

**K822823 HEMATOLOGY COMPUTER-INTERFACE, VISCOUNT**Oct 18, 1982  
27 days to decisionK822823 · Product code: **LKM** · Hematology  
Source: <https://www.510kdatabase.net/k822823/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Counter, Urine Particle (LKM)
Date received	Sep 21, 1982
Decision date	Oct 18, 1982
Days to decision	27 days
Third-party review	No

**APPLICANT**

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Company	<b>Eko Scann AB</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983

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

Device record: <https://www.510kdatabase.net/k822823/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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