

K802999 BETA-TEC QUALITATIVEDec 22, 1980
27 days to decisionK802999 · Product code: **JHI** · Chemistry
Source: <https://www.510kdatabase.net/k802999/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Visual, Pregnancy Hcg, Prescription Use (JHI) |
| Date received | Nov 25, 1980 |
| Decision date | Dec 22, 1980 |
| Days to decision | 27 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Armkel, LLC |
| Location | Mchenry, IL, US |
| 510(k) history | 68 submissions · 68 cleared · 1979-2004 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k802999/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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