

**K033861 BD PROBETEC ET LEGIONELLA PNEUMOPHILA (LP)
AMPLIFIED DNA ASSAY**Mar 9, 2004
88 days to decisionK033861 · Product code: **LQH** · Microbiology
Source: <https://www.510kdatabase.net/k033861/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dna-reagents, Legionella (LQH)
Date received	Dec 12, 2003
Decision date	Mar 9, 2004
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033861/> Data sourced from FDA 510(k) public records (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. - Generated May 24, 2026