FDA 510(k) Premarket Notification

K782065 LASER, PHOTOCOAGULATION

CLEARED

Dec 28, 1978
15 days to decision

K782065 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k782065/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Dec 13, 1978
Decision date	Dec 28, 1978
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Cavitron Corp.**
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1976-1981

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k782065/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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