FDA 510(k) Premarket Notification

K070747 QUICK VUE RSV TEST, MODELS 20193, 20199

CLEARED

Apr 23, 2007
35 days to decision

K070747 · Product code: **GQG** · Microbiology
Source: https://www.510kdatabase.net/k070747/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antigen, Cf (including Cf Controls), Respiratory Syncytial Virus (GQG)
Date received	Mar 19, 2007
Decision date	Apr 23, 2007
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Quidel Corp.**
Location	Washington, DC, US
Contact	JENNIFER HANKARD
510(k) history	93 submissions · 93 cleared · 1983-2013