FDA 510(k) Premarket Notification

K821356 DESERET CAPACITY PLUS ELECTROSURG. PAD

CLEARED

Jul 14, 1982
69 days to decision

K821356 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k821356/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 6, 1982
Decision date	Jul 14, 1982
Days to decision	69 days
Third-party review	No

APPLICANT

Company	**Warner-Lambert Co.**
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1979-2003