CODIGO : 172200

NOMBRE PACIENTE : MATEO NIÑO ARIAS **SEXO :** MASCULINO

FECHA DE NACIMIENTO : 21/12/2025 **REGISTRO CIVIL :** 1,113,072,721

NOMBRE RESPONSABLE : HINGRI LORENA ARIAS MORALES

DOC.IDENTIDAD DE LA MADRE : 1,114,727,423

FECHA TOMA DE MUESTRA : 03/02/2026 **TIPO DE MUESTRA :** TALÓN

FECHA DE IMPRESIÓN : 27/02/2026 **PESO :** 3535

TAMIZAJE NEONATAL

ANÁLISIS MUESTRA DE SANGRE

	RESULTADO	VALORES DE REFERENCIA	INTERPRETACIÓN
T.S.H Neonatal	1.20	>= 6 µl/mL talón en prematuros >= 10 µl/mL talón >= 15 µl/mL cordón	Normal
Deficiencia de G6PDH	6.10	> 2.6 U/gHb	Normal

TÉCNICA: Fluoroinmunoensayo (Delfia). *Procesado en Colombia por PREGEN.*

Hemoglobinopatías	FA	Ausencia de hemoglobinas anormales	Normal

TÉCNICA: Cromatografía Liquida de Alto Rendimiento (HPLC). *Procesado en Colombia por PREGEN.*

TAMIZAJE AMPLIADO

ESPECTROMETRIA DE MASAS EN TANDEM

Procesado en Archimedlife international medical laboratory. 1110 Vienna.

DESORDENES DE AMINOÁCIDOS

Citrulina, Metionina, Leucina, Isoleucina, Valina, Fenilalanina, Tirosina.

Ausencia de metabolitos anormales — Normal

PERFIL DE ACILCARNITINAS

C16,C18,C18:1,C16OH,C18:1OH,C8,C10:1,C5,C5DC,C4,C14,C14:1,C50H,C3,C5:1

Ausencia de metabolitos anormales — Normal

RESULTADOS NORMALES

Recuerde que estas son pruebas de tamizaje que solo indican la probabilidad de que el recién nacido tenga una de las enfermedades estudiadas por el programa y pueden requerir pruebas adicionales para la confirmación de algún diagnóstico. La sensibilidad de estas pruebas se reduce a medida que aumenta la edad del paciente, por esto es conveniente realizarlas dentro del primer mes de nacido.

REVISADO : EDUVILIA JOHANA GOMEZ
Bacterióloga
Reg. 40.936.003

PROCESADO : MARIA JOSE PINZON GARCIA
Bacterióloga
Reg. 1.015.469.392

FECHA : 27/02/2026

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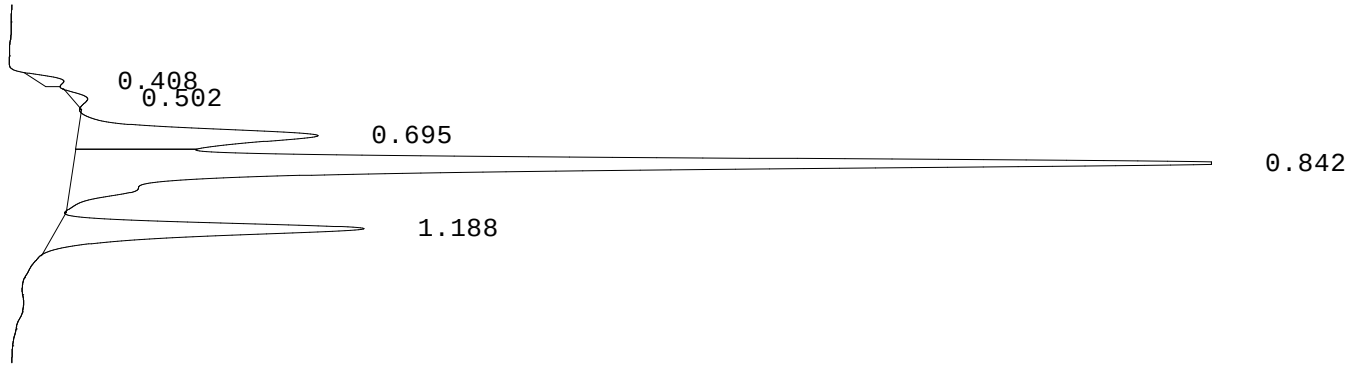
`LABORATORIO PREGEN
Carrera 15a No 106-42
BOGOTA

Batch 2068, Rack A, Plate 1, Well C05, 172200 FA
[CD26253B39840460] Feb 10, 2026 11:22:15 Pressure = 94 bar (94 to 97)



PEAK	RT	REL RT	% CONC	AREA	COMMENT
1	0.408	F 0.48	1.0%	7611	
2	0.502	F 0.59	0.7%	5468	
3	0.695	F 0.82	16.3%	123756	Acetylated F peak
4	0.842	F 1.00	65.2%	496019	Consistent with F 6
5	1.188	A 1.01	16.8%	127589	A peak
			Total Area:	760443	

Codes:
1) Wide A peak
2) Area of A peak < 80%
3) Peak area greater than expected
4) Peak after A2
5) A1c > 10%
6) HbF or variant present
7) Total sample area too small/big
8) A2 is not within normal range

baby health check©

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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	17.02.2026
Date of Sampling	03.02.2026
LAB-ID	262007354

