

Remdesivir

Reduce direct handling to a minimum and wear appropriate protective clothing		
CAUTION: High Administration Risk Rating		
CAUTION: Remdesivir is administered as a loading dose followed by a maintenance dose . Double check the correct dose has been prescribed		
Form	100mg powder for reconstitution	Store below 25°C
Reconstitution	Add 19mL WFI per 100mg vial (final concentration upon addition is 5mg/ml i.e. 100mg in 20ml) Allow the contents of the vial to settle for 2 to 3 minutes. The solution should be clear.	
Compatibility & Stability	Sodium chloride 0.9%	
Administration	IV Infusion (Preferred)	
	<ul style="list-style-type: none"> Withdraw and discard a volume of infusion fluid from the bag equal to the volume of drug solution being added Add the required volume of drug solution to the infusion bag Gently invert the bag 20 times to mix the solution in the bag Do not shake Give over 30 - 120 minutes 	
	Dose (mg) and number of Remdesivir 100mg vials	Infusion bag volume to be used (mL)
	200mg (2 vials)	250mL
	100mg (1 vial)	250mL
	Volume to be withdrawn and discarded from sodium chloride 0.9% bag 40mL 20mL	
Extravasation	<ul style="list-style-type: none"> Flush the line with at least 30mL of NaCl 0.9% to ensure that all the remdesivir solution has been administered Fluid restriction: Can use 100mL bag NaCl 0.9%. Withdraw and discard a volume of infusion fluid from the bag equal to the volume of drug solution being added 	
	Remdesivir has a low pH and may cause venous irritation and tissue damage in cases of extravasation. If a central venous access device is unavailable, administer via a large peripheral vein monitoring insertion site closely using a recognised phlebitis scoring tool. Re-site cannula at first signs of inflammation.	
Monitor	Monitor for signs of hypersensitivity and blood pressure. <ul style="list-style-type: none"> If infusion-related reactions occur, slow the infusion rate and give over 120 minutes. If significant hypersensitivity reactions occur, stop the infusion. 	
Additional Information	<ul style="list-style-type: none"> Any remaining reconstituted remdesivir for injection and / or diluted remdesivir solution for infusion should be disposed of in a purple lidded sharps bin. 	

Information provided relates to Veklury manufactured by Gilead
Last updated 28/11/25

This information has been summarised to act as a guide for those administering IV medication. The monograph should be used in conjunction with the drug data sheet and BNF for information on dose, adverse effects, cautions and contra-indications. Further information is available from Pharmacy on 22146 or 22542