

**K781761 KENDRICK EXTRICATION DEVICE**Oct 27, 1978  
11 days to decisionK781761 · Product code: **IQF** · Physical MedicineSource: <https://www.510kdatabase.net/k781761/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Cervical-thoracic, Rigid (IQF)
Date received	Oct 16, 1978
Decision date	Oct 27, 1978
Days to decision	11 days
Third-party review	No

**APPLICANT**

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Company	<b>Medi-Ked, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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

Device record: <https://www.510kdatabase.net/k781761/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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