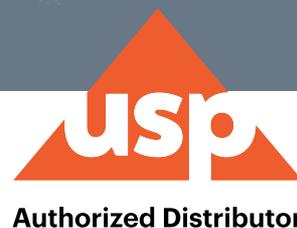
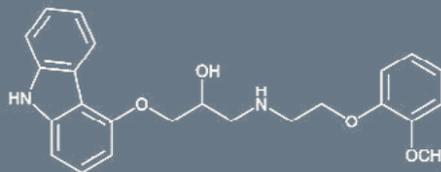


# Carvedilol

## Quality Solutions

Category: Cardiovascular



USP can support your development and manufacturing activities on Carvedilol-based medicines with these existing and upcoming standards.

## Official and Proposed Documentary Standards and Associated Physical Materials

### MONOGRAPH 1

<b>CARVEDILOL</b> Official as of 01-Jan-2021	
USP Carvedilol RS	<a href="#">USP-1096622</a>
USP Carvedilol Related Compound A RS	<a href="#">USP-1096633</a>
USP Carvedilol Related Compound B RS	<a href="#">USP-1096644</a>
USP Carvedilol Related Compound C RS	<a href="#">USP-1096655</a>
USP Carvedilol Related Compound D RS	<a href="#">USP-1096666</a>
USP Carvedilol Related Compound E RS	<a href="#">USP-1096677</a>
USP Carvedilol System Suitability Mixture RS	<a href="#">USP-1096688</a>

### MONOGRAPH 2

<b>CARVEDILOL TABLETS   NEW</b> Official as of 01-Aug-2023	
USP Carvedilol RS	<a href="#">USP-1096622</a>
<b>PHARMACEUTICAL ANALYTICAL IMPURITY(IES)*</b>	
USP N-Isopropylcarvedilol PAI I <b>NEW</b>	<a href="#">USP-1A00230</a>
USP Carvedilol Related Compound H PAI I <b>NEW</b>	<a href="#">USP-1A00890</a>
USP N-Nitroso Carvedilol Solution (1 mL (1mg/mL)) PAI I <b>NEW</b>	<a href="#">USP-1A04210</a>

\* *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.*

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