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|---------------------------------------|---------------------------------------|--|--|
| Referral Status: | | MRN: | |
| <input type="checkbox"/> New referral | <input type="checkbox"/> Order change | <input type="checkbox"/> Order Renewal | |
| Patient preferred clinic: | | | |

Tezspire™ (tezepelumab-ekko) Standard Plan of Treatment

PATIENT DEMOGRAPHICS:

| | | | |
|-------------------|------------------|-------------------|--|
| Date of Referral: | | Patient's Phone: | |
| Patient Name: | | Address: | |
| Date of Birth: | | City, State, Zip: | |
| Height in inches: | Weight: LB or KG | Gender: | Allergies: <input type="checkbox"/> See list <input type="checkbox"/> NKDA |

DIAGNOSIS: (PLEASE COMPLETE 2ND AND 3RD DIGITS TO COMPLETE ICD 10 FOR BILLING)

| | |
|--------------------------|---|
| <input type="checkbox"/> | J45.51 - Severe persistent asthma with (acute) exacerbation |
| <input type="checkbox"/> | J45.50 - Severe persistent asthma, uncomplicated |
| <input type="checkbox"/> | - Other: |

REQUESTED DOCUMENTATION:

| 1 | Insurance information | IF NO: | IF YES: |
|---|--|--|--|
| 2 | Most recent History & Physical | PLEASE STATE REQUIRED WASHOUT FROM PREVIOUS THERAPY: | LAST INJECTION DATE: |
| 3 | Full medication list | | NEXT INJECTION DATE: |
| 4 | Tried and failed therapies | | IF ORDER CHANGE: <input type="checkbox"/> Continue current order until insurance approved |
| 5 | Include any lab results/and or Pulmonary Function Tests to support diagnosis | | |

Provider Attestation for HCP administration:

| | | | |
|--------------------------|--|--------------------------|---|
| <input type="checkbox"/> | Provider attestation that the patient or caregiver are not competent or are physically unable to administer the Tezspire™ product FDA labeled for self-administration | <input type="checkbox"/> | Patient has experienced severe hypersensitivity reactions (e.g., anaphylaxis, angioedema, bronchospasm, or hypotension) to Tezspire™ within the past 6 months and requires administration and direct monitoring by a healthcare professional* |
| <input type="checkbox"/> | Patient has a history of uncontrolled disease and ordering provider attests that in their clinical opinion, it is not advisable to try the self-administered formulation of requested drug | <input type="checkbox"/> | Due to patient's weight, ordering provider attests that in their clinical opinion, it is not advisable to try the self-administered formulation of requested drug |
| <input type="checkbox"/> | The location and circumstances for self-administration are not adequate for the potential treatment of anaphylaxis should that arise. | | |

*Specific reactions: _____

MEDICATION ORDERS:

NOTE: Patient may be ineligible to receive TEZSPIRE™ (Tezepelumab-ekko) if patient has signs/symptoms of a parasitic infection, is currently being treated for a parasitic infection, or is having an acute bronchospasm and/or asthma attack.

DOSE/FREQUENCY:

Tezspire™ (Tezepelumab-ekko) 210mg every four (4) weeks via subcutaneous injection.
Administer subcutaneously to upper arm, thigh, or abdomen

OTHER FREQUENCY:

SPECIAL ORDERS:

Refills x 12 months unless noted otherwise here:

CARE ORDERS:

Provide nursing care per Palmetto Infusion Nursing Procedures and post procedure observation if indicated.

ADVERSE REACTION & ANAPHYLAXIS ORDERS:

Administer acute infusion and anaphylaxis medications per Palmetto Infusion standing adverse reaction orders, which can be found at our website or scan here.



PRESCRIBER INFORMATION:

| | |
|-------------------|--------|
| PROVIDER NAME: | PHONE: |
| ADDRESS: | FAX: |
| CITY, STATE, ZIP: | NPI: |

PRESCRIBER SIGNATURE: (No stamp signatures)

DATE

| | |
|---|------------------------|
| | |
| Dispense as written/Brand medically necessary | Substitution permitted |