

**K232779 QUEX ED**Oct 11, 2024  
396 days to decisionK232779 · Product code: **OMC** · Neurology  
Source: <https://www.510kdatabase.net/k232779/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reduced- Montage Standard Electroencephalograph (OMC)
Date received	Sep 11, 2023
Decision date	Oct 11, 2024
Days to decision	396 days
Third-party review	No
Summary / Statement	Summary
Other names	QUEX S

**APPLICANT**

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Company	<b>Qx World, Ltd.</b>
Location	Budapest, HU
Contact	Lilla Strobel
510(k) history	1 submissions · 1 cleared · 2024-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k232779/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026