

K831760 LEGIONELLA IFA KIT IAug 8, 1983
68 days to decisionK831760 · Product code: **LHL** · Toxicology
Source: <https://www.510kdatabase.net/k831760/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reagents, Antibody, Legionella, Direct & Indirect Fluorescent (LHL)
Date received	Jun 1, 1983
Decision date	Aug 8, 1983
Days to decision	68 days
Third-party review	No

APPLICANT

Company	Bionetic Laboratory Products
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1983-1984

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Device record: <https://www.510kdatabase.net/k831760/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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