

K782106 COAGULATOR, MODEL 40 AARGONFeb 8, 1979
57 days to decisionK782106 · 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

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Device record: <https://www.510kdatabase.net/k782106/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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