# **Rivaroxaban** Quality Solutions

Category: Anticoagulants



### MONOGRAPH 1

<b>RIVAROXABAN</b> Official as of 01-Dec-2022	
USP Rivaroxaban R-Enantiomer RS	<u>USP-1604541</u>
USP Rivaroxaban RS	<u>USP-1604530</u>
USP Rivaroxaban Related Compound B RS	<u>USP-1604552</u>
USP Rivaroxaban Related Compound D RS	<u>USP-1604563</u>
USP Rivaroxaban Related Compound G RS	<u>USP-1604596</u>
USP Rivaroxaban Related Compound J RS	<u>USP-1604574</u>

### MONOGRAPH 2

<b>RIVAROXABAN TABLETS  </b> NEW Official as of 01-May-2023		
USP Rivaroxaban RS	<u>USP-1604530</u>	
USP Rivaroxaban Related Compound H RS	<u>USP-1604585</u>	

\* Documents in PF Online are not official and not suitable to demonstrate compliance. They may never become official.

#### PHARMACEUTICAL ANALYTICAL IMPURITY(IES)\* USP Rivaroxaban Glycolic Acid Analog PAI USP-1A01170 USP Rivaroxaban Open Oxomorpholine Ring USP-1A02240 PAI USP Rivaroxaban Open Ring Methylamine USP-1A00650 PAI USP Rivaroxaban Phthalamide PAI USP-1A00660 USP Rivaroxaban Open Ring PAI USP-1A00670 USP Rivaroxaban Amine PAI USP-1A01160 USP Chlorothiophene Carboxylic Acid PAI USP-1A01180 USP Rivaroxaban N-Acyl Glycolic Acid PAI USP-1A01190 USP Chloro N-methylthiophene USP-1A01450 Carboxamide PAI USP Rivaroxaban Open Chain Dithiophene USP-1A01460 Analog PAI USP Rivaroxaban Desmorpholinone Analog USP-1A01470 PAI

01-Oct-2023. For publications related to **Rivaroxaban** in *PF* online after this date, please visit the integrated *USP-NF/PF* online.



\* Pharmaceutical Analytical Impurities (PAI products) are released using a process developed by USP's subject matter experts. The release process is based on internal policies, standard operating procedures, and requirements as defined by USP's Quality Management System. USP is an ISO 9001:2015 certified facility. PAI products are different from official USP Reference Standards. PAI products are not required for compendial compliance.

**Disclaimer:** USP Quality Solution Sheets are provided as a convenience and for informational purposes only. Labmix24 makes reasonable efforts to provide correct information but assumes no liability for the completeness, timeliness or accuracy of the information contained in each QSS. It is the responsibility of the user to verify the information. The US Pharmacopeia remains the official source of this information.

## www.labmix24.com

# **Rivaroxaban** Quality Solutions

Category: Anticoagulants



PHARMACEUTICAL ANALYTICAL IMPURITY(IES)*		
USP Desthiophenerivaroxaban Methylcarba- mate Analog PAI	<u>USP-1A01480</u>	
USP Methylisoindolinedione PAI	<u>USP-1A01490</u>	
USP Desthiophenerivaroxaban Phthalimodo Analog PAI	<u>USP-1A01500</u>	
USP Rivaroxaban Desthiophene Analog PAI	<u>USP-1A01510</u>	
USP Rivaroxaban N-methylphthalamido Analog PAI	<u>USP-1A01520</u>	
USP Aminophenylmorpholinone PAI	<u>USP-1A01530</u>	
USP (S)-Glycidyl Phthalimide PAI	<u>USP-1A01540</u>	
USP Rivaroxaban Open Ring S-isomer PAI	<u>USP-1A01550</u>	
USP Chlorothiophene Aldehyde PAI	<u>USP-1A01560</u>	
USP Rivaroxaban Phthalimido Analog PAI	<u>USP-1A01570</u>	
USP Dichlorothiophene Carboxylic Acid PAI	<u>USP-1A01580</u>	
USP Nitrophenylcholrothiophene Carboxyl- ate PAI	<u>USP-1A01590</u>	
USP Dihthalimido Morpholinone Analog PAI	<u>USP-1A01600</u>	
USP Aminophenylmorpholin-2-one PAI	<u>USP-1A01610</u>	
USP Rivaroxaban Ethyl Carbamate PAI	<u>USP-1A02250</u>	
USP Dichlorothienoic Acid PAI	<u>USP-1A02260</u>	
USP Rivaroxaban Phthalimide PAI	<u>USP-1A02270</u>	
USP Rivaroxaban Isomer PAI	<u>USP-1A02280</u>	
USP Rivaroxaban Phthalimido Ethyl Ester Analog PAI	<u>USP-1A02290</u>	
USP Rivaroxaban Ethylamino Analog PAI	<u>USP-1A02300</u>	

01-Oct-2023. For publications related to **Rivaroxaban** in *PF* online after this date, please visit the integrated *USP-NF/PF* online.



<sup>#</sup> Pharmaceutical Analytical Impurities (PAI products) are released using a process developed by USP's subject matter experts. The release process is based on internal policies, standard operating procedures, and requirements as defined by USP's Quality Management System. USP is an ISO 9001:2015 certified facility. PAI products are different from official USP Reference Standards. PAI products are not required for compendial compliance.

**Disclaimer:** USP Quality Solution Sheets are provided as a convenience and for informational purposes only. Labmix24 makes reasonable efforts to provide correct information but assumes no liability for the completeness, timeliness or accuracy of the information contained in each QSS. It is the responsibility of the user to verify the information. The US Pharmacopeia remains the official source of this information.

## www.labmix24.com