

K092172 EXEL I.V. EXTENSION SETJan 4, 2010
167 days to decisionK092172 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k092172/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Set, Administration, Intravascular (FPA) |
| Date received | Jul 21, 2009 |
| Decision date | Jan 4, 2010 |
| Days to decision | 167 days |
| Third-party review | No |
| Summary / Statement | Statement |

APPLICANT

| | |
|----------------|---|
| Company | Exelint Intl. Co. |
| Location | Culver City, CA, US |
| Contact | ARMAND HAMID |
| 510(k) history | 12 submissions · 12 cleared · 2000-2010 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092172/> 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 21, 2026