FDA 510(k) Premarket Notification

K261038 Xpert PFP

CLEARED

May 29, 2026
60 days to decision

K261038 · Product code: **HRS** · Orthopedic
Source: https://www.510kdatabase.net/k261038/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Mar 30, 2026
Decision date	May 29, 2026
Days to decision	60 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Newclip Technics**
Location	Haute-Goulaine, FR
Contact	Gaëlle Gourbière
Website	https://www.newcliptechnics.com
510(k) history	32 submissions · 32 cleared · 2006-2026

Newclip Technics is a French manufacturer of orthopedic osteosynthesis solutions based in Haute-Goulaine, near Nantes. The company designs, manufactures, and markets surgical instruments and implants for hospitals and clinics worldwide. The company has received FDA 510(k) clearances from total submissions since its first clearance in 2006. All submissions focus on orthopedic devices. The latest clearance in 2026 confirms the company remains actively engaged in regulatory submissions and product innovation. Recent cleared devices include patient-matched instrumentation sys...

REGULATORY CONSULTANT

Consulting firm	**The OrthoMedix Group, Inc.**
Contact	J.D. Webb

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k261038/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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