FDA 510(k) Premarket Notification

K914883 CIRCON ACMI GEN SURG INSTRU FOR MINI INVASIVE SURG

CLEARED

Jan 15, 1993
443 days to decision

K914883 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k914883/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 30, 1991
Decision date	Jan 15, 1993
Days to decision	443 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Circon Video**
Location	Stanford, CT, US
Contact	ERVIN F TAYLOR
510(k) history	14 submissions · 14 cleared · 1990-2002