FDA 510(k) Premarket Notification

K920217 MAX-E-VAC

CLEARED

Jul 23, 1992
189 days to decision

K920217 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k920217/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 16, 1992
Decision date	Jul 23, 1992
Days to decision	189 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Leisegang Medical, Inc.**
Location	Miami, FL, US
Contact	DOUGLAS KWART
510(k) history	10 submissions · 10 cleared · 1986-1999