FDA 510(k) Premarket Notification

K821659 RESPIRACULT-STREP DEEP BED CULTURE PAD

CLEARED

Aug 5, 1982
62 days to decision

K821659 · Product code: **JSH** · Microbiology
Source: https://www.510kdatabase.net/k821659/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Culture Media, Non-selective And Differential (JSH)
Date received	Jun 4, 1982
Decision date	Aug 5, 1982
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Orion Diagnostica, Inc.**
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1980-1994