

WHO Update Column



Welcome to the initiation of a new column in our Global Clinical Engineering Journal that will serve the readership as additional information source of international health technology interest, as opportunity to exchange comment and collect your feedback, and promote stronger engagement with WHO representative members.

This new added feature will be directed under the expertise of Adriana Velazquez Berumen, Senior advisor on medical devices, Department of Essential Medicines and Health Products, at the World Health Organization in Geneva, Switzerland ([url: www.who.int/medical_devices/en/](http://www.who.int/medical_devices/en/) and *e-mail: velazquezberumena@who.int*). Your feedback and promotion and sharing of the information posted in this column are welcome.

At a press conference at the United Nations in Geneva, on July 9, 2019, WHO launched the 2nd WHO Model List of Essential In Vitro Diagnostics (EDL) and the application for the 3rd EDL List has opened. (https://www.who.int/medical_devices/publications/Second_WHO_Model_List_of_Essential_In_Vitro_Diagnostics/en/). The objective of the EDL is to increase access, affordability, availability of these tests globally, to support universal health coverage and better health for all. It includes laboratory tests as well as point of care and the objective is that countries will refer to WHO lists and updated their national reference lists for public procurement or reimbursement.

The process to select these diagnostics included: assess the tests submitted for the 2nd EDL, series of consultations, including public comments and final review, analysis and discussion by Members of Strategic Advisory Group of Experts on In Vitro Diagnostics (SAGE IVD) and WHO staff.

The full report of the process will present a description of the methodologies, reviews, evidence, references and recommendations of the SAGE IVD members and will be published in September 2019 as part of WHO Technical Report Series. and will be found at: https://www.who.int/medical_devices/diagnostics/selection_in-vitro/en/.

You may find of interest the exciting update about Health Product Profile Directory <https://www.who.int/tdr/diseases-topics/product-directory/en/> a free-to-use online resource created and developed by TDR (Special Programme for Research and Training in Tropical Diseases) on behalf of WHO as a global public good to improve the efficiency of efforts to develop new products for neglected diseases and populations as well as threats to global health. It provides a searchable database of the 8-10 key characteristics used to describe desired health products, including medicines, vaccines, diagnostics and medical equipment. Links are provided to access the full Product Profile document where this is publicly available. The Directory was launched in May 2019 and will be continuously updated. You are encouraged to visit the website provided.



Finally, I would also like to share with you additional highlights from the July 2019 WHO Medical Devices Newsletter that describe career opportunities at WHO. In particular, a full time Technical Officer position is open (Position 1902257) where you can find additional information about it such as job description and how to apply as well as about more opportunities at <https://careers.who.int/career-section/ex/jobdetail.ftl?job=1902257&tz=GMT%2B02%3A00&tzname=Europe%2FBerlin>.

I am delighted with this opportunity to connected with you and to create a new platform for increasing knowledge about and communication between stake holders with interest in health technologies.

Respectfully,

Adriana

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