronic AMR data in hospitals /Total number of hospitals *100	3.2.3.1.2 Once/year	3.2.3.1.2 Hospitals/ IPC directorat e/ MOH	3.2.3.1.2 Checking	3.2.3.1.2 Not available
	3.2.3.2 Develop electronic			3.2.3.2 electronic outbreak	3.2.3.2 Better managem	3.2.3.2 Yes/No	3.2.3.2 Once	3.2.3.2 Hospitals	3.2.3.2 Checking	3.2.3.2 Not available

outbreak reporting system for data entry, analysis and generatin g reports.			reporting system available	ent and control of outbreaks					
3.2.3.3 Develop electronic reporting system for data entry, analysis and generatin g hospital/n ational reports for bundles of care.			Bundles data collection, analysis and generatin g hospitals /national reports is done through. an electronic advanced system Yes/No	3.2.3.3 Better monitorin g and auditing of bundles of care	3.2.3.3 Yes/No	3.2.3.3 Once	3.2.3.3 Hospitals /IPC directorat e	3.2.3.3 Checking	3.2.3.3 Not available
3.2.3.4 Develop electronic infection preventio n and control system capable of	3.2.3.4.1 Electronic system has a function for self diagnosis of HAIs and		3.2.3.4. % of HAIs and outbreaks in hospitals are automatic ally	3.2.3.4.1 Better tracking and managem ent of HAIs and outbreaks and	3.2.3.4.1 Number HAIs and outbreaks detected/ Total number of HAIs and outbreaks	3.2.3.4.1 Once/year	3.2.3.4.1 IPC directorat e/ Hospitals	3.2.3.4.1 Checking	3.2.3.4.1 Not available

	self- identifyin g HAI and detecting outbreaks	automatic ally detects outbreak			detected by the IT system	preventin g spread					
Strategic O		nhance IPC e	ducation in al	l health- relat	ted specialtie	s and among	HCW.				<u>.</u>
3.3.1 Include IPC as a core element in education and training of healthcar e profession als	3.3.1.1 Include infection control in undergrad uate curricula for health care students (medical, nursing, dental and pharmace utical).	3.3.1.1.1 Prepare module about IPC and AMR to be included in all health specialties in universitie s/PAAET.	-	-	3.3.1.1.1 Module is available	3.3.1.1.1 To provide all health specialties with IPC material to be taught in undergrad uate and post graduate education	3.3.1.1.1 Yes/No	3.3.1.1.1 Once	3.3.1.1.1 IPC Directorat e	3.3.1.1.1 Checking	3.3.1.1.1 Not available
		3.3.1.1.2 MOH letter to MOHE and PAAET to ask to include this modules into the undergrad s and graduate	-	-	3.3.1.1.2 Letters sent	3.3.1.1.2 To make it official and speed up the inclusion of these modules into the curricula	3.3.1.1.2 Yes/No	3.3.1.1.2 Once	3.3.1.1.2 IPC Directorat e	3.3.1.1.2 Checking	3.3.1.1.2 Not sent yet

		health students' curricula								
		3.3.1.1.3 Modules sent to Kuwait university / PAAET to include it in curricula		3.3.1.1.3 % health specialties that include IPC module	3.3.1.1.3 To incorporat e IPC training in all health specialties including dentistry	3.3.1.1.3 Health specialties programs that include IPC module/T otal health specialties programs X 100	3.3.1.1.3 Once/6 months until all programs do include IPC modules	3.3.1.1.3 Universiti es/PAAET	3.3.1.1.3 Data collection	3.3.1.1.3 Partially available
3.3.2 Enhance IPC education among undergrad uates and higher education health students	3.3.2.1 Include infection control in continuing education of healthcar e workers and mandator y prerequisi te for promotio n	3.3.2.1.1 Provide IPC courses with CE credits in universitie s, hospitals, MOH, PAAET PAAF Include CME Credit		3.3.2.1.1 Number of IPC courses/ year	3.3.2.1.1 To provide opportuni ty of IPC education among non-IPC HCW and to promote IPC as a job among students and HCW	3.3.2.1.1 Number/ year	3.3.2.1.1 Once/year	3.3.2.1.1 Universiti es/PAAET KIMS	3.2.2.1.1 Data collection	3.3.2.1.1 Not available

	10% IPC									
	3.3.2.1.2 Request that when CE credits for physicians , nurses, or pharmacis ts is needed, 10% of these credits should be about IPC.			3.3.2.1.2 % profession als have 10% of their CE credits are related to IPC	3.3.2.1.2 To provide IPC education to all HCW, and not to IPC profession als alone	3.3.2.1.2 Number of HC profession als that have CE related to IPC/Total number of profession als who are requested to have CE credits.	3.3.2.1.2 Once/year	3.3.2.1.2 Syndicates of HC profession als (MDs, nurses, pharmacis ts, dentists, physiothe rapists, technician s)	3.3.2.1.2 Data collection	3.3.2.1.2 Not available
	3.3.2.1.3 In hospitals, yearly and upon employm ent, mandator y attendanc e of IPC lectures	-	-	3.3.2.1.3 % hospitals that mandated that all employee get IPC training upon employm ent and yearly IPC lecture.	3.3.2.1.3 To update and remind and engage HCW with IPC	3.3.2.1.3 Number of hospitals that have this program /total number of hospitals	3.3.2.1.3 Once/year	3.3.2.1.3 Hospitals	3.3.2.1.3 Data collection	3.3.2.1.3 Partially available
3.3.2.2 Hygiene and IPC education provided	3.3.2.2.1 Require IPC education by drug	3.3.2.2.1.1 Request to include few topics about IPC	3.3.2.2.1.1 .1 Letter to be sent	3.3.2.2.1.1 .1 Yes/No		3.3.2.2.1.1 .1 Yes/No	3.3.2.2.1.1 .1 Once/conf erence	3.3.2.2.1.1 .1 conferenc e organizers	3.3.2.2.1.1 .1 Checking	3.3.2.2.1. 1.1 Not available

