

## Pharmacy Request for Prior Approval - Juxtapid **Recipient Information** 1. Recipient Last Name: First Name: 4. Recipient Date of Birth:\_\_\_\_\_\_5. Recipient Gender:\_ 3. Recipient ID # **Payer Information** 6. Is this a Medicaid or Health Choice Request? Medicaid: Health Choice: Prescriber Information 7. Prescribing Provider #: NPI: Or Atypical: 8. Prescriber DEA #: Requester Contact Information Name: Phone #: **Drug Information** 9. Drug Name: 10. Strength: \_\_\_ 11. Quantity per 30 Days: 12. Length of Therapy (in days): up to 30 60 90 120 180 365 other: **Clinical Information Request for Non-Preferred Drug:** 1. Has the member been diagnosed with homozygous familial hypercholesterolemia (HoFH)? Yes No 2. Is the member enrolled in the Juxtapid REMS program? 3. Is the member at least 18 years old or older? Yes No 4a. If female, has a negative pregnancy test been obtained? Yes No 5. Has a measurement of the recipient's ALT, AST, alkaline phosphatase, and total bilirubin been obtained before initiating treatment? 🔲 Yes 🔠 No 5a. ALT level:\_\_\_\_\_(U/L) 5b. AST level: \_\_\_\_\_\_(U/L) 5c. alkaline phosphatase level: \_\_\_\_\_ (U/L) 5d. Bilirubin level: (mg/dL) 6. For reauthorization: 6a. during the first year, has the member received liver-related tests (ALT and AST, at a minimum) prior to each increase in dose or monthly, whichever occurs first? Yes No 6b. after the first year, has the member received these tests at least every 3 months and before any increase in dose? $\square$ Yes $\square$ No 7. Failed two preferred drug(s). List preferred drugs failed: 7a. Allergic Reaction 7b. Drug-to-drug interaction. Please describe reaction: 8. Previous episode of an unacceptable side effect or therapeutic failure. Please provide clinical information: 9. Clinical contraindication, co-morbidity, or unique patient circumstance as a contraindication to preferred drug(s). Please provide clinical 10. Age specific indications. Please give patient age and explain: 11. Unique clinical indication supported by FDA approval or peer reviewed literature. Please explain and provide a general reference:

## (Prescriber Signature Mandatory)

12. Unacceptable clinical risk associated with therapeutic change. Please explain:

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

Fax this form to: (833) 404-2393

Signature of Prescriber:

Pharmacy PA Call Center: (833) 585-4309