

**K833397 TRI-MED 511 INFANT RESPIRATION MONITOR**Nov 28, 1983  
59 days to decisionK833397 · Product code: **CCK** · AnesthesiologySource: <https://www.510kdatabase.net/k833397/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Sep 30, 1983
Decision date	Nov 28, 1983
Days to decision	59 days
Third-party review	No

**APPLICANT**

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Company	<b>Tri-Med, Inc.</b>
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1977-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k833397/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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