

K863565 MENTOR ALLOPLASTIC SPERMATOCELENov 19, 1986
68 days to decisionK863565 · Product code: **LQS** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k863565/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spermatocele, Alloplastic (LQS)
Date received	Sep 12, 1986
Decision date	Nov 19, 1986
Days to decision	68 days
Third-party review	No

APPLICANT

Company	Mentor Corp.
Location	Mchenry, IL, US
Contact	BYRON WICKETT
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...
