

**K830572 CLEAN-FLUSH**Mar 17, 1983  
22 days to decisionK830572 · Product code: **KYZ** · General Hospital  
Source: <https://www.510kdatabase.net/k830572/>**SUBMISSION DETAILS**

---

|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)     |
| Submission type       | Traditional                            |
| Device classification | Syringe, Irrigating (non Dental) (KYZ) |
| Date received         | Feb 23, 1983                           |
| Decision date         | Mar 17, 1983                           |
| Days to decision      | 22 days                                |
| Third-party review    | No                                     |

**APPLICANT**

---

|                |   |
|----------------|---|
| Company        | <b>Burron Medical Products, Inc.</b>    |
| Location       | Mchenry, IL, US                         |
| 510(k) history | 41 submissions · 40 cleared · 1979-1987 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830572/> 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 May 27, 2026