Nursing Education for Remdesivir

• Antiviral

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- 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 9 days for patients on mechanical ventilation, or 200mg on day 1 followed by 100mg daily for 4 days for patient not on a vent.
 - The prescriber may continue remdesivir for an additional 5 days (total of 10 days' treatment) for non-vented patients at their discretion.
 - 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 and prime tubing 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
- o 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

• 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

- o Pharmacy will monitor labs daily prior to dispensing medication
- o If infusion reactions occur, discontinue administration and institute appropriate treatment if a clinically significant reaction and notify provider immediately

Monitoring

- o CMP x1 at baseline and Daily x10
- o Signs and symptoms of infusion reaction
- Excretion: Urine-74% (majority as metabolites); Feces-18%

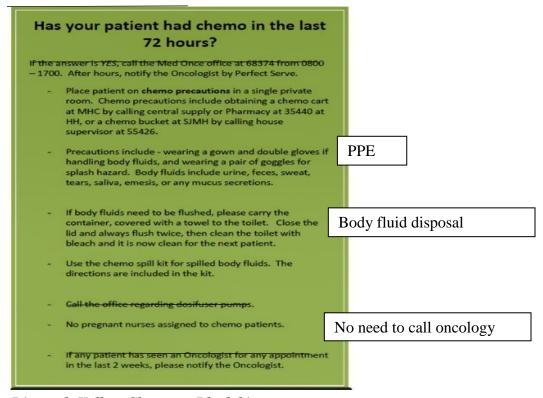
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• Special Considerations

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- o Remdesivir will be handled as a hazardous drug at SIH
- o Chemo precautions (Below are the instructions for ordered chemo precautions)



- o Disposal: Yellow Chemo or Black bins
- o IV Compatibilities: Unknown (Should only run with plain saline)

References

Gilead. (2020). Fact sheet for healthcare provider's emergency use authorization (EUA) of Remdesivir (GS-5734™). Retrieved from https://www.gilead.com/-/media/files/pdfs/remdesivir/eua-fact-sheet-for-

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