

**K802672 CARDIOACTIVE DRUGS CONTROL**Nov 24, 1980  
28 days to decisionK802672 · Product code: **DIF** · Toxicology  
Source: <https://www.510kdatabase.net/k802672/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Oct 27, 1980
Decision date	Nov 24, 1980
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Fisher Scientific Co., LLC</b>
Location	Mchenry, IL, US
510(k) history	89 submissions · 89 cleared · 1976-1993

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Device record: <https://www.510kdatabase.net/k802672/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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