FDA 510(k) Premarket Notification

K051860 FLEXIBLE INTRODUCER CANNULA, MODEL 135-1837

CLEARED

Aug 22, 2005
45 days to decision

K051860 · Product code: **MIA** · Anesthesiology
Source: https://www.510kdatabase.net/k051860/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Spinal, Short Term (MIA)
Date received	Jul 8, 2005
Decision date	Aug 22, 2005
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Epimed International, Inc.**
Location	Johnstown, NY, US
Contact	CHRISTOPHER LAKE
510(k) history	14 submissions · 14 cleared · 1998-2020