



CODIGO : 172068
NOMBRE PACIENTE : JERONIMO FIERRO MEDINA **SEXO :** MASCULINO
FECHA DE NACIMIENTO : 21/11/2025 **REGISTRO CIVIL :** 1,206,231,681
NOMBRE RESPONSABLE : TATIANA MEDINA MOSQUERA
DOC.IDENTIDAD DE LA MADRE : 1,022,992,904
FECHA TOMA DE MUESTRA : 27/01/2026 **TIPO DE MUESTRA :** TALÓN
FECHA DE IMPRESIÓN : 18/02/2026 **PESO :** 2950

TAMIZAJE NEONATAL

ANÁLISIS MUESTRA DE SANGRE

	RESULTADO	VALORES DE REFERENCIA	INTERPRETACIÓN
T.S.H Neonatal	1.19	>= 6 µl/mL talón en prematuros >= 10 µl/mL talón >= 15 µl/mL cordón	Normal
Deficiencia de G6PDH	5.90	> 2.6 U/gHb	Normal
<i>TÉCNICA: Fluoroimmunoensayo (Delfia).</i>			<i>Procesado en Colombia por PREGEN.</i>
Hemoglobinopatías	FA	Ausencia de hemoglobinas anormales	Normal
<i>TÉCNICA: Cromatografía Líquida de Alto Rendimiento (HPLC).</i>			<i>Procesado en Colombia por PREGEN.</i>

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

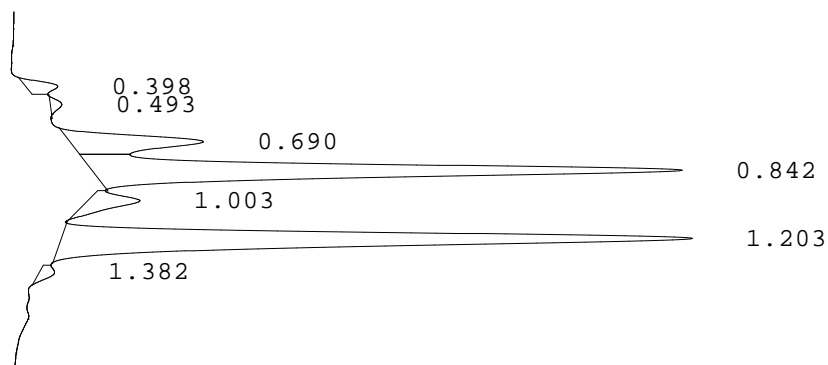
Recuerde que estas son pruebas de tamizaje que solo indican la probabilidad de que el recién nacido tenga una de las enfermedades estudiadas por el programa y pueden requerir pruebas adicionales para la confirmación de algún diagnóstico. La sensibilidad de estas pruebas se reduce a medida que aumenta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

REVISADO : EDUVILIA JOHANA GOMEZ **PROCESADO :** MARIA JOSE PINZON GARCIA **FECHA :**
Bacterióloga Bacterióloga
Reg. 40.936.003 Reg. 1.015.469.392 18/02/2026

LABORATORIO PREGEN
 Carrera 15a No 106-42
 BOGOTA

Batch 2061, Rack A, Plate 1, Well F12, 172068
 [C9678A05F6977317] Jan 30, 2026 12:29:06 Pressure = 88 bar (88 to 91)

FA



PEAK	RT	REL RT	% CONC	AREA	COMMENT
1	0.398	F 0.47	1.7%	10079	
2	0.493	F 0.58	0.7%	4015	
3	0.690	F 0.82	9.6%	57121	Acetylated F peak
4	0.842	F 1.00	39.0%	231377	Consistent with F
5	1.003	F 1.19	4.0%	23524	
6	1.203	A 1.01	44.0%	260797	A peak - REVIEW
7	1.382	S 0.89	1.0%	5964	3
Total Area:				592877	

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

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

Date of Report 17.02.2026
Sample Received 09.02.2026
Date of Sampling 27.01.2026
LAB-ID 262005989

