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CoSara Diagnostics bags CDSCO clearance for high-risk HPV multiplex test

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The multiplex test is built on the company's patented CoPrimer technology



Co-Diagnostics, a molecular diagnostics company, has announced that CoSara Diagnostics, its joint venture for manufacturing and sales in India, has received clearance by the Central Drugs Standard Control Organization (CDSCO) to manufacture and sell its SARAGENE Human Papillomavirus High-Risk (HPV-HR) Real-Time PCR test as anin vitro diagnostic (IVD).

CoSara's new multiplex test, the 12th CoSara assay to receive CDSCO approval, is built on the company's patented CoPrimer technology and designed to detect and differentiate between HPV genotypes 16 and 18, while simultaneously detecting high-risk carcinogenic HPV types 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, and 68.

CoSara has previously received CDSCO clearance for RT-PCR tests for Mycobacterium tuberculosis, malaria, hepatitis B, hepatitis C, HPV (types 16 and 18 only), two COVID-19 assays, chikungunya, dengue, a dengue/chikungunya duplex test, and a Flu A/Flu B/COVID-19 (ABC) multiplex test, all designed using the Company's patented CoPrimer technology and cleared to be manufactured and sold as IVDs in the Indian market.