

**K802030 U.I. OCTOSON**Oct 3, 1980  
43 days to decisionK802030 · Product code: **DXK** · CardiovascularSource: <https://www.510kdatabase.net/k802030/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Aug 21, 1980
Decision date	Oct 3, 1980
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Ausonics Pty , Ltd.</b>
Location	Mchenry, IL, US
510(k) history	27 submissions · 26 cleared · 1979-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k802030/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 20, 2026