

**K823026 380 PIGGYBACK CONTROLLER**Nov 22, 1982  
39 days to decisionK823026 · Product code: **LDR** · General HospitalSource: <https://www.510kdatabase.net/k823026/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Oct 14, 1982
Decision date	Nov 22, 1982
Days to decision	39 days
Third-party review	No

**APPLICANT**

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Company	<b>Imed Corp.</b>
Location	Mchenry, IL, US
510(k) history	43 submissions · 43 cleared · 1977-1996

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

Device record: <https://www.510kdatabase.net/k823026/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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