FDA 510(k) Premarket Notification

K970170 SWIFT DELIVERY PRODUCT #002-VAC EX CUP W/VAC REL VALVE

CLEARED

Mar 28, 1997
78 days to decision

K970170 · Product code: **HDB** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k970170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Extractor, Vacuum, Fetal (HDB)
Date received	Jan 9, 1997
Decision date	Mar 28, 1997
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Richard G. Lindsay**
Location	Oklahoma City, OK, US
Contact	RICHARD G LINDSAY
510(k) history	1 submissions · 1 cleared · 1997-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k970170/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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