

K831253 LANCER HEMOGLOBIN ANALYZER REAGENTSMay 9, 1983
21 days to decisionK831253 · Product code: **KMG** · Hematology
Source: <https://www.510kdatabase.net/k831253/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Purifier, Water, Ultraviolet, Medical (KMG)
Date received	Apr 18, 1983
Decision date	May 9, 1983
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Sherwood Medical Co.
Location	Mchenry, IL, US
510(k) history	191 submissions · 177 cleared · 1976-1998

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Device record: <https://www.510kdatabase.net/k831253/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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