

**K840044 SDI PREPAX DILUKIT 29-1000**Mar 9, 1984  
63 days to decisionK840044 · Product code: **JJO** · Toxicology  
Source: <https://www.510kdatabase.net/k840044/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flame Emission Photometer For Clinical Use (JJO)
Date received	Jan 6, 1984
Decision date	Mar 9, 1984
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>Scientific Distributors, Inc.</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1984-1989

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

Device record: <https://www.510kdatabase.net/k840044/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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