FDA 510(k) Premarket Notification

K902355 MCR4000 & MCR4000F CARDIOTOMY RESERVOIR

CLEARED

Aug 23, 1990
86 days to decision

K902355 · Product code: **DTP** · Cardiovascular
Source: https://www.510kdatabase.net/k902355/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Defoamer, Cardiopulmonary Bypass (DTP)
Date received	May 29, 1990
Decision date	Aug 23, 1990
Days to decision	86 days
Third-party review	No

APPLICANT

Company	**Medtronic Vascular**
Location	Walker, MI, US
Contact	LOWE, PHD
510(k) history	475 submissions · 453 cleared · 1977-2023