FDA 510(k) Premarket Notification

K810460 BURRON PERCUTANEOUS INDUCER SET

CLEARED

Mar 11, 1981
19 days to decision

K810460 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k810460/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Feb 20, 1981
Decision date	Mar 11, 1981
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**Burron Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1979-1987