

**K091022 MODIFICATION TO RESERT XL HLD HIGH LEVEL
DISINFECTANT**Apr 12, 2010
368 days to decisionK091022 · Product code: **MED** · General Hospital
Source: <https://www.510kdatabase.net/k091022/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Sterilant, Medical Devices (MED) |
| Date received | Apr 9, 2009 |
| Decision date | Apr 12, 2010 |
| Days to decision | 368 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | STERIS Corporation |
| Location | Mentor, OH, US |
| Contact | JOHN SCOVILLE JR. |
| 510(k) history | 204 submissions · 202 cleared · 1997-2026 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091022/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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