FDA 510(k) Premarket Notification

K802030 U.I. OCTOSON

CLEARED

Oct 3, 1980
43 days to decision

K802030 · Product code: **DXK** · Cardiovascular
Source: https://www.510kdatabase.net/k802030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Aug 21, 1980
Decision date	Oct 3, 1980
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Ausonics Pty , Ltd.**
Location	Mchenry, IL, US
510(k) history	27 submissions · 26 cleared · 1979-1994