

**K913696 GLOVE-CHEK PORTABLE FLUID LEAKAGE SENSING
DEVICE**Feb 18, 1992
183 days to decisionK913696 · Product code: **LDQ** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k913696/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Aug 19, 1991
Decision date	Feb 18, 1992
Days to decision	183 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Vertex Technical Group, Inc.
Location	Bethesda, MD, US
Contact	JOYCE T DAVIS
510(k) history	1 submissions · 1 cleared · 1992-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913696/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 7, 2026