

**K260181 LF Process Indicator Tape for Steam Sterilization**Feb 20, 2026  
30 days to decisionK260181 · Product code: **JOJ** · General Hospital  
Source: <https://www.510kdatabase.net/k260181/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                       |
| Submission type       | Special  |
| Device classification | Indicator, Physical/chemical Sterilization Process (JOJ) |
| Date received         | Jan 21, 2026   |
| Decision date         | Feb 20, 2026   |
| Days to decision      | 30 days  |
| Third-party review    | No   |
| Combination product   | No   |
| PCCP authorized       | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Intertape Polymer Group</b>        |
| Location       | Sarasota, FL, US                      |
| Contact        | Michael Johnson                       |
| 510(k) history | 1 submissions · 1 cleared · 2026-2026 |

**REGULATORY CONSULTANT**

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|-----------------|-----------------------|
| Consulting firm | <b>FLX Consulting</b> |
| Contact         | Mike Nolan            |

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k260181/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 13, 2026