

**K832907 INVASIVE MONITORING KITS**Jan 10, 1984  
134 days to decisionK832907 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k832907/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Aug 29, 1983
Decision date	Jan 10, 1984
Days to decision	134 days
Third-party review	No

**APPLICANT**

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Company	<b>American Pharmaseal Div. Ahsc</b>
Location	Mchenry, IL, US
510(k) history	64 submissions · 61 cleared · 1980-1987

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

Device record: <https://www.510kdatabase.net/k832907/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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