

**K021240 EIKON AUTOMATIC DIGITAL BLOOD PRESSURE MONITOR, MODEL HD-200M/HD-200**Nov 8, 2002  
204 days to decisionK021240 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k021240/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021240/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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