FDA 510(k) Premarket Notification

K823026 380 PIGGYBACK CONTROLLER

CLEARED

Nov 22, 1982
39 days to decision

K823026 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k823026/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Oct 14, 1982
Decision date	Nov 22, 1982
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**Imed Corp.**
Location	Mchenry, IL, US
510(k) history	43 submissions · 43 cleared · 1977-1996