Medical Report

Patient name	**NIÑO ARIAS MATEO**
Date of Birth	**21.12.2025**

Sample-ID	A0320565
Gender	M

Indication: Newborn Screening

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

Results:

Parameter	Value	Unit	Reference
Birth weight (g)	3535	g	-
17-hydroxyprogesterone (17OHP)	8.5	nmol/L	< 90.0
Thyroid-stimulating hormone (TSH)	1.5	µU/mL	< 15.0
Biotinidase	269.6	U	> 51.0
Galactose-1-P-uridyltransferase (GALT)	8.0	U/g Hb	> 2.5
Immunoreactive trypsinogen (IRT)	15.8	ng/mL	< 65.0
Phenylalanine	36.5	µmol/L	< 150.0
Amino acid profile	negative		-
Acylcarnitine profile	negative		-

Interpretation: NEGATIVE RESULT

Contact Details
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International Medical Laboratory+
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Patient name	**NIÑO ARIAS MATEO**	Sample-ID	A0320565
Date of Birth	**21.12.2025**	Gender	M

Results:

Amino Acids

Parameter	Value	Unit	Reference
Phenylalanine (Phe)	36.5	µmol/L	< 150.0
Phenylalanine / Tyrosine ratio (Phe/Tyr)	0.40	µmol/L	< 2.20
Tyrosine (Tyr)	92.0	µmol/L	< 200.0
Leucine (Leu)	157.3	µmol/L	< 270.0
Valine (Val)	71.2	µmol/L	< 200.0
Methionine (MET)	34.6	µmol/L	< 78.0
Methionine / Phenylalanine (Met/Phe)	0.95	µmol/L	< 1.60
Citrulline (Cit)	18.5	µmol/L	< 50.0
Ornithine (Orn)	123.8	µmol/L	< 250.0
Ornithine / Citrulline ratio (Orn/Cit)	6.69	µmol/L	1.50 - 20.00
Proline (Pro)	163.0	µmol/L	< 350.0
Alanine (Ala)	148.4	µmol/L	< 750.0
Arginine (Arg)	18.0	µmol/L	< 100.0
Aspartic acid (Asp)	103.2	µmol/L	< 100.0
Glutamic acid (Glu)	235.6	µmol/L	< 600.0
Glycamine (Gly)	187.8	µmol/L	< 700.0

Acylcarnitines

Free carnitine (C0)	13.82	µmol/L	6.00 - 100.00
acetylcarnitine (C2)	9.00	µmol/L	1.34 - 48.81
propionylcarnitine (C3)	1.53	µmol/L	0.13 - 6.60
butyryl-/isobutyrylcarnitine (C4)	0.10	µmol/L	0.03 - 0.90
isovaleryl-/2-methylbutyrylcarnitine(C5)	0.12	µmol/L	0.02 - 2.00
tiglylcarnitine (C5:1)	0.01	µmol/L	< 0.20
hydroxyvalerylcarnitine (C5OH)	0.25	µmol/L	0.02 - 0.57
glutarylcarnitine (C5DC)	0.03	µmol/L	< 0.30
hexanoylcarnitine (C6)	0.04	µmol/L	0.01 - 0.13
octanoylcarnitine (C8)	0.02	µmol/L	0.01 - 0.30
decanoylcarnitine (C10)	0.03	µmol/L	0.01 - 0.36
decenoylcarnitine (C10:1)	0.09	µmol/L	< 0.30
decadienoylcarnitine (C10:2)	0.02	µmol/L	< 0.10
dodecanoylcarnitine (C12)	0.05	µmol/L	0.10 - 0.60
myristoylcarnitine (C14)	0.06	µmol/L	0.01 - 0.57
tetradecenoylcarnitine (C14:1)	0.06	µmol/L	0.10 - 0.38
palmitoylcarnitine (C16)	0.53	µmol/L	0.62 - 7.81
3-hydroxypalmitoylcarnitine (C16OH)	0.02	µmol/L	< 0.10
stearoylcarnitine (C18)	0.19	µmol/L	0.30 - 2.40
oleylcarnitine (C18:1)	1.46	µmol/L	0.06 - 3.86
3-hydroxystearoylcarnitine (C18OH)	0.01	µmol/L	< 0.09
malonylcarnitine (C3DC)	0.02	µmol/L	< 0.50

Amino acid levels are indicators of phenylketonuria, tyrosinemia, MSUD, hydroxyprolinuria, hypermethioninemia (homocystinuria), citrullinemia, argininosuccinate aziduria, hyperargininemia, and hyperprolinemia. Acylcarnitine levels are indicators of carnitine uptake disorders, CPT-I deficiency, CPT-II deficiency, CAT deficiency, propionaciduria, methylmalonic aciduria, malonic aciduria, SCAD deficiency/ethylmalonic aciduria, isovaleric aciduria, HMG-CoA lyase deficiency, 3-methylcrotonyl-CoA carboxylase deficiency, methylglutaconiciduria, MCAD deficiency, VLCAD deficiency, LCHAD deficiency, glutaraziduria I, multiple acyl-CoA dehydrogenase deficiency (MAD deficiency/glutaraziduria II), and Beta-ketothiolase deficiency.

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.

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