FDA 510(k) Premarket Notification

K810940 THE BETA

CLEARED

Jul 16, 1981
100 days to decision

K810940 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k810940/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Apr 7, 1981
Decision date	Jul 16, 1981
Days to decision	100 days
Third-party review	No

APPLICANT

Company	**Burdick Corp.**
Location	Mchenry, IL, US
Website	https://www.cardinalhealth.com
510(k) history	38 submissions · 38 cleared · 1976-1994