

K782054 LASER, CO2 SURGICALJan 17, 1979
41 days to decisionK782054 · Product code: **HHR** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k782054/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Surgical, Gynecologic (HHR)
Date received	Dec 7, 1978
Decision date	Jan 17, 1979
Days to decision	41 days
Third-party review	No

APPLICANT

Company	Cavitron Corp.
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1976-1981

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Device record: <https://www.510kdatabase.net/k782054/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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