FDA 510(k) Premarket Notification

K920040 INTRADUCER

CLEARED

Feb 28, 1992
53 days to decision

K920040 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k920040/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jan 6, 1992
Decision date	Feb 28, 1992
Days to decision	53 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Taut, Inc.**
Location	Walker, MI, US
Contact	KENSETH
510(k) history	16 submissions · 16 cleared · 1983-2002