

**K070747 QUICK VUE RSV TEST, MODELS 20193, 20199**Apr 23, 2007  
35 days to decisionK070747 · 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	<b>Quidel Corp.</b>
Location	Washington, DC, US
Contact	JENNIFER HANKARD
510(k) history	93 submissions · 93 cleared · 1983-2013

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070747/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 8, 2026