Update on COVID-19 in vitro diagnostics listed by National Regulatory Authorities in IMDRF jurisdictions

Several National Regulatory Authorities (NRAs) have established assessment procedures for listing/authorizing COVID-19 in vitro diagnostics (IVD). The links provided below present information on IVDs authorized for use in the International Medical Device Regulators Forum (IMDRF)\(^1\) jurisdictions along with other useful information on policies and guidance.

Disclaimer: WHO does not endorse any of the lists provided by NRAs. This information is provided exclusively to assist stakeholders with identifying the links to the various lists.

1. **United States of America:**
   
   
   

   Coronavirus (COVID-19) Diagnostic Tests Hotline:
   
   For test developers and labs who have questions about the EUA process or spot shortages of testing supplies, contact our toll-free phone line 24 hours a day: 1-888-INFO-FDA (1-888-463-6332), then press star (*)

2. **Canada:**
   

3. **Japan**

   Information on products authorized by the Ministry of Health, Labour and Welfare (MHLW) of Japan can be found in the attached document.

4. **Australia**
   
   

5. **South Korea**

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\(^1\) For more information on the IMDRF please refer to this link: [http://www.imdrf.org/](http://www.imdrf.org/)

\(^2\) For agencies who provided such information to WHO/RPQ
6. Singapore:

7. China:
https://mp.weixin.qq.com/s/8nXIXJbE95hp0gcqsqj35g

8. Brazil
https://consultas.anvisa.gov.br/#/saude/q/?nomeTecnico=coronav%C3%ADrus

9. Russia:
https://www.roszdravnadzor.ru/services/misearch

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