FDA 510(k) Premarket Notification

K900371 MENTOR MODIFIED MALLEABLE PENILE PROSTHESES

CLEARED

Nov 2, 1990
280 days to decision

K900371 · Product code: **FAE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k900371/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Penile (FAE)
Date received	Jan 26, 1990
Decision date	Nov 2, 1990
Days to decision	280 days
Third-party review	No

APPLICANT

Company	**Mentor Corp.**
Location	Mchenry, IL, US
Contact	TERRY A THOMPSON
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...