

## Amvuttra Order Form (vutrisiran)

FAX TO: 972.499.9210

		PATIENT INFO	DRMATION		
Patient Name:	DO	3: Phor	e:	Sex: M F Ht: _	Wt: lbs kg
Primary Language:	Allergies: _				
Patient Preferred Locat	ion:				
<icd 10="" code="" requir<="" td=""><td>ED DIAGN</td><td>OSIS &amp; CLINIC</td><td>AL INFORMATI</td><td>ON</td><td></td></icd>	ED DIAGN	OSIS & CLINIC	AL INFORMATI	ON	
Primary ICD 10 Code	edofamilial Amyloidosis	Prescribing info		l racommonded cumple	mentation: Supplement
Loo. 1 Neuropaulic Her	edolamiliai Amyloidosis	with the recomn	nended daily allowar	nce of vitamin A. Refer	to an ophthalmologist if
				nin A deficiency occur UTTRA as soon as pos	sible. Resume dosing
				tly administered dose	·
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<u>REQUIRED</u> : Demographics & Most Recent: H&P, clinical notes, & medication list. Supporting clinical notes to include any past tried and/or failed therapies, intolerance, outcomes, or contraindications to conventional therapy.					
<u>LAB RESULTS</u> : Serum TTR, PND Scores, FAP Stage, or modified Neuropathy Impairment Scores and/or tests to support diagnosis.					
diagnosis.					
		PRESCRI	PTION		
AMVUTTRA (vutrisiran)					
Administer 25 mg by subcutaneous injection once every 3 months for one year					
Is the patient on any other disease modifying therapy? Yes No If yes, please note therapy and last dose:					
Post Treatment Observations: The patient is observed for 30 minutes following the first administration.					
<b>Adverse Events:</b> In the event of an adverse reaction occurring at a Medix Infusion suite, utilize the Medix Infusion adverse reactions protocol.					
Comments:					
					_
		_			
PRESCRIBER INFORMATION					
Prescriber Name:			_ Signature:		
Date:	NPI #:	Sr	ecialty:		
Supervising Physician:					(If Applicable)
Address:		City:		State:	Zip:
Contact Name:	Phone		Fax:	Email:	