

**K801330 ANSPACH LEG SUPPORT**Jun 26, 1980  
23 days to decisionK801330 · Product code: **CCX** · AnesthesiologySource: <https://www.510kdatabase.net/k801330/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Support, Patient Position (CCX)
Date received	Jun 3, 1980
Decision date	Jun 26, 1980
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>The Anspach Effort, Inc.</b>
Location	Mchenry, IL, US
510(k) history	60 submissions · 60 cleared · 1980-2022

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

Device record: <https://www.510kdatabase.net/k801330/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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