

K223122 MENIX®Aug 3, 2023
304 days to decisionK223122 · Product code: **GAT** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k223122/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable, Synthetic, Polyethylene (GAT)
Date received	Oct 3, 2022
Decision date	Aug 3, 2023
Days to decision	304 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	MENIX® DUO

APPLICANT

Company	S.B.M. Sas (Science & Bio Materials)
Location	Lourdes, FR
Contact	Anne Cospin-Latapie
510(k) history	2 submissions · 2 cleared · 2020-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223122/> 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 27, 2026