FDA 510(k) Premarket Notification

K880533 VITEL III CM

CLEARED

May 13, 1988
94 days to decision

K880533 · Product code: **DXH** · Cardiovascular
Source: https://www.510kdatabase.net/k880533/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Feb 9, 1988
Decision date	May 13, 1988
Days to decision	94 days
Third-party review	No

APPLICANT

Company	**Equimed Corp.**
Location	Plymouth, MN, US
Contact	DAVID C GUST
510(k) history	2 submissions · 2 cleared · 1987-1988