	outside formal teaching programs	companie s, hospitals, LTCFs, FAO, and medical/v eterinary associatio ns	in conferenc es agendas								
3.4.1 Develop a comprehe nsive program of vaccinatio n to protect HCWs from vaccine preventab le diseases	-	-	-	ions in HCWs	3.4.1 % of HCWs profession als that are vaccinate d	3.4.1 Prevent vaccine preventab le diseases	3.4.1 Number of HCW who are vaccinate d/Total number of HCW*100	3.4.1 Once/year	3.4.1 MOH Hospitals/ Public Health Directorat e	3.4.1 Checking	3.4.1 Partially available
3.4.2 Develop policies and procedure s to protect healthcar e workers from	-	-	-	-	3.4.2 Yes/No	3.4.2 Prevent occupatio nal exposure to infections among HCW	3.4.2 Yes/No	3.4.2 Once	3.4.2 Hospitals	3.4.2 Checking	3.4.2 Partially available

occupational exposure to infections. 3.4.3 Availability of Airborne Isolation Rooms in all hospitals	3.4.3.1 screening mapping and updating the recomme ndations according to the expected need according	-	-	-	3.4.3.1 Yes/No	3.4.3.1 proper managem ent of airborne infections and improving quality of care	3.4.3.1 Yes/No	3.4.3.1 Once	3.4.3.1 Hospitals	3.4.3.1 Checking	3.4.3.1 Partially available
3.4.4 Mandate to hospitals to provide airborne isolation rooms (AII) available as per the updated recomme ndations	to need	-	-	-	3.4.4 Percent hospitals have enough All rooms as per MOH recomme ndation	3.4.4 Prevent spread of airborne infections plus proper managem ent	3.4.4 Number of hospitals with enough All rooms /total number of hospitals*	3.4.4 Once/year	3.4.4 MOH Hospitals	3.4.4 Checking	3.4.4 Partially available
Strategic Ob	jective 3.5 A	dequate was	te manageme	ent in hospita	ls						
3.5.1	3.5.1.1	3.5.1.1.1	3.5.1.1.1.1	-	3.5.1.1.1.1		3.5.1.1.1.1 Yes/No	3.5.1.1.1.1 Once	3.5.1.1.1.1 MOH	3.5.1.1.1.1 Checking	3.5.1.1.1.

Put National Guidelines	National policies, strategies and plans for healthcar e waste managem ent is in place	Unify recomme ndations between KEPA and MOH	Create a committe e from MOH and KEPA that unifies the recomme ndation and the road map of all types of waste that are generated		Committe e members are nominate d				KEPA		Not available
			in healthcar e 3.5.1.1.1.2 The committe e reviews waste policies that are related to HCF and unifies recomme ndations and provides the proper ways of	-	3.5.1.1.1.2 Percent chapters that represent conflict between recomme ndations of the 2 ministries that have been reviewed and unified	3.5.1.1.1.2 Unify recomme ndation and thus proper methods of supervisio n	3.5.1.1.1.2 number of chapters edited /total number of chapters* 100	3.5.1.1.1.2 Once	3.5.1.1.1.2 MOH KEPA	3.5.1.1.1.2 Checking	3.5.1.1.1. 2 Not available

			supervisio n								
3.5.2 Supervisio n of waste disposal from public hospitals	3.5.2.1 Create a committe e for the supervisio n of waste disposal from public hospitals	-	-	-	3.5.2.1 Available and ongoing						
3.5.3 Supervisio n of waste disposal in private hospitals	3.5.3.1 Create a committe e for the supervisio n of waste disposal in private hospitals	-	-	-	3.5.3.1 Yes/No	3.5.3.1 overseein g proper waste disposal	3.5.3.1 Yes/No	3.5.3.1 Once	3.5.3.1 private hospitals	3.5.3.1 Checking	3.5.3.1 Not available
3.5.4 Waste policies are being audited by KEPA	3.5.4.1 Organizati on of supervisio n of waste disposal inside and from private hospitals	-	-	-	3.5.4.1 % Public hospitals are being audited for waste disposal.	3.5.4.1 Ensure proper waste managem ent and preventio n of environm ent contamin ation	3.5.4.1 Number of audited hospitals/ total number of hospitals *100	3.5.4.1 Once/year	3.5.4.1 KEPA	3.5.4.1 Checking	3.5.4.1 Not available
	3.5.4.2		-	-	3.5.4.2		3.5.4.2 Yes/No	3.5.4.2 Once	3.5.4.2 KEPA	3.5.4.2 Checking	3.5.4.2

Strategic O	Designate 1 unit at KEPA to oversee the waste in hospitals	o enhance in	fection preve	ntion and cor	% Private hospitals where waste managem ent is being supervise d	s in veterinar	y and agricul	ture settings			Not available
3.6.1 Availabilit y of IPC guidelines in veterinary practices	3.6.1.1 Implementation of OIE guidelines regarding IPC in veterinary field	3.6.1.1.1 Committe e in PAAF that regularly checks updates in OIE guidelines regarding IPC in veterinary field, and updates the guidelines accordingl y	-	-	3.6.1.1.1 Committe e is formed	3.6.1.1.1 To have up-to-date guidelines and to have local responsibl e bodies involved rather than internatio nal NGOs alone.	3.6.1.1.1 Guidelines availabilit y and Update at least once/3 years	3.6.1.1.1 Once/year	3.6.1.1.1 PAAF	3.6.1.1.1 Checking the guidelines	3.6.1.1.1 Guideline s available, need improved governanc e
		3.6.1.1.2 Overseein g the implemen tation of national IPC guidelines	3.6.1.1.2 Regular audits to farms	3.6.1.1.2 Put a yearly audit plan	3.6.1.1.2 % Farms that are being audited yearly for IPC measures.	3.6.1.1.2 To check the applicatio n of IPC guidance in farms	3.6.1.1.2 Number of farms that are being audited yearly for IPC	3.6.1.1.2 Once/year	3.6.1.1.2 Farms	3.6.1.1.2 Data collection	3.6.1.1.2 Not performe d

