

**K090971 BD PROBETEC NEISSERIA GONORRHOEAE (GC) QX
AMPLIFIED DNA ASSAY**Jun 5, 2009
60 days to decisionK090971 · Product code: **LSL** · Microbiology
Source: <https://www.510kdatabase.net/k090971/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dna-reagents, Neisseria (LSL)
Date received	Apr 6, 2009
Decision date	Jun 5, 2009
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Becton, Dickinson & CO
Location	Franklin Lakes, NJ, US
Contact	KATHRYN BABKA CARR
510(k) history	190 submissions · 190 cleared · 2001-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k090971/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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