

**K080072 MRI PATIENT POSITIONING DEVICES**Jun 4, 2008  
145 days to decisionK080072 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k080072/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jan 11, 2008
Decision date	Jun 4, 2008
Days to decision	145 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>CIVCO Medical Instruments Co., Inc.</b>
Location	Walker, MI, US
Contact	ARTHUR WARD
510(k) history	29 submissions · 29 cleared · 1982-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k080072/> 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 17, 2026