		in Vet field and OIE good practices					measures/ Total number of farms X 100				
3.6.2 Include IPC in licensing of Vets	3.6.2.1 Include questions about OIE IPC guidelines and vaccinatio ns in all vets licensing exams or interviews .	3.6.2.1.1 Prepare a set of questions related to IPC in Vets to be included in licensing exams and interviews .	-	-	3.6.2.1.1 Availabilit y of IPC questions in licensing exams (Yes/No)	3.6.2.1.1T o promote revision of IPC practices in preparatio n for pre- employm ent interviews and exams	3.6.2.1.1 Yes/No	3.6.2.1.1 Yearly	3.6.2.1.1 PAAF Exam material	3.6.2.1.1 Checking	3.6.2.1.1 Not available
3.6.3 Include IPC in Continuous education of Vets and profession als in Agricultur e	3.6.3.1 Prepare a slide bank related to IPC in vet and agricultur e, to be given to drug companie s that usually do the continuou s	-	-	-	3.6.3.1 Yes/No	3.6.3.1 Improve IPC knowledg e among profession als in vet/agricu Iture fields	3.6.3.1 Yes/No	3.6.3.1 Once	3.6.3.1 PAAF	3.6.3.1 Checking	3.6.3.1 Not available

	education activities.										
	3.6.3.2 Mandate that all drug company events or seminars include a specific number of slides related to IPC and AMR.	-	-	-	3.6.3.2 Mandate available or not	3.6.3.2 To oblige drug companie s that are almost the only provider of CE, to include IPC in their presentati ons.	3.6.3.2 Mandate issued: Yes/No	3.6.3.2 Once	3.6.3.2 PAAF	3.6.3.2 Checking	3.6.3.2 Not available
3.6.4 Include hygiene and infection preventio n and control as core content in training of veterinary profession als.	3.6.4.1 Close communic ation between. FAO and PAAF to put a joint yearly plan of education activities related to IPC.	-	-	-	3.6.3.2 Joint education plan between FAO and PAAF that includes IPC- related training	3.6.3.2 To provide CE by nonprofit organizati ons to eliminate the possibility of commerci al bias while teaching IPC	3.6.3.2 Plan available: Yes/No	3.6.3.2 Once/year	3.6.3.2 PAAF FAO	3.6.3.2 Checking	3.6.3.2 Not available
	3.6.4.2	3.6.4.2.1	-	-	3.6.4.2.1	3.6.4.2.1	3.6.4.2.1	3.6.4.2.1 Once	3.6.4.2.1	3.6.4.2.1 Checking	3.6.4.2.1

Include	IPC credits		Mandate	То	Mandate	Vet	Not
hygiene	mandator		available	enhance	issued	associatio	available
and	y for		or not	IPC	yes/no	n	
infection	promotio			education			
preventio	n in Vet.			and			
n and	Associatio			update			
control as	ns			among			
mandator				vets.			
У							
prerequisi							
te for							
promotio							
n of for							
the							
veterinary							
workers.							

Annex 5

A. Antimicrobial Use Strategic Plan

Activity	Sub-activity	Sub-sub-activity	Date	Milestone for strategic objective
	governance of antibiotic axis p	illar and follow up of the activit	ies of this pillar	
4.1.1 Identify who will be the				
ABX arm pillar focal point in			4.1.1	4.1
each ministry/authority and			6 months	2 years
assign the TOR				
4.1.2 Prepare clear TOR of			4.1.2	
the focal person for ABX in			2 months	
each ministry/ authority				
4.1.3 Assign the job to a				
defined person/position in			4.1.3	
each ministry/ authority, and			3 months	
nomination by the director				
of each ministry.				
4.1.4 Put TOR and			4.1.4	
reimbursement plan for the			3 months	
different tasks.				
4.1.5 Nominate the			4.1.5	
stakeholders for the task			4 months	
force of the pillar				
4.2.1 Put clear criteria for	uninterrupted access to high-q	uality antimicrobial medicines		
standards of quality safety			4.2.1 System is in place	4.2 Available
and efficiency of ABX			4.2.1 System is in place	4.2 Available
4.2.2 Put a system of				
controlling that all available			4.2.2 Already available,	
ABX in the country should			Only ABX that are licensed	
meet the international			by WHO, EMEA or FDA,	
standards listed above			are allowed in the market	

				Т
otherwise they are not				
allowed to be in the market.				
4.2.3 All ABX should be				
tested by a system that			4.2.3 Available	
identifies counterfeit prior to			4.2.3 Available	
licensing.				
4.2.4 Put regulations that				
prohibit the sale of			4.2.4 Available	
counterfeit products				
4.2.5 The system that				
controls quality assurance of				
ABX has the authority to			4.2.5 available	
stop counterfeit products			4.2.5 available	
from being licensed and				
marketed				
Strategic Objective 4.3 Develo	p and enforce legislation and re	egulations on prescription and c	lispensing of medicine including	g antibiotics
4.3.1 Develop and				
implement national essential	4.3.1.1 Finalize the essential		4244	4.2
medicine list guided by the	medicine list that is		4.3.1.1 6 months	4.3
WHO Model List of Essential	being prepared at MOH		6 months	4 years
Medicines				
4.3.2 Mandate that every			4.3.2	
institution has an essential				
medicine list			6 months	
4.3.3 Each institution has an	4.3.3.1 Mandate from MOH			
essential medicine list and	that once the			
this is checked in	national essential medicine		4.3.3.1	
accreditation standards	list is available,		4.3.3.1 8 months	
include checking the	all hospitals including public		0 1110111115	
availability of the essential	and private hospitals should			
medicine list	formulate their own list			
	4.3.3.2 Include			
	the availability of essential	4.3.3.2.1 Private hospital	4.3.3.2.1	
	medicine list into	renewal license	8 months	
	the license renewal			

