

K811223 VASOVASOSTOMY STABILIZER PLATFORM DEVMay 21, 1981
17 days to decisionK811223 · Product code: **KOA** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k811223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Surgical Instruments, G-u, Manual (and Accessories) (KOA)
Date received	May 4, 1981
Decision date	May 21, 1981
Days to decision	17 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	Biomet, Inc.
Location	Mchenry, IL, US
Website	http://www.biomet.com/
510(k) history	441 submissions · 419 cleared · 1978-2026

Biomet, Inc. is an orthopedic medical device manufacturer based in McHenry, US. The company specializes in surgical implants, fixation systems, and trauma solutions. Biomet has maintained a strong FDA 510(k) regulatory record since its first clearance in 1978. The company has received FDA 510(k) clearances from total submissions. Orthopedic devices represent 88% of its submission portfolio, reflecting the company's core focus on joint reconstruction, trauma fixation, and surgical instrumentation. The latest clearance in 2024 demonstrates continued regulatory activity and ...
