

**K840022 HEART CATHETERIZATION TRAY SET**Feb 4, 1984  
30 days to decisionK840022 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k840022/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Jan 5, 1984
Decision date	Feb 4, 1984
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Medi-Plast Intl., Inc.</b>
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1981-1996

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

Device record: <https://www.510kdatabase.net/k840022/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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