HER2 Digital Manual Read, 1/14

January 2014—Royal Philips received 510(k) clearance from the FDA to market its HER2/neu IHC Digital Manual Read product in the U.S.

Using Royal Philips' system, pathologists can now benefit from digital imaging advancements for scoring Hercep-Test-stained tumor tissue slides digitized by Philips' ultra-fast scanner and made accessible through Philips' advanced image viewing and analysis management system.

The Philips HER2/neu IHC Digital Manual Read is based on the Philips Digital Pathology Solution platform and was commercially introduced in Europe and Asia Pacific in 2012.

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