FDA 510(k) Premarket Notification

K782106 COAGULATOR, MODEL 40 AARGON

CLEARED

Feb 8, 1979
57 days to decision

K782106 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k782106/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Dec 13, 1978
Decision date	Feb 8, 1979
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Lasertek OY**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1979-1984