



KYNAMRO® (mipomersen sodium) injection is only available through KYNAMRO Risk Evaluation and Mitigation Strategy (REMS).

In order to prescribe KYNAMRO, a prescriber must:

1. Complete the KYNAMRO REMS prescriber training by reviewing the materials in the KYNAMRO REMS Prescriber Education and Enrollment Kit.
2. Complete this one-time KYNAMRO REMS *Prescriber Enrollment Form*.
3. Complete and submit a KYNAMRO REMS *Prescription Authorization Form* for each new prescription.

**Complete this enrollment form and submit to KYNAMRO REMS by fax at 877-778-9008**

Prescriber Information (All information required)				
Name (first, middle, last)		Credentials <input type="checkbox"/> MD <input type="checkbox"/> DO <input type="checkbox"/> NP <input type="checkbox"/> PA <input type="checkbox"/> Other _____		
Name of Institution/Practice Name		Prescriber Specialty (Board Certification): <input type="checkbox"/> Cardiology <input type="checkbox"/> Endocrinology <input type="checkbox"/> Family Medicine <input type="checkbox"/> Internal Medicine <input type="checkbox"/> Other [please specify] _____		
Practice Setting: <input type="checkbox"/> Hospital-Based Practice <input type="checkbox"/> Private/Group Practice				
Practice Address				
City	State	Zip Code	Preferred Method of Contact <input type="checkbox"/> Fax <input type="checkbox"/> Phone	
Email Address	Office Phone Number		Office Contact Name	Office Fax Number
Primary State License Number/State of Issue			National Provider Identification (NPI) Number	

**Prescriber Attestation**

**By signing this form, I attest that:**

- I understand that KYNAMRO is indicated as an adjunct to lipid-lowering medications and diet to reduce low density lipoprotein-cholesterol (LDL-C), apolipoprotein B (apo B), total cholesterol (TC), and non-high density lipoprotein-cholesterol (non-HDL-C) in patients with homozygous familial hypercholesterolemia (HoFH).
- I understand that KYNAMRO is only available through KYNAMRO REMS and that I must comply with the program requirements in order to prescribe KYNAMRO.
- I have completed the KYNAMRO REMS Prescriber Training.
- I understand that there is a risk of hepatotoxicity associated with KYNAMRO.
- I understand that serum ALT, AST, alkaline phosphatase, and total bilirubin must be measured before initiating therapy with KYNAMRO.
- I understand that during the first year of treatment with KYNAMRO, liver-related laboratory tests (ALT and AST at a minimum) must be measured monthly.
- I understand that after the first year, these parameters should be measured at least every 3 months.
- I agree that personnel from the KYNAMRO REMS Program may contact me to gather further information or resolve discrepancies or to provide other information related to KYNAMRO or KYNAMRO REMS.
- I will complete and submit a KYNAMRO REMS *Prescription Authorization Form* for each new prescription.
- I agree that Kastle Therapeutics, its agents, and contractors such as the pharmacy providers may contact me via phone, mail, or email to survey me on the effectiveness of the program requirements for KYNAMRO REMS.

Prescriber Signature \_\_\_\_\_

Date \_\_\_\_\_

Print Name \_\_\_\_\_

**Questions? Contact KYNAMRO REMS**

Phone: 877-596-2676 | Fax: 877-778-9008 | [www.KynamroREMS.com](http://www.KynamroREMS.com)

The KYNAMRO Prescription Authorization Form is available at [www.KynamroREMS.com](http://www.KynamroREMS.com)

**Please see Prescribing Information for KYNAMRO.**

KYNAMRO is a registered trademark of Kastle Therapeutics.

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