FDA 510(k) Premarket Notification

K832882 DIF-STET

CLEARED

Dec 6, 1983
102 days to decision

K832882 · Product code: **DQD** · Cardiovascular
Source: https://www.510kdatabase.net/k832882/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Aug 26, 1983
Decision date	Dec 6, 1983
Days to decision	102 days
Third-party review	No

APPLICANT

Company	**Intersect Systems, Inc.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1983-2002