

K813204 SERACULT OCCULT BLOOF TESTJan 28, 1982
76 days to decisionK813204 · Product code: **KHE** · Hematology
Source: <https://www.510kdatabase.net/k813204/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reagent, Occult Blood (KHE)
Date received	Nov 13, 1981
Decision date	Jan 28, 1982
Days to decision	76 days
Third-party review	No

APPLICANT

Company	Propper Mfg. Co., Inc.
Location	Mchenry, IL, US
510(k) history	25 submissions · 25 cleared · 1977-2012

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Device record: <https://www.510kdatabase.net/k813204/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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