FDA 510(k) Premarket Notification

K842007 CENTRIMED SYSTEM 1

CLEARED

Aug 3, 1984
78 days to decision

K842007 · Product code: **KFM** · Cardiovascular
Source: https://www.510kdatabase.net/k842007/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Blood, Cardiopulmonary Bypass, Non-roller Type (KFM)
Date received	May 17, 1984
Decision date	Aug 3, 1984
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Centrimed Corp.**
Location	Walker, MI, US
510(k) history	4 submissions · 4 cleared · 1984-1985