

**K172450 TENS device-HeadTerm, eEspress**Sep 13, 2018  
395 days to decisionK172450 · Product code: **PCC** · Neurology  
Source: <https://www.510kdatabase.net/k172450/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Electrical, Transcutaneous, For Migraine (PCC)
Date received	Aug 14, 2017
Decision date	Sep 13, 2018
Days to decision	395 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Wat Medical Technology (Ningbo) Co., Ltd.</b>
Location	Ningbo, CN
Contact	Rodney Zhang
510(k) history	2 submissions · 2 cleared · 2018-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172450/> 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 June 18, 2026