

Each vial contains: Remdesivir 100 mg. **Dosage:** As directed by the Physician.

See enclosed prescribing information for reconstitution instructions and complete information on dosage and administration.

Store below 30°C until required for use. Reconstitute with 19 mL of sterile water for injection and diluted into 0.9% saline prior to administration by intravenous infusion.

After reconstitution, the storage time before administration should not exceed 4 hours at room temperature (20°C to 25°C) or 24 hours at refrigerated temperature (2°C to 8°C).

Preservative Free

Note: Parenteral products should be inspected visually for particulate matter prior to administration.

KEEP OUT OF THE REACH AND SIGHT OF CHILDREN.

Reg.No. 1C 45/63 (NG)



Remdesivir

100

for Injection 100 mg/vial

DESREM^{TI}

Lyophilized Powder for Injection for IV Infusion

Discard Unused Portion Single-Dose Vial Sterile



DESREM is manufactured under a license from Gilead Sciences, Inc.



Manufactured by:

Gland Pharma Limited

Plot No. 42 to 52, Survey No. 166, 171, 172 & 177, TSIIC, Phase-III, IDA, Pashamylaram (V), Patancheru (M), Sangareddy District, Telangana State - 502307, India

Imported by:

Viatris Pharmaceuticals (Thailand) Limited Bangkok, Thailand

Mfg. Lic. No.: TS/SGY/2020-65910 TM - Trade Mark under registration

100 mg

Remdesivir for Injection 100 mg/vial

DESREM[™]

Lyophilized Powder for Injection for IV Infusion

Discard Unused Portion Single-Dose Vial Sterile

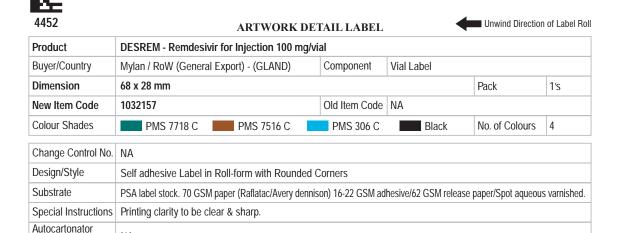


ARTWORK DETAIL LABEL							
PRODUCT	Remdesivir for Injection 100 mg/vial						
BUYER / COUNTRY	Mylan /ROW (GLAND - Dundigal)						
DIMENSION	48 x 48 x 88 mm	COMPONENT	Carton	PACK	1's	NO. OF COLOURS	4
COLOUR SHADES	PMS 7718 C PM	IS 7516 C	PMS 306 C		Black		
VERSION & DATE	Ver. 0; Date : 18.08.2021						
specia. Institutable limit Charedition Only. Codes shall be assigned during commercial artwork preparation.							





200%



Caution to the printer: Before processing, please ensure that the ARTWORK received for printing is exactly in line with APPROVED ARTWORK provided to you. In case of any FONTS/DESIGN are Mis-matching with the APPROVED ARTWORK, please inform PDC for further action. DO NOT MAKE ANY CHANGE TO THE ARTWORK WITHOUT WRITTEN INSTRUCTIONS FROM PDC.

1.3.1.1 labelling-Gland Rev. 0, Ver. 2; Date: 02-10-2020

Requirements