

K980483 MENTOR SUSPEND SLINGApr 30, 1998
80 days to decisionK980483 · Product code: **FTL** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k980483/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Feb 9, 1998
Decision date	Apr 30, 1998
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mentor Corp.
Location	Mchenry, IL, US
Contact	DONNA A CRAWFORD
510(k) history	61 submissions · 61 cleared · 1977-2013

Mentor Corp. is a surgical aesthetics and medical device company based in McHenry, US. Now part of Johnson & Johnson MedTech, the brand supplies products to plastic surgeons and specialists worldwide. Mentor has received FDA 510(k) clearances from total submissions since its first clearance in 1977. The company's regulatory record spans General & Plastic Surgery, Gastroenterology & Urology, Obstetrics & Gynecology, and Radiology device categories. The latest clearance was recorded in 2013, reflecting the company's historical significance in surgical device innovation. Men...

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