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| Patient Name: Patient ID: | | | _ Date: _ Patient Date Of Birth: | 10/1 | 10/11/2024 | | | | |
|-------------------------------------|---|---|--|------|---|---|---|--|--|
| Pat | ient Group No: | NPI#: | Patient Phone: | Spe | Physician Name: Specialty: Physician Office Telephone | | | | |
| Phy | sician Office Address: | | | | | | • | | |
| Dru | g Name (specify drug) | | | | | | | | |
| Quantity: Route of Administration: | | • | _ Expected Length of Therapy: | _ | | | | | |
| | | | | | | | | | |
| | | | | | | | | | |
| | | | | | | | | | |
| Ple | | e answer for each applica | able question. | | | | | | |
| 1. | What is the diagnosis? | | | | _ | | | | |
| | Myelofibrosis, myelofi checked, go to 2) | brosis-associated anemia o | r myeloproliferative neoplasms (If | | Ш | | | | |
| | Other, please specify. | (If checked, no further que | stions) | | | | | | |
| 2. | Is the patient currently re | eceiving treatment with the | requested medication? | Y | | N | | | |
| 3. | Has there been an improtoxicity while on the curr | ovement in symptoms witho ent regimen? | ut any evidence of unacceptable | Y | | N | | | |
| 4. | What is the clinical settir | ng in which the requested m | nedication will be used? | | | | | | |
| | Accelerated phase or | blast phase myeloproliferat | ive neoplasm (If checked, go to 5) | | | | | | |
| | Low-risk myelofibrosis | s (If checked, go to 6) | | | | | | | |
| | Intermediate or high-ressential thrombocyth | isk primary or secondary (p emia) myelofibrosis (If chec | ost-polycythemia vera or post- cked, go to 6) | | | | | | |
| | High-risk myelofibrosi | s (If checked, go to 8) | | | | | | | |
| | Myelofibrosis-associa | ted anemia (If checked, go | to 9) | | | | | | |
| 5. | How will the requested r | medication be used? | | | | | | | |
| | As a single agent (If c | hecked, no further question | s) | | | | | | |
| | In combination with az | zacitidine (If checked, no fui | rther questions) | | | | | | |
| | In combination with de | ecitabine (If checked, no fur | ther questions) | | | | | | |
| | Other, please specify. | (If checked, no further que | stions) | | | | | | |
| 6. | What is the patient's pre or chart note(s) with pre | | CTION REQUIRED: Attach test resu | ılts | | | | | |
| | 50,000 or less (If ched | • | | | | | | | |
| | Greater than 50.000 (| If checked, no further quest | ions) | | П | | | | |

| | Unknown (If checked, no further questions) | | | | |
|-------|--|---|--|-----|--|
| | ACTION REQUIRED: Submit supporting documentation | | | | |
| 7. | Which of the following applies to the patient's disease? | | | | |
| | Symptomatic low-risk myelofibrosis (MF) (If checked, no further questions) | | | | |
| | Intermediate or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocythemia) myelofibrosis (MF) (If checked, no further questions) | | | | |
| | Other, please specify. (If checked, no further questions) | | | | |
| 8. | Does the patient have a symptomatic disease (e.g., splenomegaly and other disease-related symptoms)? | Y | | N 🔲 | |
| 9. | Does the patient have symptomatic splenomegaly and/or constitutional symptoms (e.g., fatigue, night sweats, fever, weight loss)? | Y | | N 🔲 | |
| and t | st that the medication requested is medically necessary for this patient. I further attest that the informat rue, and that the documentation supporting this information is available for review if requested by the classic | | | | |

plan sponsor, or, if applicable a state or federal regulatory agency.

Prescriber (Or Authorized) Signature and Date

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