FDA 510(k) Premarket Notification

K063814 BP-200 PLUS

CLEARED

Mar 7, 2007
75 days to decision

K063814 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k063814/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 22, 2006
Decision date	Mar 7, 2007
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Schiller AG**
Location	Baar, CH
Contact	RETO KUETTEL
510(k) history	16 submissions · 16 cleared · 1985-2023