FDA 510(k) Premarket Notification

K071887 SPORTMESH OR ARTELON TISSUE REINFORCEMENT

CLEARED

Sep 18, 2007
71 days to decision

K071887 · Product code: **FTL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k071887/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Jul 9, 2007
Decision date	Sep 18, 2007
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Artimplant AB**
Location	Washington, DC, US
Contact	TERRY S POWELL
510(k) history	7 submissions · 7 cleared · 2003-2007