

K840799 V.D.T.-GONORRHEA TEST KITJun 7, 1984
105 days to decisionK840799 · Product code: **LIO** · Microbiology
Source: <https://www.510kdatabase.net/k840799/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Specimen Collection (LIO)
Date received	Feb 23, 1984
Decision date	Jun 7, 1984
Days to decision	105 days
Third-party review	No

APPLICANT

Company	M.G. Laboratories, Inc.
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k840799/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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