

**K992638 DINAMAP PRO SERIES MONITOR, MODELS 100, 200,
300, 400**Feb 24, 2000
202 days to decisionK992638 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k992638/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI) |
| Date received | Aug 6, 1999 |
| Decision date | Feb 24, 2000 |
| Days to decision | 202 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Critikon Company, LLC |
| Location | Mchenry, IL, US |
| Contact | THOMAS ENGLISH |
| 510(k) history | 51 submissions · 51 cleared · 1979-2000 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992638/>; 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 26, 2026