	conditions of private hospitals	checklist should include the availability of essential medicine list		
4.3.4 Re imbursement plan is based on essential medicine list in public and private hospitals			4.3.4 8 months	
	MS program in primary healtho	are		
4.4.1 Create a technical group that puts national ABX use guidelines for outpatient care based on International guidelines and National AMR profiles: can be a subcommittee from the national task force for the ABX pillar for proper use of ABX	4.4.1.1 Put TOR and renewal conditions of the focal group of stakeholders		4.4.1.1 3 months	4.4 5 years
	4.4.1.2 Assign the members of this committee		4.4.1.2 3 months	
	4.4.1.3 Put TOR and official renewal time for the guidelines committee		4.4.1.3 3 months	
4.4.2 Guidelines in primary healthcare are available and are regularly updated			4.4.2 6 months	
	4.4.2.1 Training workshop about ABX guidelines in primary healthcare		4.4.2.1 5 years	
4.4.3 Apply AMS in primary healthcare: Monitoring ABX consumption and Link prescription to diagnoses	4.4.3.1 Project ABX consumption documentation and surveillance in primary healthcare	4.4.3.1.1 Put a proposal for this project	4.4.3.1.1 8 months	

	4.4.3.2 Start collecting data and link it to diagnosis 4.4.3.3 Data analysis and feedback to prescribers 4.4.3.4 Employ more pharmacists in primary healthcare 4.4.3.5 Improve manpower with professionals for the	4.4.3.5.1 Train available	4.4.3.2 10 months 4.4.3.3 1 year 4.4.3.4 1 year	
	AMS project in primary healthcare	pharmacists on AMS and clinical pharmacy principles	18 months	
		4.4.3.5.2 Employ new clinical pharmacist specialized in AMS and 1 full time clinical microbiologist in primary healthcare directorate	4.4.3.5.2 18 months	
4.4.4 Update the digital system that is available in primary healthcare to integrate the ABX guidelines into the electronic system in the form of pop-ups			4.4.4 2 years	
Strategic Objective 4.5 Apply	AMS In Public and Private hospi	tals		
4.5.1 Improve and measure appropriate use of antimicrobial agents in healthcare	4.5.1.1 Nominate who will be the AMS focal person (Could be the ABX pillar focal person)		4.5.1.1 3 months	4.5 5 years

4.5.2 Identify the National AMS Task force (Could be a subcommittee of the ABX pillar Task force) the National AMS Technical team	4.5.2.1 AMS technical group is assigned (subgroup from the ABX technical committee. +other specialists like IT, etc)		4.5.2.1 3 months	
	4.5.2.2 TOR of the technical groups are well defined They include: -guidelines for inpatient and outpatient careput or endorse diagnostic pathways, treatment and national AMS targets		4.5.2.2 3 months	
4.5.3 Put national guidelines either <i>de novo</i> or Endorse existing international guidelines with regular updates	4.5.3.1 MOH puts national guidelines for common community acquired infections (CAI)		4.5.3.1 6 months	
	4.5.3.2 Hospitals put institutional guidelines for treatment of Healthcare associated (HAI) according to their local epidemiology, with the help of taskforce members	4.5.3.2.1 Mandate for hospitals to issue HAI treatment guidelines based on local epidemiology and with the help of taskforce members	4.5.3.2.1 6 months	
		4.5.3.2.2 Organize the work of taskforce members with the different hospitals to assist in putting their own HAI guidelines 4.5.3.2.3	4.5.3.2.2 8 months 4.5.3.2.3	

		HAI treatment guidelines should be reviewed every year	once/ year
	4.5.3.3 Dissemination and implementation of these treatment guidelines in hospital practice.	4.5.3.3.1 Workshops for discussion and endorsement of these guidelines with practicing physicians all over the country	4.5.3.3.1 1 year
		4.5.3.3.2 Distribution of educational material in the form of booklets, brochures, posters, &pocket cards, or equivalent softcopies available on mobile phones	4.5.3.3.2 1 year
		4.5.3.3.3 Post the guidelines on the MOH website and integrate them into the intranet of each hospital	4.5.3.3.3 1yr, 3 months
		4.5.3.3.4 Integrate the guidelines into the electronic medical record, in the form of pop-ups	4.5.3.3.4 2 years
4.5.4 Make AMS programs mandatory in each hospital (presence of AMS team/committee)	4.5.4.1 Mandate by MOH for this issue		4.5.4.1 1 y 6 months
	4.5.4.2 Build AMS manpower	4.5.4.2.1 Train microbiologists, clinical pharmacists and Infectious	4.5.4.2.1 1 y 6 months

		Disease specialists on AMS		
Stratogic Objective 4.6 Quality	control of antimicrobials used	through national workshops		
Strategic Objective 4.6 Quanty	Control of antimicropiais used	in annuals and plants		
4.6 ABX used in the				
agriculture/ veterinary				
are under the same control				
as those used in humans			4.6 Already available	4.6 available
Pharmaceutical and Herbal				
Medicines Registration and				
Control Administration/The				
Dug Inspection Control				
	t the use of critically important	ABX to human health in the ve	terinary field	
4.7.1				
Establish the list of			4.7.1	4.7
critically important			3 months	2 years
antibiotics for humans in the				,
animal sector.				
4.7.2			472	
Determine from this list which antibiotics should be			4.7.2	
restricted for use to humans			3 months	
4.7.3				
Issue a decree that the				
critically important ABX to				
human health are not to be			4.7.3	
used in animals whether in			6 months	
individual treatment or in				
herd therapy				
4.7.4	4.7.4.1			
Prohibit the importation of	Decree		4.7.4.1	
these critically	that prohibits the		6 months	
important ABX for	importation and sale of		o months	
vet use	these ABX in the vet field			
Strategic Objective 4.8 Organi	ze the use of antimicrobial age	nts in terrestrial and aquatic ani	mals and agriculture.	

