

**K020288 PADPRO, MODEL 2603**Feb 27, 2002  
30 days to decisionK020288 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k020288/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                     |
| Submission type       | Special  |
| Device classification | Automated External Defibrillators (non-wearable) (MKJ) |
| Date received         | Jan 28, 2002   |
| Decision date         | Feb 27, 2002   |
| Days to decision      | 30 days  |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Padpro, LLC</b>                    |
| Location       | Deer Field, IL, US                    |
| Contact        | DANIEL KAMM                           |
| 510(k) history | 5 submissions · 5 cleared · 2001-2002 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020288/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 3, 2026