

K230104 Anthogyr AXIOM® BL X3 implants, Anthogyr Multi-Unit components for AXIOM® BLOct 16, 2023
276 days to decisionK230104 · Product code: DZE · Dental
Source: <https://www.510kdatabase.net/k230104/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 13, 2023
Decision date	Oct 16, 2023
Days to decision	276 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Anthogyr
Location	Sallanches, FR
Contact	Estelle Salle
510(k) history	10 submissions · 10 cleared · 2004-2024

REGULATORY CONSULTANT

Consulting firm	Straumann USA, LLC
Contact	Jennifer M. Jackson

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230104/>; Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026