FINAL Report Date: May 8, 2023 **Octave MS Disease Activity Test Report** PATIENT NAME DOB SEX INTERNAL PT ID CURRENT DMT

Octave MS Disease Activity Test

TEST REQUESTED

CLINIC FAX

Not Provided

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la.	no	Eacto	1

Jane Fostei

36.3

Disease Activity Score

TRFID

MCU1610

Sanctum Sanctorum

CLINIC NAME

0.00	-	Ee	otor	

ORDERING PHYSICIAN

Stephen Strange

AGE AT SAMPLE DRAWN

Jun 9, 1985 F ST-0000060 Kesimpta

YEAR OF DIAGNOSIS Not Provided

TAVF

COLLECTION DATE Oct 11, 2021

Mock Data - No PHI

CLINIC PHONE 212-555-1212

2.5 Low

Patient has a Low Disease Activity (DA) Score. Generally, this indicates disease activity is well controlled as evidenced by a high probability of minimal or no radiographic worsening. This Low DA score has decreased by 1.5 units from the previous DA score, which was in the Low category.



Disease Activity & Pathway Scores: Current and Historical Results

Collection Date	DA Score	Immunomodu	lation Score Neuroinflamma	ation Score Myelin Biol	ogy Score Neuroaxonal In	tegrity Score
10/11/2021	(2.5 L)	2.0	2.5	2.5	2.5	
9/01/2020	(4.0 L)	4.5	4.5	2.5	5.0	
8/29/2019	(3.0 L)	3.0	3.0	1.5	3.0	
9/10/2018	(8.0 H)	7.0	7.5	8.0	9.0	
9/21/2017	(5.5 M)	5.5	5.5	4.0	7.0	
9/27/2016	(8.5 H)	7.0	8.0	7.0	10.0	
DA Score Categories:	Low (L): 1.0 - 4.0	Moderate (M): 4.5 - 7.0	High (H): 7.5 - 10.0			

Test Description: The Octave MS Disease Activity Test measures the concentrations of 18 serum proteins. An algorithm is applied that utilizes subsets of the protein concentrations (adjusted for age and sex) to calculate four Disease Pathway Scores that reflect key hallmarks of multiple sclerosis pathophysiology: Immunomodulation, Neuroinflammation, Myelin Biology and Neuroaxonal Integrity. The individual biomarkers and the four Disease Pathway scores are used to determine the overall Disease Activity Score. The scale of each score is scaled from 1.0 to 10.0 with intervals of 0.5. Prior to 01May2023, MSDA scores were derived from an earlier iteration of the algorithm. The current version of the algorithm was validated for disease activity assessments and results from the two algorithm versions were demonstrated to be equivalent. Test results are intended to aid in the assessment of disease activity in patients with MS when used in conjunction with standard clinical and radiographic assessments. This test is not intended or validated to diagnose MS.

The Octave MS Disease Activity Test is intended for clinical use. Octave Bioscience Inc. developed the MS Disease Activity Test and determined its performance characteristics. It has been analytically and clinically validated and is offered as a Lab Developed Test. It has not been cleared or approved by the US Food and Drug Administration (FDA). The Octave Clinical Laboratory is certified under the Clinical Laboratory Improvement Act of 1988 (CLIA) as qualified to perform high complexity clinical testing and is a College of American Pathology (CAP) Accredited Laboratory.	LABORATORY DIRECTOR	^{clia} №	LABORATORY ID N°
	Russell Kerschmann, MD	05D2168340	CDF-00354252
Octave Bioscience 1440 OBrien Drive, Suite B, Menlo Park, CA 94025 w	/ww.octavebio.com Phone (65	0) 459-0942	1/2

FINAL Report Date: May 8, 2023 Octave MS Disease Activity Test Report



PATIENT NAME			DOB	SEX	INTERNAL PT ID	CURRENT DMT	YEAR OF DIAGNOSIS	
Jane Foster			Jun 9, 1985	F	ST-00000060	Kesimpta	Not Provided	
AGE AT SAMPLE DRAWN	TRFID	TEST REQUESTED			COLLECTION DATE			
36.3	MCU1610	Octave MS Disease	Activitv Test		Oct 11, 2021			

Biomarker Pathway Categories



Please Note: Individual biomarker results are expressed to the hundredths place and are required inputs into the algorithms used to calculate the Disease Activity Score and the four Pathway Scores. Clinical interpretation of individual biomarker levels and the four disease pathway scores, which have different weights in the algorithms, has not been established.

(*) These 95% reference ranges (expressed in two significant figures) were established from 1645 patient samples tested during method validation at the Octave Bioscience Clinical Laboratory.

(1) Subject's biomarker level relative to levels in MS patient samples from which the MS ranges were determined.

Individual Biomarker Results

Mock Data - No PHI

Biomarker	Pathways	Concentration	MS Range*	Percentile ⁽¹⁾
NfL Neurofilament light	NI	4.90 pg/mL	3.5 - 42 _{pg/mL}	10 th
GFAP Glial Fibrillary Acidic Protein	NI	54.39 pg/mL	24 - 220 pg/mL	25 th
SERPINA9 Serpin Family A Member 9	NI	32.06 pg/mL	12 - 160 _{pg/mL}	34 th
FLRT2 Leucine-rich repeat transmembrane protein	NI	90.38 pg/mL	63 - 180 _{pg/mL}	30 th
CNTN2 Contactin 2	NI	0.91 ng/mL	0.65 - 3.3 ng/mL	14 th
PRTG Protogenin	NI	119.22 pg/mL	71 - 180 _{pg/mL}	56 th
OPN Osteopontin	NI M	19.52 ng/mL	9.5 - 39 ng/mL	56 th
MOG Myelin Oligodendrocyte Glycoprotein	M	14.85 pg/mL	12 - 47 _{pg/mL}	7 th
CXCL9 Monokine Induced by Gamma Interferon	MN	50.23 pg/mL	17 - 250 _{pg/mL}	60 th
CXCL13 C-X-C Motif Chemokine Ligand 13	(M N	40.24 pg/mL	22 - 190 _{pg/mL}	30 th
CD6 Cluster of Differentiation 6	MN	74.14 pg/mL	46 - 250 _{pg/mL}	20 th
CCL20 MIP 3-alpha	N	4.43 pg/mL	2.1 - 52 pg/mL	26 th

Inversely Correlated with Disease Activity

APLP1 ⁽²⁾ Amyloid Beta Precursor Like Protein 1		8.19 ng/mL	5.5 - 22 ng/mL	19 th
OPG ⁽²⁾ Osteoprotegerin	NI	0.54 ng/mL	0.41 - 1.4 ng/mL	17 th
TNFRSF10A ⁽²⁾ TRAIL-R1	NI N	5.34 pg/mL	2.8 - 9.7 pg/mL	61 st
TNFSF13B ⁽²⁾ BAFF		5.00 ng/mL	2.3 - 10 ng/mL	66 th
IL-12B ⁽²⁾ Interleukin 12B		134.68 pg/mL	28 - 280 pg/mL	76 th
CDCP1 ⁽²⁾ CUB domain-containing protein 1	IM	91.86 pg/mL	28 - 230 _{pg/mL}	69 th

(2) Biomarker is significantly inversely correlated with disease activity, therefore a lower concentration was associated with a higher level of disease activity in validation studies.

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