

**K880230 SIXALYZER**Mar 16, 1988  
56 days to decisionK880230 · Product code: **IKK** · Physical MedicineSource: <https://www.510kdatabase.net/k880230/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Jan 20, 1988
Decision date	Mar 16, 1988
Days to decision	56 days
Third-party review	No

**APPLICANT**

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Company	<b>Lukens Corp.</b>
Location	Mchenry, IL, US
Contact	LUTZ KAUFFMANN
510(k) history	11 submissions · 11 cleared · 1977-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880230/>, 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