

**K770767 IN VITRO DIAG. FOR URIC ACID**Jul 5, 1977  
68 days to decisionK770767 · Product code: **CDO** · Chemistry  
Source: <https://www.510kdatabase.net/k770767/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Acid, Uric, Uricase (u.v.) (CDO)
Date received	Apr 28, 1977
Decision date	Jul 5, 1977
Days to decision	68 days
Third-party review	No

**APPLICANT**

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Company	<b>Smithkline Diagnostics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	42 submissions · 42 cleared · 1976-1996

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Device record: <https://www.510kdatabase.net/k770767/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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