FDA 510(k) Premarket Notification

K833227 SONICAID MODEL D206E AIR EMBOLI-DETECT

CLEARED

Oct 27, 1983
72 days to decision

K833227 · Product code: **LBA** · Toxicology
Source: https://www.510kdatabase.net/k833227/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Procainamide Control Materials (LBA)
Date received	Aug 16, 1983
Decision date	Oct 27, 1983
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Sonicaid, Inc.**
Location	Mchenry, IL, US
510(k) history	18 submissions · 17 cleared · 1977-1988