HPRA
An tÚdarás Rialála Táirgí Sláinte
Health Products Regulatory Authority

Agenda for
The Application of the *In Vitro* Diagnostic Medical Devices Regulation (IVDR)

9:30 am – 1 pm, Monday 9 May 2022

SESSION	TOPIC	PRESENTER
1	Welcome and Introduction	Niall MacAleenan, Director of Medical Devices, HPRA
	European and National Implementation of the IVDR	Sinead Duggan, Communications and Policy Lead, HPRA
First Q&A Session		
2	Post Market Surveillance, Vigilance and Market Surveillance Requirements	Elaine Kilbane, IVD Operations Manager, HPRA
	HPRA IVD Inspection Activities	Jennifer Roche, Medical Device Inspection Operations Manager, HPRA
Second Q&A Session & Short Break		
3	Performance Studies	Philip Kelly, IVDR Project Manager, HPRA
	Registration Requirements and Certificates of Free Sale	Andrea Hanson, Market Analysis Lead, HPRA
Third Q&A Session and Close of Event		