FDA 510(k) Premarket Notification

K021835 BRENNEN MEDICAL SURGICAL MESH, GLUCAMESH/GLUCATEX

CLEARED

Oct 17, 2002
135 days to decision

K021835 · Product code: **FTL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k021835/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Jun 4, 2002
Decision date	Oct 17, 2002
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Brennen Medical, Inc.**
Location	St, Paul, MN, US
Contact	PHILLIP B LAWIN
510(k) history	17 submissions · 16 cleared · 1994-2006