

K832089 MODEL 150K LASER COAGULATORJan 30, 1984
221 days to decisionK832089 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k832089/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026