FDA 510(k) Premarket Notification

K870059 MENTOR NIPPLE RECONSTRUCTION DEVICE

CLEARED

Feb 24, 1987
48 days to decision

K870059 · General & Plastic Surgery
Source: https://www.510kdatabase.net/k870059/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jan 7, 1987
Decision date	Feb 24, 1987
Days to decision	48 days
Third-party review	No

APPLICANT

Company	**Mentor Corp.**
Location	Mchenry, IL, US
Contact	LYNN BRECKENRIDGE
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...