

**K830609 INFLATION DEVICE**Mar 24, 1983  
27 days to decisionK830609 · Product code: **DXT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k830609/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Feb 25, 1983
Decision date	Mar 24, 1983
Days to decision	27 days
Third-party review	No

**APPLICANT**

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Company	<b>Interventional Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-1988

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Device record: <https://www.510kdatabase.net/k830609/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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