FDA 510(k) Premarket Notification

K802205 ECG/RESPIRATION STIMULATOR

CLEARED

Oct 3, 1980
22 days to decision

K802205 · Product code: **BZQ** · Anesthesiology
Source: https://www.510kdatabase.net/k802205/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Sep 11, 1980
Decision date	Oct 3, 1980
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Healthdyne, Inc.**
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1978-1995