4.8.1 Put (or endorse) guidelines regarding the use of ABX in animals and agriculture			4.8.1 8 months	4.8 5 years
4.8.2 Disseminate these guidelines among veterinarians	4.8.2.1 Booklets/ Brochures/app that summarize the key points in these guidelines.		4.8.1.1 18 months	
	4.8.2.1 Regular workshops for veterinary specialists regarding the application of these guidelines		4.8.2.1 1 year	
4.8.3 Restrict selling ABX for animal and agriculture health to exclusively prescriptions by vets	4.8.3.1 Decree that vet pharmacies should dispense ABX only according to vet prescriptions, and should submit these prescriptions with the evidence of selling to PAAF.		4.8.3.1 2 years	
4.8.4 Control the use of ABX in animals	4.8.4.1. Detection of the ABX residues in milk, meat, poultry and eggs		4.8.4.1. Already available	4.8.4.1. Already available
Strategic Objective 4.9 Capaci 4.9.1 Provide training	ty building among vets in antim	icrobial use and AMR		
courses for vets for ABX use according to guidelines and provide certificate.			4.9.1 2 years	4.9 4 years
4.9.2 Study baseline consumption imported antibiotics in this field	4.9.2.1 Pharmaceutical Dep in PAAF Committee that will study and document this consumption		4.9.2.1 2 years	

and compa	re it to		
internation	ial benchmark		

B. Antimicrobial Use Monitoring Plan

Activity	Sub-activity	Sub-sub- activity	Indicator	Purpose	Calculation	Frequency	Data source	Method	Baseline
Strategic Obje	ctive 4.1 Ensure	governance of	antibiotic axis p	illar and follow	up of the activit	ies of this pillar			
4.1.1 Identify									
who will be									
the ABX arm									
pillar focal			4.1.1						
point in each			available						
ministry			avallable						
/authority									
and assign									
the TOR									
4.1.2									
Prepare				4.1.2 proper					
clear TOR of				organization			4.1.2		
the focal			4.1.2	and follow-	4.1.2	4.1.2	AMR	4.1.2	4.1.2
person for			Yes/No	up on the	Yes/No	Once	committee	Checking	Not available
ABX in each				work in this			Committee		
ministry				pillar					
authority									
4.1.3			4.1.3	4.1.3 proper	4.1.3		4.1.3		
Assign the			Proportion	monitoring	number of		AMR		
job to a			of	and	ministries /	4.1.3 once	Inter-	4.1.3	4.1.3
defined			ministries/au	surveillance	authorities	4.1.3 0116	ministerial	checking	not available
person/posit			thorities	on the work	with		executive		
ion in each			where ABX	in this pillar	assigned		committee		

ministry authority, and nomination by the director of each ministry/ authority.			arm focal person is assigned		focal persons/tota I number of concerned ministries/au thorities*10 0				
4.1.4 Put TOR And reimbursem ent plan for the different tasks.			4.1.4 Yes/No	4.1.4 proper organization and follow-up on the work of this axis	4.1.4 Yes/No	4.1.4 Once	4.1.4 MOH NMCG/ Focal person	4.1.4 Checking	4.1.4 Not available
4.1.5 Nominate the stakeholders for the task force of the pillar			4.1.5 ABX use task force group is formed Yes/No	4.1.5 assign responsibiliti es to specific persons regarding the work in this pillar	4.1.5 Yes/No	4.1.5 Once	4.1.5 MOH PAAF KEPA PAFN	4.1.5 Checking	4.1.5 Not available
	ctive 4.2 Ensure	uninterrupted a	access to high-q	uality antimicro	bial medicines				
4.2.1 Put clear criteria for standards of quality safety and efficiency of ABX			4.2.1 System is in place						
4.2.2 Put a system of controlling			4.2.2 Already available						267

that all					
available					
ABX in the					
country					
should					
meet the					
international					
standards					
listed above					
otherwise					
they are not					
allowed to					
be in the					
market.					
4.2.3 All ABX					
should be					
tested by a		4226			
system that		4.2.3 System			
identifies		already in			
counterfeit		place			
prior to					
licensing.					
4.2.4 Put					
regulations		4.2.4 System			
that prohibit		already in			
the sale of		place			
counterfeit		piace			
products					
4.2.5 The					
system that					
controls		4.2.5 System			
quality		already			
assurance of		In place			
ABX has the					
authority to					
stop					

	ctive 4.3 Develo	p and enforce le	egislation and re	egulations on pr	escription and c	dispensing of m	edicine includinį	g antibiotics	
4.3.1 Develop and implement national essential medicine list guided by the WHO Model List of Essential Medicines	4.3.1.1 Finalize the essential medicine list that is being prepared at MOH		4.3.1.1 List is issued Yes/No	4.3.1.1 regulate prescription and dispensing of medications including antimicrobial s	4.3.1.1 Yes/No	4.3.1.1 Once	4.3.1.1 MOH Central Medical Stores Directorate	4.3.1.1 Checking	4.3.1.1 task in progress
4.3.2 Mandate that every institution has an essential medicine list			4.3.2 Yes/No	4.3.2 regulate prescription and dispensing of medications including antimicrobial s on the facility level	4.3.2 Yes/No	4.3.2 Once	4.3.2 MOH Undersecret ary general ABX focal person	4.3.2 Checking	4.3.2 Not available
4.3.3 Each institution has an essential medicine list and this is checked in accreditation standards	4.3.3.1 Mandate from MOH that once the national essential medicine list is available,		4.3.3.1 Yes/No	4.3.3.1 put a legal frame for the prescription and dispensing of medicines including	4.3.3.1 Yes/No	4.3.3.1 Once	4.3.3.1 MOH Undersecret ary general ABX focal person	4.3.3.1 Checking	4.3.3.1 Not available

