

Nursing Education for Remdesivir

- Antiviral
- Authorized for emergency use in the United States and administered intravenously by health care providers, as appropriate, to treat suspected or laboratory-confirmed COVID-19 in adults and children hospitalized with severe disease. Severe disease is defined as patients with low blood oxygen levels or needing oxygen therapy or more intensive breathing support such as a mechanical ventilator.
- Ordering is restricted to Infectious Disease only
- Adult Administration:
 - Usual adult dose is 200mg on day 1 followed by 100mg daily for 4 days.
 - Administer as an IV infusion of 250mL over 120 minutes. Flush line with at least 30 mL NS after remdesivir infusion is complete
 - Pharmacy will mix medication with normal saline prior to administration.
- ****Important Information ****
 - Patient or designee must receive patient information prior to administration of the drug; Fact Sheet for Patients (Available in English and Spanish); To be printed directly from link in order.
 - Patient or designee must be aware that drug is not FDA approved and they may decline the use of the drug if they wish
 - Patient or designee must sign ***Informed Consent for Remdesivir.***
 - Specific consent can be printed directly from link in order. After consent has been signed, it will need to be scanned and uploaded into Epic.
 - Ordering provider (Infection Disease physician) will obtain and must sign the consent
 - *To eliminate touch points, verbal or telephone consent may be obtained and form may be signed outside of the patient room with 2 nurse's witness and signatures*
- Adverse Reactions:
 - Hepatic: Increased serum alanine aminotransferase (FDA 2020a), increased serum aspartate aminotransferase (FDA 2020a)
 - Miscellaneous: Infusion related reaction (including hypotension, nausea, vomiting, diaphoresis, and shivering) (FDA 2020a)
- Warnings/Precautions
 - Pharmacy will monitor labs daily prior to dispensing medication
 - If infusion reactions occur, discontinue administration and institute appropriate treatment if a clinically significant reaction and notify provider immediately
- Monitoring
 - CMP x1 at baseline and Daily x5
 - Signs and symptoms of infusion reaction
- Excretion: Urine-74% (majority as metabolites); Feces-18%
- IV Compatibilities: Unknown (Should only run with plain saline)

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References

- Johnston, K.S. (2020, December). Drug monograph: Remdesivir (Veklury-Gilead).
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