

**K240108 Wrap Accessory Electrodes (GMX-ABSBELT01)**Jun 10, 2024  
146 days to decisionK240108 · Product code: **GXY** · Physical MedicineSource: <https://www.510kdatabase.net/k240108/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jan 16, 2024
Decision date	Jun 10, 2024
Days to decision	146 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	GMX-ABSBELT02; GMX-WRIST01; GMX-ELBOW01; GMX-LEG01, GMX-ANKLE01, GMX-KNEE01; GMX-FOOT PAD01; GMX-GLOVES01; SHOULDER01; GMX-NCK01)

**APPLICANT**

---

Company	<b>Gymmax Technology Shenzhen Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Benson Wang
510(k) history	1 submissions · 1 cleared · 2024-2024

**REGULATORY CONSULTANT**

---

Consulting firm	<b>Imd Medical &amp; Drug Technology Service Institutions</b>
Contact	Salon Chen

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.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k240108/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 14, 2026