include checking the availability of the essential medicine list	all hospitals including public and private hospitals should formulate their own list			antimicrobial s					
	4.3.3.2 Include the availability of essential medicine list into the license renewal conditions of private hospitals	4.3.3.2.1 Private hospital renewal license checklist should include the availability of essential medicine list	4.3.3.2.1 Licensing checklist includes the item of essential medicine list or not	4.3.3.2.1 having an essential list is an obligation and represents a standard of good practice of medicine	4.3.3.2.1 Yes/No	4.3.3.2.1 Once	4.3.3.2.1 MOH/ Medical Licensing Dep	4.3.3.2.1 Checking	4.3.3.2.1 Not available
4.3.4 Re imbursemen t plan is based on essential medicine list in public and private hospitals			4.3.4 % of reimbursed ABX that are listed in the essential medicine list	4.3.4 making essential list an obligation as a standard of good practice of medicine	4.3.4 number of reimbursed antibiotics/t otal number of available antibiotics*1 00	4.3.4 once/year	4.3.4 MOH	4.3.4 data collection	4.3.4 Not available
4.4.1 Create	ctive 4.4 Start A	MS program in	primary healtho	are	l	l	4.4.1.1		
a technical group that puts national ABX use guidelines	4.4.1.1 Put TOR and renewal conditions of the		4.4.1.1 TOR put Yes or no	4.4.1.1 organize the AMS work that this taskforce is	4.4.1.1 Yes/No	4.4.1.1 Once	MOH/ National Committee of proper ABX use	4.4.1.1 Checking	4.4.1.1 Not available

for outpatient care based on International guidelines and National AMR profiles: can be a subcommitte e from the national task force for the ABX pillar for proper use	focal group of stakeholders		responsible					
of ABX	4.4.1.2 Assign the members of this committee	4.4.1.2 TOR put Yes or no	4.4.1.2 oversee AMS work in this sector	4.4.1.2 Yes/No	4.4.1.2 Once	4.4.1.2 MOH National Committee of proper ABX use.	4.4.1.2 Checking	4.4.1.2 Not available
	4.4.1.3 Put TOR and official renewal time for the guidelines committee	4.4.1.3 TOR put Yes or no	4.4.1.3 oversee guideline developmen t	4.4.1.3 Yes/No	4.4.1.3 Once	4.4.1.3 MOH National Committee of proper ABX use.	4.4.1.3 Checking	4.4.1.3 Not available
4.4.2 Guidelines in primary		4.4.2 Guidelines available	4.4.2 ensure judicious	4.4.2 Yes/No	4.4.2 Once	4.4.2 Primary Healthcare	4.4.2 Checking	4.4.2 Not available

healthcare are available and are regularly updated			Yes or no	antimicrobial use in primary healthcare			Directorate/ technical committee/ National Committee of proper ABX use.		
	4.4.2.1 Training workshop about ABX guidelines in primary healthcare		4.4.2.1 Proportion of primary healthcare physician who attend every year	4.4.2.1 primary care physicians rationally prescribe antimicrobial s	4.4.2.1 number of primary care physicians who attend the training workshops/t otal number of primary healthcare physicians *100	4.4.2.1 Once/year	4.4.2.1 Primary healthcare/ technical committee	4.4.2.1 Checking	4.4.2.1 Not available
4.4.3 Apply AMS in primary healthcare: Monitoring ABX consumption and Link prescription to diagnoses	4.4.3.1 Project ABX consumption documentati on and surveillance in primary healthcare	4.4.3.1.1 Put a proposal for this project	4.4.3.1.1 Project is put	4.4.3.1.1 follow-up on managemen t of antimicrobial s in primary healthcare	4.4.3.1.1 Yes/No	4.4.3.1.1 Once	4.4.3.1.1 Primary healthcare and AMR committee	4.4.3.1.1 Checking	4.4.3.1.1 Not available
J	4.4.3.2 Start collecting data and link it to diagnosis		4.4.3.2 Proportion of primary healthcare centers that are feeding	4.4.3.2 assessment of the practice in these centers and	4.4.3.2 Number of primary healthcare centers that are feeding	4.4.3.2 Once/year	4.4.3.2 National Committee for ABX use /Primary	4.4.3.2 Data Collection	4.4.3.2 Not available

		data into the project	thus ability to give feedback to prescribers	data into the project/Total number of primary healthcare centers *100		healthcare Directorate		
4.4.3.3 Data analysis and feedback to prescribers		4.4.3.3 Proportion of primary healthcare centers that receive annual report with feedback about ABX prescription	4.4.3.3 improvemen t of practice and thus standard of care and proper application of AMS in this sector	4.4.3.3 Number of primary healthcare centers that receive annual report with feedback about ABX prescription/ Total number of primary Health care centers *100	4.4.3.3 Once/year	4.4.3.3 primary healthcare directorate	4.4.3.3 Data collection	4.4.3.3 Not available
4.4.3.4 Employ more pharmacists in primary healthcare		4.4.3.4 Number of pharmacists that are employed	4.4.3.4 Surveil, monitor on the proper use of antibiotics in primary healthcare	4.4.3.4 Yes/NO	4.4.3.4 Once	4.4.3.4 Primary healthcare directorate/ Pharmaceuti cal Services Directorate	4.4.3.4 Checking	4.4.3.4 Not available
4.4.3.5 Improve manpower with professionals for the AMS	4.4.3.5.1 Train available pharmacists on AMS and clinical	4.4.3.5.1 Number of pharmacists in primary healthcare who have a	4.4.3.5.1 oversee AMS practices by qualified pharmacists	4.4.3.5.1 Yes/No	4.4.3.5.1 Once	4.4.3.5.1 Primary healthcare directorate /Pharmaceut	4.4.3.5.1 Checking	4.4.3.5.1 Not available

	project in primary healthcare	pharmacy principles	certificate in AMS training				ical Services Directorate		
		4.4.3.5.2 Employ new clinical pharmacist specialized in AMS and 1 full time clinical microbiologi st in primary healthcare directorate	4.4.3.5.2 Clinical pharmacists are employed, Yes/No	4.4.3.5.2 oversee AMS practices in primary healthcare by qualified professionals (pharmacist and microbiologi st)	4.4.3.5.2 Yes/No	4.4.3.5.2 Once	4.4.3.5.2 Primary healthcare directorate /Pharmaceut ical Services Directorate	4.4.3.5.2 Checking	4.4.3.5.2 Not available
4.4.4 Update the digital system that is available in primary healthcare to integrate the ABX guidelines into the electronic system in the form of pop-ups			4.4.4 updated electronic system available or not	4.4.4 facilitate AMS application through user friendly software	4.4.4 Yes/No	4.4.4 Once	4.4.4 Primary Heath care directorate/ /Information System Dep.	4.4.4 Checking	4.4.4 Not available
Strategic Obje	ctive 4.5 Apply	AMS In Public a	nd Private hospi	tals					
4.5.1	4.5.1.1		4.5.1.1	4.5.1.1	4.5.1.1 Yes/No	4.5.1.1 Once	4.5.1.1 MOH	4.5.1.1 Checking	4.5.1.1 Not available

