FDA 510(k) Premarket Notification

K021240 EIKON AUTOMATIC DIGITAL BLOOD PRESSURE MONITOR, MODEL HD-200M/HD-200

CLEARED

Nov 8, 2002
204 days to decision

K021240 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k021240/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Apr 18, 2002
Decision date	Nov 8, 2002
Days to decision	204 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Eikon Healthcare Device Corp.**
Location	Hsin-Chu City,, TW
Contact	JEN KE-MIN
510(k) history	2 submissions · 2 cleared · 2002-2002