FDA 510(k) Premarket Notification

CLEARED

K944817 NEUTRALECT ADULT STANDARD, RETURN (REM), PAEDIATRIC STANDARD, RETURN (REM)

Nov 9, 1994
41 days to decision

K944817 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k944817/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 29, 1994
Decision date	Nov 9, 1994
Days to decision	41 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Msb , Ltd.**
Location	Marlborough, Wiltshire, GB
Contact	MARTIN BEAUMONT
510(k) history	7 submissions · 7 cleared · 1994-1998