SENSITIZATION WEBINAR FOR MANUFACTURERS OF WHO PREQUALIFIED IN VITRO DIAGNOSTICS (IVDs) ON THE WHO COLLABORATIVE REGISTRATION PROCEDURE (CRP)

25 June 2026

13:00 to 14:30PM CEST

Virtually by Zoom

Register here

For more information please email: crp@who.int

World Health Organization

Special Access Program (SAP)
Regulation and Safety Unit (REG)
Regulation and Prequalification Department (RPQ)

WEBINAR OVERVIEW

The WHO Collaborative Registration Procedure (CRP) for in vitro diagnostics (IVDs) provides an efficient reliance-based pathway to support accelerated national registration of WHO-prequalified IVDs. This webinar will provide manufacturers with updates on the expanding participation of countries in the CRP for IVDs, recent revisions to WHO guidance supporting CRP implementation, and practical insights on effectively navigating the process.

Target audience

- WHO-prequalified IVD manufacturers
- Manufacturers with IVD products under WHO PQ assessment

Agenda

13:00 – 13:10 CEST	Opening Remarks *Marie Valentin, Sr. Technical Officer, WHO HQ/RPQ/REG/ SAP*
13:10 – 14:05 CEST	WHO CRP for IVDs: Overview, implementation updates, process, and support mechanisms *Agnes Sitta Kijo, Technical Officer, WHO HQ/RPQ/REG/RSS*
14:05 – 14:25 CEST	Interactive Q&A Session *All*
14:25 – 14:30 CEST	Wrap-up and Closing Remarks *Marie Valentin, Sr. Technical Officer, WHO HQ/RPQ/REG/ SAP*

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