

**K981527 MENTOR PATHWAY INTRODUCER/DILATOR**Jul 9, 1998  
72 days to decisionK981527 · Product code: **QPB** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k981527/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suction, Lipoplasty For Removal (QPB)
Date received	Apr 28, 1998
Decision date	Jul 9, 1998
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mentor Corp.</b>
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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