

**K792032 LEIF CENTRIFUGAL CYTOLOGY BUCKET**Nov 30, 1979  
51 days to decisionK792032 · Product code: **IFB** · Pathology  
Source: <https://www.510kdatabase.net/k792032/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cytocentrifuge (IFB)
Date received	Oct 10, 1979
Decision date	Nov 30, 1979
Days to decision	51 days
Third-party review	No

**APPLICANT**

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Company	<b>Robert C. Leif, Ph.D.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979

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

Device record: <https://www.510kdatabase.net/k792032/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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