



Final agenda
Meeting of the Immunization Strategic Advisory Group of Experts
6-9 November 2007
Crowne Plaza Hotel, Geneva

Tuesday 6 November 2007

Time	Session	Purpose of session, target outcomes and questions for SAGE	
09:00	Welcome - introduction		20mn
09:20	Report from IVB Director including status of implementation of recommendations Presentation: J-M Okwo-Bele, WHO, 20mn Discussion 20mn	FOR INFORMATION - Update on key WHO activities including on implementation of recommendations from external review of IVB advisory committees - Report to SAGE on achievement of previous recommendations	40mn
10:00	Priorities, major policy and implementation issues: reports from AMRO, EURO, and WPRO (analytical and with emphasis on challenges, specific focus on large countries and implementation/impact of SAGE recommendations) Presentation: C. Ruiz Matus, AMRO, 20mn Discussion 20mn	FOR INFORMATION Provide SAGE with background information in defining global immunization priorities	2hrs
10:40	Coffee/tea break	Break	30mn
11:10	Priorities, major policy and implementation issues: reports from AMRO, EURO, and WPRO (ctd) Presentation: A. Lobanov, EURO, 20mn Discussion 20mn Presentation: Y. Baoping, WPRO, 20mn Discussion 20mn		
12:30	Lunch	Break	1hr 30mn

14:00	Report from the GAVI secretariat Presentation: J. Lob-Levyt, GAVI, 20mn Discussion 20mn	FOR INFORMATION	40mn
14:40	Reports from other Advisory Committees in Immunization GACVS report of June 2007 meeting and agenda of December 2007 meeting and report the October 2007 meeting of the Expert Committee on Biological Standardization (ECBS) GACVS reports provided ahead of meeting ECBS report D. Wood, WHO, 10mn Comments and feed-back 20mn	FOR INFORMATION	30mn
15:10	Pneumococcal conjugate Target Product Profile Introduction: Overview on AMC, TPP production process J. Hombach, WHO, 5mn Pneumococcal Global Serotype Distribution Project K. O'Brien, John Hopkins Bloomberg School of Public Health 15 mn Presentation of TPP document, D. Goldblatt, Chair of ad hoc expert advisory committee 20mn	FOR DECISION -Review the draft target product profile for AMC-eligible pneumococcal vaccines, as proposed by the <i>ad hoc</i> expert advisory committee. -Endorse the document with or without modifications. If considerable modifications will be deemed necessary, the final document should be endorsed by SAGE in a written process after the meeting, to allow for the TPP to be submitted to the independent assessment committee of the AMC secretariat by the end of November 2007.	2hrs 20mn
15:40	Coffee/tea break	Break	30mn
16:10	Pneumococcal conjugate Target Product Profile (ctd) Discussion 1hr 40mn		
18:00	Cocktail		

Wednesday 7 November 2007

08:30	Polio eradication Polio eradication: progress & Risks to the New Milestones, S. Cochi, Chair ACPE, 20mn Discussion 40mn Post-eradication Risk Management: deliberations & findings of the NIH consultation, K. Chumakov, US FDA, 20mn Discussion 40mn	FOR INFORMATION AND DISCUSSION -Update on status/prospects for eradication, including implications of the ongoing, increasingly difficult financing problems -Report from the consultation that NIH hosted in September and discussion on use of IPV for routine immunization. This discussion will feed into a later discussion by ACPE	2hrs
10:30	Coffee/tea break	Break	30mn

