

K901237 GLOVE LEAK DETECTORAug 21, 1990
159 days to decisionK901237 · Product code: **LDQ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k901237/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Mar 15, 1990
Decision date	Aug 21, 1990
Days to decision	159 days
Third-party review	No

APPLICANT

Company	Utah Medical Products, Inc.
Location	Mchenry, IL, US
Contact	O GOODMAN
510(k) history	38 submissions · 38 cleared · 1979-2015

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Device record: <https://www.510kdatabase.net/k901237/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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