FDA 510(k) Premarket Notification

K802423 PHADEBACT HAEMOPHILUS TEST

CLEARED

Nov 12, 1980
40 days to decision

K802423 · Product code: **GRO** · Microbiology
Source: https://www.510kdatabase.net/k802423/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Fluorescent, All Types, Hemophilus Spp. (GRO)
Date received	Oct 3, 1980
Decision date	Nov 12, 1980
Days to decision	40 days
Third-party review	No

APPLICANT

Company	**Pharmacia, Inc.**
Location	Mchenry, IL, US
510(k) history	129 submissions · 126 cleared · 1976-1998