

**K831583 GP-303**Jul 19, 1983  
63 days to decisionK831583 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k831583/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k831583/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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