

Dr. MARIA JOSE PINZON GARCIA  
RED COLOMBIANA DE MEDICINA GENETICA SAS - PREGEN  
BOGOTA  
CARRERA 15 A # 106 - 42  
11001 BOGOTA  
Colombia

Date of Report 27.02.2026  
Sample Received 19.02.2026  
Date of Sampling 06.02.2026  
LAB-ID 262006434

## Medical Report

Patient name	<b>CALLE OROZCO IVANNA</b>	Sample-ID	A0321988
Date of Birth	<b>31.01.2025</b>	Gender	F

**Indication:** Newborn Screening

**Method(s):** Immunoassay, Tandem mass spectrometry from Dried Blood Spot

**Results:**

Parameter	Value	Unit	Reference
Birth weight (g)	3490	g	-
17-hydroxyprogesterone (17OHP)	8.5	nmol/L	< 90.0
Thyroid-stimulating hormone (TSH)	0.7	µU/mL	< 15.0
Biotinidase	-	U	> 51.0
Galactose-1-P-uridyltransferase (GALT)	-	U/g Hb	> 2.5
Immunoreactive trypsinogen (IRT)	16.8	ng/mL	< 65.0
Phenylalanine	23.4	µmol/L	< 150.0
Amino acid profile	negative		-
Acylcarnitine profile	negative		-

**Interpretation:** Not enough material. Please send a 2nd DBS card for re-testing of Biotinidase and GALT. All other tests were negative.

Please note: Inconspicuous biochemical results cannot exclude any metabolic disease with certainty. We recommend genetic testing if clinical symptoms suggest any inborn error of metabolism. For further information please contact us: [info@archimedlife.com](mailto:info@archimedlife.com).

Patient name	<b>CALLE OROZCO IVANNA</b>
Date of Birth	<b>31.01.2025</b>

Sample-ID	A0321988
Gender	F

## Results:

Parameter	Value	Unit	Reference
Phenylalanine (Phe)	23.4	µmol/L	< 150.0
Phenylalanine / Tyrosine ratio (Phe/Tyr)	0.41	µmol/L	< 2.20
Tyrosine (Tyr)	56.6	µmol/L	< 200.0
Leucine (Leu)	82.1	µmol/L	< 270.0
Valine (Val)	39.7	µmol/L	< 200.0
Methionine (MET)	29.0	µmol/L	< 78.0
Methionine / Phenylalanine (Met/Phe)	1.24	µmol/L	< 1.60
Citrulline (Cit)	6.8	µmol/L	< 50.0
Ornithine (Orn)	86.7	µmol/L	< 250.0
Ornithine / Citrulline ratio (Orn/Cit)	12.75	µmol/L	1.50 - 20.00
Proline (Pro)	90.4	µmol/L	< 350.0
Alanine (Ala)	131.2	µmol/L	< 750.0
Arginine (Arg)	2.8	µmol/L	< 100.0
Aspartic acid (Asp)	31.0	µmol/L	< 100.0
Glutamic acid (Glu)	296.6	µmol/L	< 600.0
Glycamine (Gly)	154.4	µmol/L	< 700.0
Free carnitine (C0)	11.03	µmol/L	6.00 - 100.00
acetylcarnitine (C2)	8.90	µmol/L	1.34 - 48.81
propionylcarnitine (C3)	0.71	µmol/L	0.13 - 6.60
butyryl-/isobutyrylcarnitine (C4)	0.13	µmol/L	0.03 - 0.90
isovaleryl-/2-methylbutyrylcarnitine(C5)	0.11	µmol/L	0.02 - 2.00
tiglylcarnitine (C5:1)	0.01	µmol/L	< 0.20
hydroxyvalerylcarnitine (C5OH)	0.14	µmol/L	0.02 - 0.57
glutarylacarnitine (C5DC)	0.02	µmol/L	< 0.30
hexanoylcarnitine (C6)	0.05	µmol/L	0.01 - 0.13
octanoylcarnitine (C8)	0.03	µmol/L	0.01 - 0.30
decanoylcarnitine (C10)	0.02	µmol/L	0.01 - 0.36
decenoylcarnitine (C10:1)	0.08	µmol/L	< 0.30
decadienoylcarnitine (C10:2)	0.02	µmol/L	< 0.10
dodecanoylcarnitine (C12)	0.07	µmol/L	0.10 - 0.60
myristoylcarnitine (C14)	0.14	µmol/L	0.01 - 0.57
tetradecenoylcarnitine (C14:1)	0.11	µmol/L	0.10 - 0.38
palmitoylcarnitine (C16)	1.41	µmol/L	0.62 - 7.81
3-hydroxypalmitoylcarnitine (C16OH)	0.02	µmol/L	< 0.10
stearoylcarnitine (C18)	0.59	µmol/L	0.30 - 2.40
oleylcarnitine (C18:1)	5.20	µmol/L	0.06 - 3.86
3-hydroxystearoylcarnitine (C18OH)	0.01	µmol/L	< 0.09
malonylcarnitine (C3DC)	0.02	µmol/L	< 0.50

Please note: Inconspicuous negative biochemical results cannot exclude any inborn error of metabolism or endocrine disorder with certainty in newborns. We recommend any follow-up or genetic testing if any clinical symptoms are present.

**Authorized By:** Assoc.-Prof. Dr. Andrea-Romana KASPER, MD, PhD  
[Specialist for Pediatrics, Neonatology and Nutrition]

Report was electronically signed and approved.