

**K790034 INTRODUCERS**Feb 1, 1979  
29 days to decisionK790034 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k790034/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jan 3, 1979
Decision date	Feb 1, 1979
Days to decision	29 days
Third-party review	No

**APPLICANT**

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Company	<b>Clinical Instruments Corp.</b>
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1977-1991

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

Device record: <https://www.510kdatabase.net/k790034/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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