FDA 510(k) Premarket Notification

K901237 GLOVE LEAK DETECTOR

CLEARED

Aug 21, 1990
159 days to decision

K901237 · Product code: **LDQ** · General & Plastic Surgery
Source: 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