

**K813556 MOD. 587 BLOOD PRESSURE & PULSE MONITOR**Jan 22, 1982  
31 days to decisionK813556 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k813556/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 22, 1981
Decision date	Jan 22, 1982
Days to decision	31 days
Third-party review	No

**APPLICANT**

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Company	<b>Bio Mega Diagnostic, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1976-1987

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k813556/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026