FDA 510(k) Premarket Notification

K061030 BIOBLANKET SURGICAL MESH

CLEARED

May 9, 2006
25 days to decision

K061030 · Product code: **FTM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k061030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical (FTM)
Date received	Apr 14, 2006
Decision date	May 9, 2006
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Kensey Nash Corporation**
Location	Exton, PA, US
Contact	DEBORAH RACIOPPI
510(k) history	6 submissions · 6 cleared · 2006-2013