FDA 510(k) Premarket Notification

K061892 PROPATCH SOFT TISSUE REPAIR MATRIX

CLEARED

Nov 22, 2006
142 days to decision

K061892 · Product code: **FTM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k061892/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Mesh, Surgical (FTM)
Date received	Jul 3, 2006
Decision date	Nov 22, 2006
Days to decision	142 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cryolife, Inc.**
Location	Kennesaw, GA, US
Contact	JOHN D FERROS
510(k) history	12 submissions · 12 cleared · 2006-2019