FDA 510(k) Premarket Notification

K071543 HALO360' COAGULATION CATHETER

CLEARED

Jun 29, 2007
24 days to decision

K071543 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k071543/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 5, 2007
Decision date	Jun 29, 2007
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Barrx Medical, Incorporated**
Location	Washington, Dc, DC, US
Contact	VIORICA FILIMON
510(k) history	13 submissions · 13 cleared · 2005-2010