FDA 510(k) Premarket Notification

CLEARED

K260181 LF Process Indicator Tape for Steam Sterilization

Feb 20, 2026
30 days to decision

K260181 · Product code: **JOJ** · General Hospital
Source: https://www.510kdatabase.net/k260181/

SUBMISSION DETAILS

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

Company	**Intertape Polymer Group**
Location	Sarasota, FL, US
Contact	Michael Johnson
510(k) history	1 submissions · 1 cleared · 2026-2026

REGULATORY CONSULTANT

Consulting firm	**FLX Consulting**
Contact	Mike Nolan

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