FDA 510(k) Premarket Notification

K822092 CAMERON-MILLER #40-1516

CLEARED

Sep 21, 1982
67 days to decision

K822092 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k822092/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 16, 1982
Decision date	Sep 21, 1982
Days to decision	67 days
Third-party review	No

APPLICANT

Company	**Cameron-Miller, Inc.**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1977-1994