FDA 510(k) Premarket Notification

K852259 IVENT INTRAVENOUS ADMINSTRATION SET

CLEARED

Jun 7, 1985
14 days to decision

K852259 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k852259/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 24, 1985
Decision date	Jun 7, 1985
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Ivent Corp.**
Location	Walker, MI, US
Contact	CHRIS OZIMEK
510(k) history	3 submissions · 3 cleared · 1983-1985