

**K820201 DISPOS. MONITORING SYS/LI ET AL**Feb 12, 1982  
17 days to decisionK820201 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k820201/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Jan 26, 1982
Decision date	Feb 12, 1982
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Linton Biomed Corp.</b>
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k820201/> 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 29, 2026