FDA 510(k) Premarket Notification

K831583 GP-303

CLEARED

Jul 19, 1983
63 days to decision

K831583 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k831583/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	May 17, 1983
Decision date	Jul 19, 1983
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Hirata Sangyo Co. USA, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1982-1986