FDA 510(k) Premarket Notification

K902788 CLEM TM

CLEARED

Sep 12, 1990
78 days to decision

K902788 · Product code: **FIB** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k902788/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Protector, Transducer, Dialysis (FIB)
Date received	Jun 26, 1990
Decision date	Sep 12, 1990
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Arbo Medical, Inc.**
Location	Wilton, CT, US
Contact	DAWN MOORE
510(k) history	15 submissions · 15 cleared · 1989-1995