

K770830 PLATELET AGITATORJul 11, 1977
66 days to decisionK770830 · Product code: **GLE** · Hematology
Source: <https://www.510kdatabase.net/k770830/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixer, Blood Tube (GLE)
Date received	May 6, 1977
Decision date	Jul 11, 1977
Days to decision	66 days
Third-party review	No

APPLICANT

Company	Helmer Labs, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1977-1981

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Device record: <https://www.510kdatabase.net/k770830/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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