FDA 510(k) Premarket Notification

K810755 SQUINN VITATEK 500 SERIES PHYSIOL. MONI

CLEARED

Apr 3, 1981
14 days to decision

K810755 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k810755/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 20, 1981
Decision date	Apr 3, 1981
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Vitatek**
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1982