

**K820788 CLIRANS PRIMING SET**Apr 6, 1982  
14 days to decisionK820788 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k820788/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 23, 1982
Decision date	Apr 6, 1982
Days to decision	14 days
Third-party review	No

**APPLICANT**

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Company	<b>Terumo Medical Corp.</b>
Location	Elkton, MD, US
510(k) history	143 submissions · 143 cleared · 1980-2011

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

Device record: <https://www.510kdatabase.net/k820788/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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