

**K810460 BURRON PERCUTANEOUS INDUCER SET**Mar 11, 1981  
19 days to decisionK810460 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k810460/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Feb 20, 1981
Decision date	Mar 11, 1981
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>Burron Medical Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1979-1987

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

Device record: <https://www.510kdatabase.net/k810460/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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