FDA 510(k) Premarket Notification

K922878 MODEL GT1ST CRYOSURGICAL INSTRUMENT

CLEARED

Sep 2, 1992
79 days to decision

K922878 · Product code: **GEH** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k922878/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryosurgical, Accessories (GEH)
Date received	Jun 15, 1992
Decision date	Sep 2, 1992
Days to decision	79 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Gyne-Tech Instrument Corp.**
Location	Burbank, CA, US
Contact	KERMIT FLOYD
510(k) history	3 submissions · 3 cleared · 1992-1996