

**K893420 MASTERLAB PFT AND MASTERLAB BODY BOX**Jun 23, 1989  
57 days to decisionK893420 · Product code: **CCM** · AnesthesiologySource: <https://www.510kdatabase.net/k893420/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Pressure (CCM)
Date received	Apr 27, 1989
Decision date	Jun 23, 1989
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Quinton, Inc.</b>
Location	Mchenry, IL, US
Contact	RON R DUCK
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/k893420/>, 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