

K833888 PENASEJan 3, 1984
55 days to decisionK833888 · 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

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Device record: <https://www.510kdatabase.net/k833888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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