FDA 510(k) Premarket Notification

# K840503 SDL LISTERIA MONOCYTOGENES CONTROL

**CLEARED**

Mar 23, 1984
46 days to decision

K840503 · 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 | **Scientific Device Laboratory, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 28 submissions · 28 cleared · 1984-1994 |