FDA 510(k) Premarket Notification

CLEARED

K241731 Outlook Surgical Versa One System (8900139)

Aug 11, 2025
420 days to decision

K241731 · Product code: **EOB** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k241731/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nasopharyngoscope (flexible Or Rigid) (EOB)
Date received	Jun 17, 2024
Decision date	Aug 11, 2025
Days to decision	420 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Resnent, LLC**
Location	Bloomington, IL, US
Contact	Willard Noyes
510(k) history	1 submissions · 1 cleared · 2025-2025

REGULATORY CONSULTANT

Consulting firm	**Mededge Consulting**
Contact	Billi-Jo Pfalzgraf

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