

**K840841 RENEAU UNIT**Aug 7, 1984  
165 days to decisionK840841 · Product code: **CBF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k840841/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chamber, Hyperbaric (CBF)
Date received	Feb 24, 1984
Decision date	Aug 7, 1984
Days to decision	165 days
Third-party review	No

**APPLICANT**

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Company	<b>Reneau, Inc.</b>
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1984-1984

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

Device record: <https://www.510kdatabase.net/k840841/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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