

K800542 100 ML SNYDER HEMOVACApr 2, 1980
23 days to decisionK800542 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800542/> 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 May 28, 2026