

K840338 HAEMOPHILUS I.D. II TRIPLATEApr 17, 1984
83 days to decisionK840338 · Product code: **JSI** · Microbiology
Source: <https://www.510kdatabase.net/k840338/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Culture Media, Selective And Differential (JSI)
Date received	Jan 25, 1984
Decision date	Apr 17, 1984
Days to decision	83 days
Third-party review	No

APPLICANT

Company	Remel Co.
Location	Mchenry, IL, US
510(k) history	137 submissions · 137 cleared · 1979-2001

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Device record: <https://www.510kdatabase.net/k840338/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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