

**K831359 MODIFIED NEW YORK MEDIUM**May 18, 1983  
22 days to decisionK831359 · Product code: **JTY** · Microbiology  
Source: <https://www.510kdatabase.net/k831359/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Culture Media, For Isolation Of Pathogenic Neisseria (JTY)
Date received	Apr 26, 1983
Decision date	May 18, 1983
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Gibco Laboratories Life Technologies, Inc.</b>
Location	Mchenry, IL, US
510(k) history	67 submissions · 67 cleared · 1976-1987

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

Device record: <https://www.510kdatabase.net/k831359/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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