

K890109 SAFEPETTE DISPENSERFeb 23, 1989
44 days to decisionK890109 · Product code: **JKA** · Chemistry
Source: <https://www.510kdatabase.net/k890109/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA) |
| Date received | Jan 10, 1989 |
| Decision date | Feb 23, 1989 |
| Days to decision | 44 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Safe-Tec Clinical Products, Inc. |
| Location | Ivyland, PA, US |
| Contact | G KENDRICK |
| 510(k) history | 2 submissions · 2 cleared · 1988-1989 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890109/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026