FDA 510(k) Premarket Notification

K813556 MOD. 587 BLOOD PRESSURE & PULSE MONITOR

CLEARED

Jan 22, 1982
31 days to decision

K813556 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k813556/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 22, 1981
Decision date	Jan 22, 1982
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**Bio Mega Diagnostic, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1976-1987