

**K021835 BRENNEN MEDICAL SURGICAL MESH,  
GLUCAMESH/GLUCATEX**

Oct 17, 2002  
135 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k021835/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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