FDA 510(k) Premarket Notification

# K830127 QUIK-DIP BLOOD STAIN

**CLEARED**

Feb 15, 1983
34 days to decision

K830127 · Product code: **IAF** · Pathology
Source: https://www.510kdatabase.net/k830127/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Wright's Stain (IAF) |
| Date received | Jan 12, 1983 |
| Decision date | Feb 15, 1983 |
| Days to decision | 34 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Mercedes Scientific Co., Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 2 submissions · 2 cleared · 1983-1983 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k830127/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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