

**K833561 LIGHT REFLECTION RHEOGRAPHY**Jan 8, 1985  
454 days to decisionK833561 · Product code: **JOM** · CardiovascularSource: <https://www.510kdatabase.net/k833561/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Photoelectric, Pneumatic Or Hydraulic (JOM)
Date received	Oct 12, 1983
Decision date	Jan 8, 1985
Days to decision	454 days
Third-party review	No

**APPLICANT**

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Company	<b>Instru-Med, Inc.</b>
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1985-1985

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

Device record: <https://www.510kdatabase.net/k833561/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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