

**K960473 AESCULAP POWER SYSTEM HANDPIECE**Mar 7, 1996  
35 days to decisionK960473 · Product code: **GET** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k960473/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)                  |
| Submission type       | Traditional   |
| Device classification | Motor, Surgical Instrument, Pneumatic Powered (GET) |
| Date received         | Feb 1, 1996   |
| Decision date         | Mar 7, 1996   |
| Days to decision      | 35 days   |
| Third-party review    | No  |
| Summary / Statement   | Summary   |

**APPLICANT**

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|----------------|---|
| Company        | <b>Aesculap, Inc.</b>                     |
| Location       | Burlingame, CA, US                        |
| Contact        | MARY E HOLDEN                             |
| 510(k) history | 207 submissions · 201 cleared · 1991-2025 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960473/> 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 June 28, 2026