

**K813413 LECTIN H**Mar 4, 1982  
87 days to decisionK813413 · Product code: **KSI** · Pathology  
Source: <https://www.510kdatabase.net/k813413/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lectins And Protectins (KSI)
Date received	Dec 7, 1981
Decision date	Mar 4, 1982
Days to decision	87 days
Third-party review	No

**APPLICANT**

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Company	<b>American Dade</b>
Location	Mchenry, IL, US
510(k) history	149 submissions · 149 cleared · 1980-1987

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Device record: <https://www.510kdatabase.net/k813413/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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