

**K862412 INTESTOPLANT**Nov 13, 1986  
141 days to decisionK862412 · Product code: **LLD** · General HospitalSource: <https://www.510kdatabase.net/k862412/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intraperitoneal (LLD)
Date received	Jun 25, 1986
Decision date	Nov 13, 1986
Days to decision	141 days
Third-party review	No

**APPLICANT**

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Company	<b>Burron Medical Products, Inc.</b>
Location	Mchenry, IL, US
Contact	TRACEY YAKABOW
510(k) history	41 submissions · 40 cleared · 1979-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k862412/>; 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 27, 2026