

**K760504 PLETHYSMOGRAPH (MDI SYSTEM II)**Aug 26, 1976  
3 days to decisionK760504 · Product code: **JOM** · CardiovascularSource: <https://www.510kdatabase.net/k760504/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Photoelectric, Pneumatic Or Hydraulic (JOM)
Date received	Aug 23, 1976
Decision date	Aug 26, 1976
Days to decision	3 days
Third-party review	No

**APPLICANT**

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Company	<b>Marketing Development Intl.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1976-1976

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k760504/>; 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 July 10, 2026