

**K823761 ERYTHROCYTE & STAND SEDIMENT. RATE TUBE**Jan 21, 1983  
38 days to decisionK823761 · Product code: **JPH** · Hematology  
Source: <https://www.510kdatabase.net/k823761/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Test, Erythrocyte Sedimentation Rate (JPH)
Date received	Dec 14, 1982
Decision date	Jan 21, 1983
Days to decision	38 days
Third-party review	No

**APPLICANT**

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Company	<b>Cordis Corp.</b>
Location	Mchenry, IL, US
Website	<a href="https://cordis.com">https://cordis.com</a>
510(k) history	315 submissions · 281 cleared · 1976-2014

Cordis Corp. is a medical device manufacturer based in McHenry, US. The company specializes in interventional cardiovascular and gastroenterology devices. Cordis has a substantial FDA 510(k) regulatory history spanning from 1976 to 2014. The company received FDA 510(k) clearances from total submissions. Its portfolio focuses primarily on cardiovascular devices and gastroenterology stent systems, including percutaneous transluminal angioplasty catheters, emboli capture guidewires, and self-expanding biliary stent systems. Notable cleared products include the FLEXSTENT Bili...

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