FDA 510(k) Premarket Notification

# K813204 SERACULT OCCULT BLOOF TEST

**CLEARED**

Jan 28, 1982
76 days to decision

K813204 · 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 |