

**K810940 THE BETA**Jul 16, 1981  
100 days to decisionK810940 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k810940/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Apr 7, 1981
Decision date	Jul 16, 1981
Days to decision	100 days
Third-party review	No

**APPLICANT**

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Company	<b>Burdick Corp.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.cardinalhealth.com">https://www.cardinalhealth.com</a>
510(k) history	38 submissions · 38 cleared · 1976-1994

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