

K936201 URINARY ALBUMIN CALIBRATORMar 11, 1994
72 days to decisionK936201 · Product code: **JIS** · Chemistry
Source: <https://www.510kdatabase.net/k936201/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Calibrator, Primary (JIS) |
| Date received | Dec 29, 1993 |
| Decision date | Mar 11, 1994 |
| Days to decision | 72 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Dupont Medical Products |
| Location | Wilmington, DE, US |
| Contact | REBECCA S AYASH |
| 510(k) history | 28 submissions · 28 cleared · 1991-1995 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k936201/> 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 May 20, 2026