FDA 510(k) Premarket Notification

K023173 ANDREWS INTRODUCER, MODEL ASI01

CLEARED

Dec 18, 2002
86 days to decision

K023173 · Product code: **MDM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k023173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, Surgical, General Use (MDM)
Date received	Sep 23, 2002
Decision date	Dec 18, 2002
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Andrews Surgical Innovations, Ltd.**
Location	Co. Dublin, IE
Contact	EMMET ANDREWS
510(k) history	1 submissions · 1 cleared · 2002-2002