FDA 510(k) Premarket Notification

K823330 DIALATORS PC-203 & 202

CLEARED

Jan 26, 1983
79 days to decision

K823330 · Product code: **FKX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k823330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Nov 8, 1982
Decision date	Jan 26, 1983
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Medigroup**
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1983-1983