FDA 510(k) Premarket Notification

K801850 CHARLES PNEUMATIC INTRAOCULAR SCISSORS

CLEARED

Sep 16, 1980
43 days to decision

K801850 · Product code: **HQE** · Ophthalmic
Source: https://www.510kdatabase.net/k801850/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Aug 4, 1980
Decision date	Sep 16, 1980
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Cooper Medical Devices Corp.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1979-1981