## **AVISE® Test Requisition** Provider Relations: 888.452.1522





STEP	Patient & Provider I	nformation (I	Required	)
Patient Full Name:		Provider Name:		vider Details
City: State:	ZIP:	NPI #: Practice Name: Address:		
		Phone:		State: ZIP: Fax:
Attach a copy of front and back BILLING INFORMATION: Insurance	Patient 🗆 Lab	Lab Name:		ZIP:
MEDICARE only Hospital: Non-hospital particular Non-hospital particular Non-hospital particular STEP 2 Provide ICD-10 Diagnosis Code consistent with the patient's m	Diagnosis: ICD-10 (	-		patient's condition and
	//	/	/	
3         Date Specimen(s) Collected (required)         Date Specimen(s) Collected (required)         10 mL whole blood EDTA (lavender tube)         5 mL serum SST (tiger top tube)         AVISE Lupus (included with AVISE CTD)         ENA       Thyroid         U1RNP       TPO         0 U1RNP       TG         RNP70       TG         Ro60       RA         RF IgA       IgM         RF IgA       IgM         Add AVISE SLE Prognostic if AVISE Index is POSITIVE	red):       /       Time         D mL whole blood EDTA (lavender tube)       5 mL serum SST (tiger top tube)         AVISE Lupus consists of 10       analytes, including 2 CB-CAPs         (EC4d & BC4d) and 8       autoantibodies (ANA, anti-dsDNA, anti-Smith, anti-CCP, anti-Centromere protein B, anti-Jo-1, anti-Scl70, and anti-SSB/La), to aid the differential diagnosis of Lupus.         Add AVISE SLE Prognostic if AVISE Index is POSITIVE	of collection:	bignostic ube) G □ IgG JM □ IgM JA □ IgA Differ avender tube) ube)	<pre>(full name): AVISE Vasculitis-AAV S mL serum SST (tiger top tube) Anti-PR3 Anti-GBM Anti-MPO ANCA (IFA) ANTISE MTX S mL whole blood EDTA (lavender tube) Current dose: mg/week Injection Or Number of pills/week AVISE HCQ S mL whole blood EDTA (lavender tube) Current dose: mg/day Specimen should be collected at least 4 hours </pre>
AVISE Anti-CarP 5 mL serum SST (tiger top tube) Anti-Histone 5 mL serum SST (tiger top tube)	AVISE APS         5 mL serum SST (tiger top tube)         aCL       β2 GP1       PS/PT         IgG       IgG       IgG         IgG       IgG       IgG         IgM       IgM       IgM         IgA       IgA       IgA	Current dose: Specimen should be collecte after last dose	mg/day dd at least 4 hours mg/week	after last dose
In the event test orders contain overlapping analyt	es, those analytes will be reported on eac	h test report but will not be	performed more t	han once.
STEP	Medically N	ecessary		
Physician signature:	are) reasonable and medically necessa	Date:		

AVISE Specimen Requirements				
Order Type	Tube Requirements	Specimen Requirements		
AVISE Blood Tests	<b>One</b> - 10 mL whole blood EDTA (lavender tube) <b>One</b> - 5 mL serum SST (tiger top tube)	<ul> <li>EDTA should be drawn first</li> <li>Properly dispose of all contaminated materials in accordance with local disposal procedures</li> </ul>		
	AVISE Specimen	Submission		
PREPARE SPECIN	IEN COLLECTION KIT FOR SHIPPING:			
Ship specimens Mo	nday through Friday on same day blood is drawn, r	priority overnight delivery, using pre-printed shipping label.		
	I pack in one of the cooler wells.			
	n(s) in Bio-Hazard specimen bag and place bag inside multiple patients may be included in the same bo			
3. Replace foam coo transportation ki		d insurance card copies on top of cooler before closing outer		
4. Place kit inside	plastic carrier bag and affix shipping label to bag.			
5. Contact carrier i for assistance.	ndicated on the prepaid shipping label for pick-u	p or call Exagen Provider Relations at 888.452.1522		



## **QUESTIONS?**

Call 888.452.1522 or visit www.AviseTest.com or email shipping@exagen.com to place a kit order.

AVISE tests are used for clinical purposes, not to be regarded as investigational or for research. Results are not intended to be used as sole means for clinical diagnosis and patient management decisions. The following AVISE tests (AVISE CarP, AVISE CB-CAPs, AVISE CTD, AVISE Lupus, AVISE HCQ, AVISE MTX, AVISE SLE Monitor, AVISE SLE Prognostic) were developed, and performance characteristics were determined by Exagen Inc. as Laboratory Developed Tests (LDTs). The Exagen laboratory is certified under the Clinical Laboratory Amendments of 1988 (CLIA) and accredited by the College of American Pathologists (CAP) as qualified to perform high-complexity clinical laboratory testing, and FDA approval or clearance is not necessary.

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Exagen Inc. | 1261 Liberty Way | Vista, California 92081 Tel: 888.452.1522 | Fax: 888.452.8344 | www.AviseTest.com