

**K810323 ACRYTOL, MOUNTING MEDIUM**Mar 6, 1981  
28 days to decisionK810323 · Product code: **KEP** · Pathology  
Source: <https://www.510kdatabase.net/k810323/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Mounting, Oil Soluble (KEP)
Date received	Feb 6, 1981
Decision date	Mar 6, 1981
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Surgipath</b>
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k810323/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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