

**K130273 PANPAC VAGINAL DILATORS**Aug 8, 2013  
185 days to decisionK130273 · Product code: **HDX** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k130273/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vaginal (HDX)
Date received	Feb 4, 2013
Decision date	Aug 8, 2013
Days to decision	185 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Panpac Medical Corporation</b>
Location	Shi-Chih City, Taipei Hsien, TW
Contact	Yen-Ming Pan
510(k) history	12 submissions · 12 cleared · 2010-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130273/> 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 20, 2026