

K834067 LEGIONELLA DIFFERENTIATION DISKJan 3, 1984
39 days to decisionK834067 · Product code: **GRO** · Microbiology
Source: <https://www.510kdatabase.net/k834067/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Fluorescent, All Types, Hemophilus Spp. (GRO)
Date received	Nov 25, 1983
Decision date	Jan 3, 1984
Days to decision	39 days
Third-party review	No

APPLICANT

Company	Remel Co.
Location	Mchenry, IL, US
510(k) history	137 submissions · 137 cleared · 1979-2001

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Device record: <https://www.510kdatabase.net/k834067/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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