FDA 510(k) Premarket Notification

K081882 MODIFICATION TO XYLOS SURGICAL MESH

CLEARED

Jul 11, 2008
9 days to decision

K081882 · Product code: **FTM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k081882/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical (FTM)
Date received	Jul 2, 2008
Decision date	Jul 11, 2008
Days to decision	9 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Xylos Corporation**
Location	Washington, DC, US
Contact	GERRY ANN OSTER
510(k) history	9 submissions · 8 cleared · 1998-2011