Survey – Questions for UN Organizations
As mandated by WHA74 and for input towards EB 150.
August 2021

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<thead>
<tr>
<th>UN Organization</th>
<th>UNICEF</th>
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<tbody>
<tr>
<td>Contact information</td>
<td><a href="mailto:Pbollen@unicef.org">Pbollen@unicef.org</a></td>
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<tr>
<td>Website</td>
<td><a href="http://www.UNICEF.org">www.UNICEF.org</a></td>
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WHO thanks you very much for your participation in the meeting to discuss the standardization of medical devices nomenclature, as mandated by WHA74.

Please find below the survey for the UN Organizations, as mentioned during our meeting. Please provide all the evidence required for every statement provided (for example, website screenshots, links to your website or attach the documents required).

1. Is your organization currently using one or more Medical Device Nomenclature? If yes, please indicate:
   a. which one(s) do you use?
      - EMDN
      - GMDN X
      - UMDNS X
      - UNSPSC X (UNGM)
   Other, specify:
   Developed by your own: Yes: , No: X

2. In which language(s) is your nomenclature available?
   - English X
   - Spanish
   - French
   - Russian
   - Chinese
   - Arabic

3. Please provide a copy of your license agreement including any subscription or other costs associated with the use of your nomenclature.
   - GMDN subscription
   - ECRI subscription excluding access to the UMDNS database

4. Is the subscription free? For each level and in what conditions?
   - GMDN at not cost
   - ECRI (excluding access to UMDNS) at approx. US$ 4,500 / year
e. What are the limitation(s) of the nomenclature(s) used?

<table>
<thead>
<tr>
<th>Limitations / Nomenclature</th>
<th>GMDN</th>
<th>EMDN</th>
<th>UNSPSC</th>
<th>UMDNS</th>
<th>Other</th>
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<tbody>
<tr>
<td>Language</td>
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<td>Definitions</td>
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<td>Other (specify)</td>
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2. Does your organization see the need of using a standardized nomenclature? Please elaborate.

I can’t speak for the organisation; I use these for the identification of products during tender processes.

3. Please describe the reasons your Agency uses the Nomenclature
   a. Planning and evaluation
   b. Budget and financing
   c. Procurement
      X
   d. Maintenance (CMMS)
   e. Medical Technologies quality assurance and surveillance
   f. Medical Technologies Assessment
   g. Complete follow up to supply chain for medical devices
   h. Other (specify):

4. Please describe how codes / nomenclature are used by your final users (beneficiaries)?

I am not aware how final users use nomenclature, or even if they use nomenclature.

5. How do you reconcile when your final users use different nomenclature than your UN Agency, is using?

Have not encountered such a situation.

6. Do you map nomenclatures? If yes, how do you map them?

I do not, I cannot speak of the organisation

7. For UNDP (only): How do you structure the “United Nations Standard Product and Services Codes (UNSPSC)” for medical devices? Can you describe the methodology to assign codes, terms, definitions, and hierarchy? Please provide examples.
8. **For UNDP (only):** Please describe your relationship with GS1 in developing and maintaining the UNSPSC

9. According to your experience with Health Technology Providers, do they use their own nomenclature? If yes, do you map UNSPSC to the nomenclature used by the Health Technology Provider? How do you map them?
   No mapping is conducted

10. According to your experience with final users (beneficiaries), if the receipt country has not developed their own nomenclature or adopted an international relevant nomenclature (EMDN, GMDN, UMDNS, for example), what is the methodology to implement UNSPSC with final users (beneficiaries)?
   No experience with beneficiaries.

11. Have you had any experience working with Unique Device Identifiers? If so, please describe any nomenclature-related work in this area.
   No

Please kindly fill the survey and send to [medicaldevices@who.int](mailto:medicaldevices@who.int) before 26th August,

Many thanks