



Acknowledgment Letter

8/21/2020

Pinnacle IVD Corporation
315 Deaderick Street, St #1550
Nashville, TN 37238 UNITED
STATES

Dear Charles Balentine:

The Center for Devices and Radiological Health (CDRH) of the Food and Drug Administration (FDA) has received your submission. This submission has been assigned the unique document control number below. All future correspondence regarding this submission should be identified prominently with the number assigned and should be submitted to the Document Control Center at the above letterhead address. Failure to do so may result in processing delays. If you believe the information identified below is incorrect, please contact the Office of Product Evaluation and Quality (OPEQ) submission support at (301) 796-5640 or OPEQSubmissionSupport@fda.hhs.gov.

Submission Number: PEUA201683
Received: 8/20/2020
Applicant: Pinnacle IVD Corporation
Device: Pinnacle Cov-ID RAD Rapid Antigen Detection Test

We will notify you when the review of this document has been completed or if any additional information is required. For information about CDRH review regulations and policies, please refer to <http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/default.htm>.

Sincerely yours,

Center for Devices and Radiological Health