FDA 510(k) Premarket Notification

K260572 Marrow Marxman

CLEARED

Aug 4, 2026
166 days to decision

K260572 · Product code: **KNW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k260572/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Feb 19, 2026
Decision date	Aug 4, 2026
Days to decision	166 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Medimetrix, LLC**
Location	Germantown, TN, US
Contact	Brian Austin
510(k) history	1 submissions · 1 cleared · 2026-2026

REGULATORY CONSULTANT

Consulting firm	**MRC Global**
Contact	Christine Scifert

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