FDA 510(k) Premarket Notification

# K890844 MICROSEPT 60/20

**CLEARED**

Jun 7, 1989
106 days to decision

K890844 · Product code: **CAK** · General Hospital
Source: https://www.510kdatabase.net/k890844/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Microfilter, Blood Transfusion (CAK) |
| Date received | Feb 21, 1989 |
| Decision date | Jun 7, 1989 |
| Days to decision | 106 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Arbor Technologies, Inc.** |
| Location | Ann Arbor, MI, US |
| Contact | DAWN I MOORE |
| 510(k) history | 10 submissions · 10 cleared · 1986-1997 |