FDA 510(k) Premarket Notification

K905240 VESSELOOPS

CLEARED

Feb 6, 1991
78 days to decision

K905240 · Product code: **KDC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k905240/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Nov 20, 1990
Decision date	Feb 6, 1991
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Sparta Surgical Corp.**
Location	Hayward, CA, US
Contact	CARLETON F KIMBER
510(k) history	7 submissions · 6 cleared · 1989-1997