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Material No. : 1000364 Name Of Material : CARVEDILOL EP

Manufactured By : SYMED LABS LIMITED UNIT-II Inspection Lot No. : 010003260514

Supplied By : SYMED LABS LTD. Specification No. : 10012201

: 2CAN0030319 Batch No.

Version No. : 5
Date Of Receipt. : 01.04.2019
Manufacturing Date : MAR-2019 Advice Sheet No : 010003260514 Quantity : 50.000 KG Sample Quantity : 0.04 KG Retest Date : APR-2020

Analysis as per : RMT/320 Analytical Report No.: ARD11F0350 CEP No. : Not Applicable Date Of Report : 24.04.2019

Test	Limit	Results
Appearance	White or almost white, crystalline powder.	White, crystalline powder.
Solubility	Practically insoluble in water, sparingly soluble in methylene chloride, slightly soluble in ethanol (96 per cent). It is practically insoluble indilute acids.	Practically insoluble in water, sparingly soluble in methylene chloride, slightly soluble in ethanol (96 per cent). It is practically insoluble in dilute acids.
Identification: (A) By IR	The IR spectrum of the test sample should concordant with IR spectrum ofreference/working standard.	The IR spectrum of the test sample is concordant with IR spectrum of working standard.
Identification: (B) By P-XRD*	The XRD Diffractogram pattern of Carvedilol sample shouldhave characteristic peaks of Carvedilol form-II comprising2Ø angle values at 5.9°, 14.9°, 17.6°, 18.5° and 24.4° ±0.2° measured inX-ray Diffractometer.	The XRD Diffractogram pattern of Carvedilol sample is have characteristic peaks of Carvedilol form-II comprising 20 angle values at 5.7848°, 14.7767°, 17.4713°, 18.3848° and 24.2723° measured in X-ray Diffractometer.

Reviewed By	Approved By
Name:Sanjay Pandey	Name:Gopal Vora
Date:22.04.2019	Date:24.04.2019

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Material No. : 1000364

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Name Of Material : CARVEDILOL EP

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CEP No. : Not Applicable Date Of Report : 24.04.2019

Test	Limit	Results
Related substances (By HPLC)#:(A) Impurity A	Not more than 0.20 %	Below disregard limit
Related substances (By HPLC):(B) Impurity D	Not more than 0.15 %	Below disregard limit
Related substances (By HPLC): (C) Impurity C	Not more than 0.02 %	Not detected
Related substances (By HPLC):(D) Unspecified impurities	Not more than 0.10 %	Below disregard limit
Related substances (By HPLC)#:(E) Sum of impurities other than C	Not more than 0.50 %	Below disregard limit
Heavy Metals	Not more than 10 ppm.	Less than 10 ppm
Loss on drying#	Not more than 0.50 % w/w	0.24 % W/W
Sulfated Ash	Not more than 0.1 % w/w	0.06 % W/W
Assay	Not less than 99.0 % and not more than 101.0 % (on dried basis).	100.0 %
Residual solvents (By GC): (A) Methanol *	Not more than 1000 ppm	3 ppm

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Date:22.04.2019	Date:24.04.2019

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: 2CAN0030319 Version No. : 5 Batch No.

Date Of Receipt. Advice Sheet No : 010003260514 : 01.04.2019 Manufacturing Date : MAR-2019 Quantity : 50.000 KG Sample Quantity : 0.04 KG Retest Date : APR-2020 Analysis as per : RMT/320 Analytical Report No.: ARD11F0350

: Not Applicable Date Of Report : 24.04.2019

Test Limit Results Residual solvents (By GC): (B) Not more than 3000 ppm 213 ppm Ethyl acetate * 89 % Particle size (By Malvern): Not less than 70 % (A) Below 50 μ * 99 % Particle size (By Malvern): Not less than 95 % (B) Below 100 µ *

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: 2CAN0030319 : 5 Batch No.

Version No. Date Of Receipt. : 01.04.2019 Advice Sheet No : 010003260514 Manufacturing Date : MAR-2019 Quantity : 50.000 KG Sample Quantity : 0.04 KG Retest Date : APR-2020 Analysis as per : RMT/320 Analytical Report No.: ARD11F0350

CEP No. : Not Applicable Date Of Report : 24.04.2019

Remarks: Complies as per 10012201 Specification.

Sampling Plan : Z001

* Additional In-house tests.# Stringent limit than Ph.Eur.

Reviewed By	Approved By
Name:Sanjay Pandey	Name:Gopal Vora
Date:22.04.2019	Date:24.04.2019

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