FDA 510(k) Premarket Notification

K062036 MICRUS MICROCOIL SYSTEM PRESIDIO-18, MODEL PC4

CLEARED

Aug 25, 2006
37 days to decision

K062036 · Product code: **HCG** · Neurology
Source: https://www.510kdatabase.net/k062036/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Jul 19, 2006
Decision date	Aug 25, 2006
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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