ARUP LABORATORIES | www.aruplab.com
500 Chipeta Way, Salt Lake City, Utah 84108-1221
phone: (801) 583-2787, toll free: (800) 522-2787
Julio C. Delgado, MD, MS, Director of Laboratories

Client: ARUP Physician Services
321 TESTING ANSR EXTRACT
Salt Lake City, NY 84108
UNITED STATES

Physician: TEST,

Patient: 21596 TEST, 2C8/2C9 1

DOB
Gender: Female
Patient Identifiers: [redacted]
Visit Number (FIN): [redacted]
Collection Date: 5/21/2019 10:17

CYP2C8 and CYP2C9

ARUP test code 3001501

2C8/2C9 Specimen	Whole Blood
CYP2C8 Genotype	Neg/Neg
CYP2C9 Genotype	Neg/Neg
2C8/2C9 Interpretation	See Note

Interpretation: No impaired CYP2C8 variants were detected, which is consistent with functional *1 alleles. This result predicts the normal metabolizer phenotype.

Interpretation: No impaired CYP2C9 variants were detected, which is consistent with functional *1 alleles. This result predicts the normal metabolizer phenotype.

This result has been reviewed and approved by Pinar Bayrak-Toydemir, M.D., Ph.D.

H=High, L=Low, *=Abnormal, C=Critical

BACKGROUND INFORMATION: CYP2C8 and CYP2C9

CHARACTERISTICS: The cytochrome P450 (CYP) isozymes 2C8 and 2C9 are involved in the metabolism of many drugs. Variants in the genes that code for CYP2C8 and CYP2C9 may influence pharmacokinetics of substrates, and may predict or explain non-standard dose requirements, therapeutic failure or adverse reactions.
INHERITANCE: Autosomal co-dominant.
CAUSE: CYP2C8 and CYP2C9 gene variants affect enzyme expression or activity.
VARIANTS TESTED:
(Variants are numbered according to NM_000770 transcript for CYP2C8 and NM_000771 transcript for CYP2C9)
Negative: No variants detected is predictive of the *1 functional alleles (CYP2C8 or CYP2C9).

CYP2C8*1C: rs17110453, c.-370T>G
CYP2C8*2: rs11572103, c.805A>T
CYP2C8*3: rs10509681, c.1196A>G
CYP2C8*4: rs1058930, c.792C>G

CYP2C9*2: rs1799853, c.430C>T
CYP2C9*3: rs1057910, c.1075A>C
CYP2C9*4: rs56165452, c.1076T>C
CYP2C9*5: rs28371686, c.1080C>G
CYP2C9*6: rs9332131, c.817delA
CYP2C9*8: rs7900194, c.449G>A
CYP2C9*11: rs28371685, c.1003C>T

CLINICAL SENSITIVITY: Drug-dependent.
METHODOLOGY: Polymerase chain reaction (PCR) and fluorescence monitoring.
ANALYTICAL SENSITIVITY AND SPECIFICITY: Greater than 99 percent.
LIMITATIONS: Only the targeted CYP2C8 and CYP2C9 variants will be detected by this panel, and assumptions about phase and content are made to assign alleles. Publically available sources such as the www.pharmvar.org or www.pharmgkb.org provide guidance on phenotype predictions and allele frequencies. Diagnostic errors can occur due to rare sequence variations. Risk of therapeutic failure or adverse reactions with CYP2C8 or CYP2C9 substrates may be affected by genetic and non-genetic factors that are not detected by this test. This result does not replace the need for therapeutic drug or clinical monitoring.

Test developed and characteristics determined by ARUP Laboratories. See Compliance Statement C: aruplab.com/CS

VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
2C8/2C9 Specimen	19-141-104513	5/21/2019 10:17:00 AM	5/21/2019 11:26:00 AM	5/22/2019 8:59:00 AM
CYP2C8 Genotype	19-141-104513	5/21/2019 10:17:00 AM	5/21/2019 11:26:00 AM	5/22/2019 8:59:00 AM
CYP2C9 Genotype	19-141-104513	5/21/2019 10:17:00 AM	5/21/2019 11:26:00 AM	5/22/2019 8:59:00 AM
2C8/2C9 Interpretation	19-141-104513	5/21/2019 10:17:00 AM	5/21/2019 11:26:00 AM	5/22/2019 8:59:00 AM

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical