

K252040 BD Vacutainer® Plasma Separator Tubes (PST™), BD Vacutainer® Sodium Heparin Blood Collection TubesMar 13, 2026
256 days to decisionK252040 · Product code: JKA · Chemistry
Source: <https://www.510kdatabase.net/k252040/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA) |
| Date received | Jun 30, 2025 |
| Decision date | Mar 13, 2026 |
| Days to decision | 256 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Becton, Dickinson and Company |
| Location | Franklin Lakes, NJ, US |
| Contact | Angela Mariani |
| Website | https://www.bd.com |
| 510(k) history | 134 submissions · 134 cleared · 2010-2026 |

Becton, Dickinson and Company is an American multinational medical technology company headquartered in Franklin Lakes, New Jersey. BD manufactures and sells medical devices, instrument systems, and reagents globally. The company maintains a strong FDA 510(k) regulatory record with FDA 510(k) clearances from total submissions spanning 2010 to 2026. BD's cleared devices span multiple categories including microbiology systems, blood collection products, and general hospital devices. The company's latest clearance in 2026 reflects continued innovation and regulatory engagement...