FDA 510(k) Premarket Notification

K801336 MODEL 900K PHOTOCOAGULATOR

CLEARED

Jul 28, 1980
54 days to decision

K801336 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k801336/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Jun 4, 1980
Decision date	Jul 28, 1980
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Coherent Medical Division**
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1990