FDA 510(k) Premarket Notification

K810409 MODEL C2100-SERIES OPHTHALMIC CRYOPROBE

CLEARED

Mar 20, 1981
30 days to decision

K810409 · Product code: **GEH** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k810409/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryosurgical, Accessories (GEH)
Date received	Feb 18, 1981
Decision date	Mar 20, 1981
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Valleylab, Inc.**
Location	Mchenry, IL, US
510(k) history	94 submissions · 93 cleared · 1976-2003

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k810409/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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