Improve and measure appropriate use of antimicrobial agents in healthcare	Nominate who will be the AMS focal person (Could be the ABX pillar focal person)	AMS focal person is nominated	oversee AMS work in hospitals					
4.5.2 Identify the National AMS Task force (Could be a subcommitte e of the ABX pillar Task force) the National AMS Technical team	4.5.2.1 AMS technical group is assigned (subgroup from the ABX technical committee. +other specialists like IT, etc)	4.5.2.1 Task force members are nominated Yes/No	4.5.2.1 oversee AMS work in hospitals	4.5.2.1 Yes/No	4.5.2.1 Once	4.5.2.1 MOH	4.5.2.1 Checking	4.5.2.1 Not available
	4.5.2.2 TOR of the technical groups are well defined They include: -guidelines for inpatient and outpatient care.	4.5.2.2 TOR put	4.5.2.2 oversee AMS work in hospitals	4.5.2.2 Yes/No	4.5.2.2 Once	4.5.2.2 MOH	4.5.2.2 Checking	4.5.2.2 Not available

	-put or endorse diagnostic pathways, treatment and national AMS targets								
4.5.3 Put national guidelines either de novo or Endorse existing international guidelines with regular updates	4.5.3.1 MOH puts national guidelines for common community acquired infections (CAI)		4.5.3.1 CAI guidelines developed Yes/No	4.5.3.1 ensure proper prescribing and use of antimicrobial s	4.5.3.1 Yes/No	4.5.3.1 Once	4.5.3.1 AMS Task force Group	4.5.3.1 Checking	4.5.3.1 Not available
	Hospitals put institutional guidelines for treatment of Healthcare associated infections (HAI) according to their local epidemiolog y, with the help of taskforce members	4.5.3.2.1 Mandate for hospitals to issue HAI treatment guidelines based on local epidemiolog y and with the help of taskforce members	4.5.3.2.1 Mandate issued Yes/No	4.5.3.2.1 putting HAI guideline is obligatory	4.5.3.2.1 Yes/No	4.5.3.2.1 Once	4.5.3.2.1 MOH/task force	4.5.3.2.1 Checking	4.5.3.2.1 Not available
		4.5.3.2.2	4.5.3.2.2	4.5.3.2.2	4.5.3.2.2	4.5.3.2.2	4.5.3.2.2	4.5.3.2.2	4.5.3.2.2

	Organize the work of taskforce members with the different hospitals to assist in putting their own HAI guidelines	Proportion of hospitals that have HAI guidelines based on their own epidemiolog y.	to see if hospitals abide by developing institutional HAI guidelines	number of hospitals with HAI guidelines/T otal number of hospitals*10 0	Once/year	MOH/ Hospitals	data collection	Partially available in some hospitals
	4.5.3.2.3 HAI treatment guidelines should be reviewed every year	4.5.3.2.3 Proportion of hospitals that have guidelines updated	4.5.3.2.3 continuous follow-up and reduction of inappropriat e antibiotic use	4.5.3.2.3 number of hospitals with HAI guidelines updated/Tot al number of hospitals*10	4.5.3.2.3 Once/year	4.5.3.2.3 MOH/ Hospitals	4.5.3.2.3 data collection	4.5.3.2.3 Partially available in some hospitals
4.5.3.3 Disseminatio n and implementat ion of these treatment guidelines in hospital practice.	4.5.3.3.1 Workshops for discussion and endorsemen t of these guidelines with practicing physicians all over the country	4.5.3.3.1 Number of regions where guidelines workshops have been performed	4.5.3.3.1 proper application of guidelines recommend ations	4.5.3.3.1 Yes/No	4.5.3.3.1 Once/year	4.5.3.3.1 MOH	4.5.3.3.1 Checking	4.5.3.3.1 Not available

4.5.3.3.2 Distribution of educational material in the form of booklets, brochures, posters, &pocket cards, or equivalent softcopies available on mobile phones	4.5.3.3.2 Proportion of hospitals that have the softcopies available on the mobile phones of staff	4.5.3.3.2 facilitate disseminatio n and implementat ion of these treatment guidelines in hospital practice	4.5.3.3.2 Number of hospitals that have the softcopies available on the mobile phones of staff/Total number of hospitals*10 0	4.5.3.3.2 Once/year	4.5.3.3.2 MOH/Hospit als	4.5.3.3.2 data collection	4.5.3.3.2 not available
4.5.3.3.3 Post the guidelines on the MOH website and integrate them into the intranet of each hospital	4.5.3.3.3 Proportion of hospitals that have the guidelines posted on their intranet	4.5.3.3.3 facilitate access to these treatment guidelines	4.5.3.3.3 Number of hospitals that have the guidelines posted on their intranet/Tot al number of hospitals*10	4.5.3.3.3 Once/year	4.5.3.3.3 MOH/Hospit als	4.5.3.3.3 data collection	4.5.3.3.3 not available
4.5.3.3.4 Integrate the guidelines into the electronic medical record, in	4.5.3.3.4 Proportion of hospitals that have guidelines recommend ations integrated in	4.5.3.3.4 remind prescribers on a regular basis of recommend ations and facilitate	4.5.3.3.4 Number of hospitals that have guidelines recommend ations integrated in	4.5.3.3.4 Once/year	4.5.3.3.4 Hospitals	4.5.3.3.4 data collection	4.5.3.3.4 Not available

