FDA 510(k) Premarket Notification

K800542 100 ML SNYDER HEMOVAC

CLEARED

Apr 2, 1980
23 days to decision

K800542 · Product code: **GCY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k800542/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Suction, Single Patient Use, Portable, Nonpowered (GCY)
Date received	Mar 10, 1980
Decision date	Apr 2, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Snyder Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1978-1984