FDA 510(k) Premarket Notification

K833888 PENASE

CLEARED

Jan 3, 1984
55 days to decision

K833888 · Product code: **MDB** · Microbiology
Source: https://www.510kdatabase.net/k833888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Blood Culturing (MDB)
Date received	Nov 9, 1983
Decision date	Jan 3, 1984
Days to decision	55 days
Third-party review	No

APPLICANT

Company	**Oxoid U.S.A., Inc.**
Location	Mchenry, IL, US
510(k) history	93 submissions · 93 cleared · 1980-1989