FDA 510(k) Premarket Notification

CLEARED

K030358 KD-622 FULLY AUTOMATIC ELECTRONIC BLOOD PRESSURE MONITOR

Oct 31, 2003
270 days to decision

K030358 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k030358/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Feb 3, 2003
Decision date	Oct 31, 2003
Days to decision	270 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Kodon(Tianjin)Electronic&Electrical Apparatus Co.,**
Location	Tianjin, CN
Contact	LIU YI
510(k) history	9 submissions · 9 cleared · 2003-2007