

**K801336 MODEL 900K PHOTOCOAGULATOR**Jul 28, 1980  
54 days to decisionK801336 · Product code: **HQF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k801336/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Jun 4, 1980
Decision date	Jul 28, 1980
Days to decision	54 days
Third-party review	No

**APPLICANT**

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Company	<b>Coherent Medical Division</b>
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1990

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Device record: <https://www.510kdatabase.net/k801336/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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