

**K811334 MYCOSEL**May 29, 1981  
17 days to decisionK811334 · Product code: **JSI** · Microbiology  
Source: <https://www.510kdatabase.net/k811334/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k811334/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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