FDA 510(k) Premarket Notification

K840799 V.D.T.-GONORRHEA TEST KIT

CLEARED

Jun 7, 1984
105 days to decision

K840799 · 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