

**K840503 SDL LISTERIA MONOCYTOGENES CONTROL**Mar 23, 1984  
46 days to decisionK840503 · Product code: **GSI** · Microbiology  
Source: <https://www.510kdatabase.net/k840503/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antigens, Slide And Tube, All Types, Listeria Monocytogenes (GSI)
Date received	Feb 6, 1984
Decision date	Mar 23, 1984
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Scientific Device Laboratory, Inc.</b>
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1984-1994

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

Device record: <https://www.510kdatabase.net/k840503/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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