FDA 510(k) Premarket Notification

K960695 MODEL 200C ELECTROSURGICAL PROBE

CLEARED

Apr 30, 1996
70 days to decision

K960695 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k960695/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 20, 1996
Decision date	Apr 30, 1996
Days to decision	70 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Zomed Intl.**
Location	Mountain View, CA, US
Contact	THOMAS C WEHMAN, PH.D.
510(k) history	7 submissions · 7 cleared · 1995-1996