

**K860097 THERAPEUTIC DRUG (TDM) CALIBRATOR**Feb 11, 1986  
32 days to decisionK860097 · Product code: **DIF** · Toxicology  
Source: <https://www.510kdatabase.net/k860097/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Jan 10, 1986
Decision date	Feb 11, 1986
Days to decision	32 days
Third-party review	No

**APPLICANT**

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Company	<b>Instrumentation Laboratory CO</b>
Location	Mchenry, IL, US
Contact	GEORGE A LYNA
510(k) history	321 submissions · 320 cleared · 1976-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860097/>; 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 June 15, 2026