

**K900741 BIODEX BACK SYSTEM ATTACHMENT**Feb 26, 1990  
10 days to decisionK900741 · Product code: **IKK** · Physical MedicineSource: <https://www.510kdatabase.net/k900741/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Feb 16, 1990
Decision date	Feb 26, 1990
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Biodex Corp.</b>
Location	Center Moriches, NY, US
Contact	JAMES REISS
510(k) history	7 submissions · 7 cleared · 1986-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900741/>, 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 June 30, 2026