FDA 510(k) Premarket Notification

K830022 S.M.P. CONTRAST MEDIA SET

CLEARED

Jan 21, 1983
17 days to decision

K830022 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k830022/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 4, 1983
Decision date	Jan 21, 1983
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Omega Technologies, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983