

BD Kiestra ReadA device listed with the FDA

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May 2020—BD announced that its BD Kiestra ReadA is device listed with the FDA. The BD Kiestra ReadA incubation and imaging system, which is available as a standalone instrument, aims to help improve operational efficiency in clinical microbiology laboratories by automating routine plate management tasks and delivering accuracy through standardized digital image acquisition.

[BD](#), 201-847-6800



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