FDA 510(k) Premarket Notification

K960808 CONMED SELECT ONE LAPAROSCOPY ELECTRODE

CLEARED

Mar 21, 1996
22 days to decision

K960808 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k960808/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 28, 1996
Decision date	Mar 21, 1996
Days to decision	22 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Conmedcorp**
Location	Dayton, OH, US
Contact	IRA D DUESLER, JR.
510(k) history	92 submissions · 92 cleared · 1981-2010