

**K903379 PYMAH PRE-GAGED BALANCED TEMPA-KUFF**Oct 19, 1990  
81 days to decisionK903379 · Product code: **DXQ** · CardiovascularSource: <https://www.510kdatabase.net/k903379/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	Jul 30, 1990
Decision date	Oct 19, 1990
Days to decision	81 days
Third-party review	No

**APPLICANT**

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Company	<b>Pymah Corp.</b>
Location	Somerville, NJ, US
Contact	L KONOPKO
510(k) history	17 submissions · 17 cleared · 1988-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903379/>; 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