FDA 510(k) Premarket Notification

K813413 LECTIN H

CLEARED

Mar 4, 1982
87 days to decision

K813413 · Product code: **KSI** · Pathology
Source: https://www.510kdatabase.net/k813413/

SUBMISSION DETAILS

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

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