

**K864629 MODIFIED DUAL LUMEN NEEDLE**Dec 16, 1986  
21 days to decisionK864629 · Product code: **LBW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k864629/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Dialysis, Single Needle (co-axial Flow) (LBW)
Date received	Nov 25, 1986
Decision date	Dec 16, 1986
Days to decision	21 days
Third-party review	No

**APPLICANT**

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Company	<b>Quinton, Inc.</b>
Location	Mchenry, IL, US
Contact	MARY-LEE DONOGHUE
510(k) history	164 submissions · 160 cleared · 1976-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k864629/>; 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 20, 2026