FDA 510(k) Premarket Notification

K772350 PATIENT ALERT MODEL 555

CLEARED

Mar 9, 1978
76 days to decision

K772350 · Product code: **DXH** · Cardiovascular
Source: https://www.510kdatabase.net/k772350/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Dec 23, 1977
Decision date	Mar 9, 1978
Days to decision	76 days
Third-party review	No

APPLICANT

Company	**Intermedics, Inc.**
Location	Mchenry, IL, US
510(k) history	211 submissions · 201 cleared · 1977-1996