FDA 510(k) Premarket Notification

K832089 MODEL 150K LASER COAGULATOR

CLEARED

Jan 30, 1984
221 days to decision

K832089 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k832089/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Jun 23, 1983
Decision date	Jan 30, 1984
Days to decision	221 days
Third-party review	No

APPLICANT

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