FDA 510(k) Premarket Notification

K960549 MITYVAC DIRECTIONAL VACUUM EXTRACTOR, DISPOSABLE

CLEARED

Aug 29, 1996
203 days to decision

K960549 · Product code: **HDB** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k960549/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Extractor, Vacuum, Fetal (HDB)
Date received	Feb 8, 1996
Decision date	Aug 29, 1996
Days to decision	203 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Neward Enterprises, Inc.**
Location	Mchenry, IL, US
Contact	ALLYSON CARMACK
510(k) history	8 submissions · 8 cleared · 1984-1996