FDA 510(k) Premarket Notification

K831766 MICROSTAR VCI

CLEARED

Oct 14, 1983
135 days to decision

K831766 · Product code: **FKX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k831766/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Jun 1, 1983
Decision date	Oct 14, 1983
Days to decision	135 days
Third-party review	No

APPLICANT

Company	**Medionics International , Ltd.**
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1982-1992