FDA 510(k) Premarket Notification

K782054 LASER, CO2 SURGICAL

CLEARED

Jan 17, 1979
41 days to decision

K782054 · Product code: **HHR** · Obstetrics & Gynecology
Source: 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