FDA 510(k) Premarket Notification

K811964 VISITEC NEEDLE RANGE (CARDIOVASCULAR)

CLEARED

Jul 20, 1981
10 days to decision

K811964 · Product code: **DRC** · Cardiovascular
Source: https://www.510kdatabase.net/k811964/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Trocar (DRC)
Date received	Jul 10, 1981
Decision date	Jul 20, 1981
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Visitec Co.**
Location	Walker, MI, US
510(k) history	49 submissions · 49 cleared · 1979-1995