FDA 510(k) Premarket Notification

K862412 INTESTOPLANT

CLEARED

Nov 13, 1986
141 days to decision

K862412 · Product code: **LLD** · General Hospital
Source: https://www.510kdatabase.net/k862412/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intraperitoneal (LLD)
Date received	Jun 25, 1986
Decision date	Nov 13, 1986
Days to decision	141 days
Third-party review	No

APPLICANT

Company	**Burron Medical Products, Inc.**
Location	Mchenry, IL, US
Contact	TRACEY YAKABOW
510(k) history	41 submissions · 40 cleared · 1979-1987