

**K791049 SYNALON ROLLER BOTTLE**Jun 28, 1979  
22 days to decisionK791049 · Product code: **KJC** · Pathology  
Source: <https://www.510kdatabase.net/k791049/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bottle, Roller, Tissue Culture (KJC)
Date received	Jun 6, 1979
Decision date	Jun 28, 1979
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Dynatech Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	44 submissions · 44 cleared · 1978-1996

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

Device record: <https://www.510kdatabase.net/k791049/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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