		the form of pop-ups	the electronic medical records system	implementat ion of AMS at the bedside	the electronic medical records system /Total number of hospitals*10 0				
4.5.4 Make AMS programs mandatory in each hospital (presence of AMS team/ committee)	4.5.4.1 Mandate by MOH for this issue		4.5.4.1 Mandate issued Yes/No	4.5.4.1 standardize the application of AMS in hospitals and make it obligatory	4.5.4.1 Yes/No	4.5.4.1 Once	4.5.4.1 MOH	4.5.4.1 Checking	4.5.4.1 Not available
	4.5.4.2 Build AMS manpower	4.5.4.2.1 Train microbiologi sts, clinical pharmacists and Infectious Disease specialists on AMS through national workshops	4.5.4.2.1 Percent microbiologi sts and clinical pharmacists that have attended at least 1 AMS workshop	4.5.4.2.1 build a strong AMS workforce for better implementat ion of AMS standards in hospitals	4.5.4.2.1 Number of microbiologi sts and clinical pharmacists that have attended at least 1 AMS workshop/T otal number of these professionals *100	4.5.4.2.1 Once/year	4.5.4.2.1 Task force	4.5.4.2.1 Data collection	4.5.4.2.1 Not available
Strategic Obje	ctive 4.6 Quality	y control of anti		in animals and	plants		1	Ī	
			4.6 Already available						

4.6 ABX used in the agriculture/ veterinary are under the same control as those used in humans Pharmaceuti cal and Herbal Medicines Registration and Control Administrati on/The Dug Inspection Control	ective 4 7 Restric	t the use of criti	cally important	ARX to human	health in the ve	terinary field			
4.7.1 Establish the list of critically important antibiotics for humans in the animal sector.			4.7.1 List put Yes/No	4.7.1 limit the use of critically important ABX in the animal sector.	4.7.1 Yes/No	4.7.1 Once	4.7.1 ABX task force	4.7.1 Checking	4.7.1 Not available
4.7.2 Determine from this list which antibiotics should be restricted for			4.7.2 List put Yes/No	4.7.2 limit the use of critically important ABX in the animal sector.	4.7.2 Yes/No	4.7.2 Once	4.7.2 ABX task force	4.7.2 Checking	4.7.2 Not available

use to humans									
4.7.3 Issue a decree that the critically important ABX to human health are not to be used in animals whether in individual treatment or in herd therapy			4.7.3 Decree issued Yes /No	4.7.3 put a legal frame work for restricting the use of critically important antibiotics	4.7.3 Yes/No	4.7.3 Once	4.7.3 ABX task force	4.7.3 Checking	4.7.3 Not available
4.7.4 Prohibit the importation of these critically important ABX for vet use	4.7.4.1 Decree that prohibits the importation and sale of these ABX in the vet field		4.7.4.1 Decree is issued Yes/No	4.7.4.1 put a legal framework and reinforce the prohibition of importation and sale of critically important ABX in the vet field	4.7.4.1 Yes/No	4.7.4.1 Once	4.7.4.1 PAAF	4.7.4.1 Checking	4.7.4.1 Not available
	ctive 4.8 Organi	ze the use of an			and aquatic ani	mals and agricu	ilture.		
4.8.1 Put (or endorse) guidelines			4.8.1 Guidelines are established	4.8.1 ensure proper antimicrobial	4.8.1 Yes/No	4.8.1 Once	4.8.1 PAAF	4.8.1 Checking	4.8.1 Not available

regarding the use of ABX in animals and agriculture		Yes/No	use in these fields					
4.8.2 Disseminate these guidelines among veterinarians	4.8.2.1 Booklets/ Brochures/a pp that summarize the key points in these guidelines.	4.8.1.1 Proportion of Vet clinics that have easy access to these updated guidelines	4.8.1.1 facilitate access and implementat ion of guidelines recommend ations	4.8.1.1 Number of Vet clinics that have easy access to these updated guidelines/to tal number of vet clinics*100	4.8.1.1 Once/year	4.8.1.1 PAAF	4.8.1.1 data collection	4.8.1.1 not available
	4.8.2.1 Regular workshops for veterinary specialists regarding the application of these guidelines	4.8.2.1 proportion of veterinary specialists attending these workshops	4.8.2.1 proper implementat ion of these guidelines	4.8.2.1 NUMBER of veterinary specialists attending these workshops/t otal number of specialists*1	4.8.2.1 Once/year	4.8.2.1 PAAF	4.8.2.1 data collection	4.8.2.1 Not available
4.8.3 Restrict selling ABX for animal and agriculture health to exclusively	4.8.3.1 Decree that vet pharmacies should dispense ABX only according to vet	4.8.3.1 Decree is issued Yes/No	4.8.3.1 put a legal framework for restricting sale of antibiotics in these fields	4.8.3.1 Yes/No	4.8.3.1 Once	4.8.3.1 PAAF	4.8.3.1 Checking	4.8.3.1 Not available

prescriptions by vets	prescriptions , and should submit these prescriptions with the evidence of selling to PAAF.								
4.8.4 Control the use of ABX in animals	4.8.4.1. Detection of the ABX residues in milk, meat, poultry and eggs		4.8.4.1. Already available	4.8.4.1. Abx use in food is controlled					
	ctive 4.9 Capaci	ty building amo	ng vets in antim	icrobial use and	I AMR			T	
4.9.1 Provide training courses for vets for ABX use according to guidelines and provide certificate.			4.9.1 Proportion of vets that have a yearly certificate	4.9.1 make sure that vets properly prescribe and dispense antimicrobial s	4.9.1 number of vets who attend the certification training/tota I number of vets *100	4.9.1 once/year	4.9.1 PAAF	4.9.1 data collection	4.9.1 Not available
4.9.2 Study baseline consumption imported antibiotics in this field	Pharmaceuti cal Dep in PAAF Committee that will study and document this consumption and compare it to		4.9.2.1 Report issued Yes/No	4.9.2.1 see how much vets are abiding to guidelines and to be able to compare to international data	4.9.2.1 Yes/No	4.9.2.1 Once	4.9.2.1 PAAF	4.9.2.1 Checking	4.9.2.1 Not available

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international				
benchmark				

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