Thursday 8 November 2007

08:30	<p>Potential Uses of WHO H5N1 Vaccine Stockpile and H5N1 vaccine</p> <p>Overview, K. Fukuda, WHO, 10 mn Questions, 10mn</p> <p>Indonesia thoughts and concerns, T. Yoga Adimata, Director Control of Direct Transmitted Diseases, Directorate General Disease Control and Environmental Health, Ministry of Health, 10 mn</p> <p>Questions, 10 mn</p> <p>Immunogenicity of human H5N1 vaccines, M. Zambon, Enteric, Respiratory & Neurological Virus Laboratory, Health Protection Agency, 30 mn</p> <p>Safety of human H5N1 Vaccines, S. Salmaso, Istituto Superiore di Sanita, 30 mn</p> <p>Discussion on two previous presentations 30 mn</p> <p>Potential use of H5N1 vaccine to contain an H5N1 influenza pandemic, N. Ferguson, Imperial College, 15 mn</p> <p>Discussion, 15 mn</p> <p>Report of informal consultation on technical specifications for an international H5N1 vaccine stockpile, D. Wood, WHO, 20 mn</p> <p>Discussion, 15 mn</p> <p>General discussion:</p> <p><u>How WHO Should Use Its Stockpile of H5N1 Vaccine?</u></p> <p><u>Additional questions related to H5N1 vaccine being posed to WHO</u></p>	<p>FOR INFORMATION AND DECISION</p> <p><u>FOR DECISION:</u></p> <p>How WHO Should Use Its Stockpile of H5N1 Vaccine</p> <ol style="list-style-type: none"> Should WHO reserve part of its stockpile for pandemic containment operations? If yes, how much? Should WHO reserve part of its stockpile to provide H5N1 vaccine urgently to a defined group of developing countries if an H5N1 pandemic develops? <ol style="list-style-type: none"> If yes, how should recipient countries be defined? If yes, how much vaccine should be allocated to each of the recipient countries? If yes, how much vaccine should be stored for this purpose? Should WHO store the vaccine in a single stockpile or in multiple ones located regionally? <p><u>FOR DISCUSSION: Additional questions related to H5N1 vaccine being posed to WHO</u></p> <ol style="list-style-type: none"> In the current non-pandemic period, should WHO support use of H5N1 vaccine to prevent avian H5N1 infections in countries affected by H5N1? In the current non-pandemic period, should WHO support use of H5N1 vaccine to either immunologically "prime" or "pre-immunize" selected or whole populations against the possibility that an H5N1 pandemic might occur? Should WHO develop global guidance on who should receive pandemic vaccine or should this be 	5hrs 30mn
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		left to countries to decide for themselves?	
10:00	Coffee/tea break	Break	30mn
10:30	Influenza pandemic vaccine (ctd)		
12:30	Lunch break	Break	1hr
13:30	Influenza pandemic vaccine (ctd)		
15:30	Coffee/tea break	Break	30mn
16:00	Updating of position paper on rabies Rabies pathogenesis and evidence base for post exposure prophylaxis C. Hanlon, Kansas State Veterinary Diagnostic Laboratory, 15 mn Rational and evidence base for the safety, efficacy and cost-effectiveness of intradermal immunization, M. Warrell, University of Oxford, 15 mn Practical experience on the use of Intradermal rabies vaccination in Lanka (invited comments from O. Wilamaratne, Medical Research Institute , Colombo, no slide presentation, 5 mn) Proposed updating of the vaccine position paper, B. Bjorvatn, Narestoe, 10 mn Discussion 45 mn	FOR DECISION	1hr 30mn

Friday 9 November 2007

08:30	Use of polysaccharide pneumococcal vaccine Summary of the proposed updated position paper and draft recommendations Presentation: A. Reingold, SAGE member and Chair of SAGE working Group, 30mn Discussion 1hr	FOR DISCUSSION Review the evidence with respect to the following questions/issues and to propose to SAGE related recommendations on vaccine use, with a view to update the 2003 position paper. 1. Review of available data on burden of pneumococcal disease in adults and in high risk groups 2. Effectiveness of 23-valent pneumococcal	1hr 30 mn
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		polysaccharide vaccine (PPV23) in elderly population 3. Effectiveness of PPV23 in high risk groups 4. Safety and efficacy of repeated doses of PPV23 in older children and adults 5. Use of PPV23 to boost kids previously immunized with the 7-valent pneumococcal conjugate vaccine (PCV7) 6. Impact of giving PPV23 to pregnant women/women of reproductive age in order to prevent pneumococcal disease in infants in the first few months of life 7. Consideration of the potential use of PPV23 to mitigate pandemic influenza disease	
10:00	Immunization safety - Cross Departmental report Regional perspectives: progress and challenges (5mn each) AFRO, R. Macauley AMRO, C. Ruiz Matus EMRO, E. Mohsni EURO, A. Lobanov, SEARO, H. Wibisono WPRO, Y Baoping Update on injection safety and integrated infection control strategies in health care settings S. Khamassi , WHO, 15 mn Health care waste management, Y. Chartier, WHO, 15mn Discussion 1hr	FOR INFORMATION AND DISCUSSION	2hrs
10:30	Coffee/tea break	Break	30mn
11:00	Immunization safety - Cross Departmental report (ctd)		
12:30	Closing		20mn
12:50	End of meeting		