

**K223918 AltaTrack Equipment R1.2, AltaTrack Guidewire,  
AltaTrack 3D Hub**Jun 8, 2023  
161 days to decisionK223918 · Product code: **DQK** · Cardiovascular  
Source: <https://www.510kdatabase.net/k223918/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Dec 29, 2022
Decision date	Jun 8, 2023
Days to decision	161 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medical Systems Nederland B.V.</b>
Location	Best, NL
Contact	Jeanette Becker
510(k) history	103 submissions · 102 cleared · 2005-2026

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223918/> 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 13, 2026