FDA 510(k) Premarket Notification

K053160 MICRUS MODIFIED MICROCOIL 18-SYSTEM, CERECYTE, MODEL CSP & CHE

CLEARED

Dec 7, 2005
23 days to decision

K053160 · Product code: **HCG** · Neurology
Source: https://www.510kdatabase.net/k053160/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Nov 14, 2005
Decision date	Dec 7, 2005
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Micrus Endovascular Corporation**
Location	Sunnyvale, CA, US
Contact	MARGARET WEBBER
510(k) history	23 submissions · 23 cleared · 2005-2011