FDA 510(k) Premarket Notification

K811334 MYCOSEL

CLEARED

May 29, 1981
17 days to decision

K811334 · Product code: **JSI** · Microbiology
Source: https://www.510kdatabase.net/k811334/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Culture Media, Selective And Differential (JSI)
Date received	May 12, 1981
Decision date	May 29, 1981
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Acumedia Manufacturers, Inc.**
Location	Mchenry, IL, US
510(k) history	177 submissions · 177 cleared · 1981-1988