Medical Report

Patient name	FIERRO MEDINA JERONIMO	Sample-ID	A0362076
Date of Birth	21.11.2025	Gender	M

Indication: Newborn Screening

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

Results:

Parameter	Value	Unit	Reference
Birth weight (g)	2950	g	-
17-hydroxyprogesterone (17OHP)	<5.0	nmol/L	< 90.0
Thyroid-stimulating hormone (TSH)	0.7	μU/mL	< 15.0
Biotinidase	207.7	U	> 51.0
Galactose-1-P-uridyltransferase (GALT)	8.1	U/g Hb	> 2.5
Immunoreactive trypsinogen (IRT)	20.2	ng/mL	< 65.0
Phenylalanine	49.6	μmol/L	< 150.0
Amino acid profile	negative		-
Acylcarnitine profile	negative		-

Interpretation: NEGATIVE RESULT

Patient name	FIERRO MEDINA JERONIMO
Date of Birth	21.11.2025

Sample-ID	A0362076
Gender	M

Results:

Parameter	Value	Unit	Reference
Phenylalanine (Phe)	49.6	µmol/L	< 150.0
Phenylalanine / Tyrosine ratio (Phe/Tyr)	0.64	µmol/L	< 2.20
Tyrosine (Tyr)	77.3	µmol/L	< 200.0
Leucine (Leu)	140.7	µmol/L	< 270.0
Valine (Val)	91.2	µmol/L	< 200.0
Methionine (MET)	26.8	µmol/L	< 78.0
Methionine / Phenylalanine (Met/Phe)	0.54	µmol/L	< 1.60
Citrulline (Cit)	18.0	µmol/L	< 50.0
Ornithine (Orn)	138.2	µmol/L	< 250.0
Ornithine / Citrulline ratio (Orn/Cit)	7.68	µmol/L	1.50 - 20.00
Proline (Pro)	102.7	µmol/L	< 350.0
Alanine (Ala)	152.0	µmol/L	< 750.0
Arginine (Arg)	27.2	µmol/L	< 100.0
Aspartic acid (Asp)	96.7	µmol/L	< 100.0
Glutamic acid (Glu)	207.0	µmol/L	< 600.0
Glycamine (Gly)	223.9	µmol/L	< 700.0
Free carnitine (C0)	25.41	µmol/L	6.00 - 100.00
acetylcarnitine (C2)	21.04	µmol/L	1.34 - 48.81
propionylcarnitine (C3)	2.77	µmol/L	0.13 - 6.60
butyryl-/isobutyrylcarnitine (C4)	0.21	µmol/L	0.03 - 0.90
isovaleryl-/2-methylbutyrylcarnitine(C5)	0.14	µmol/L	0.02 - 2.00
tiglylcarnitine (C5:1)	0.02	µmol/L	< 0.20
hydroxyvalerylcarnitine (C5OH)	0.18	µmol/L	0.02 - 0.57
glutaryl carnitine (C5DC)	0.03	µmol/L	< 0.30
hexanoylcarnitine (C6)	0.07	µmol/L	0.01 - 0.13
octanoylcarnitine (C8)	0.05	µmol/L	0.01 - 0.30
decanoylcarnitine (C10)	0.04	µmol/L	0.01 - 0.36
decenoylcarnitine (C10:1)	0.11	µmol/L	< 0.30
decadienoylcarnitine (C10:2)	0.02	µmol/L	< 0.10
dodecanoylcarnitine (C12)	0.03	µmol/L	0.10 - 0.60
myristoylcarnitine (C14)	0.07	µmol/L	0.01 - 0.57
tetradecenoylcarnitine (C14:1)	0.09	µmol/L	0.10 - 0.38
palmitoylcarnitine (C16)	0.53	µmol/L	0.62 - 7.81
3-hydroxypalmitoylcarnitine (C16OH)	0.01	µmol/L	< 0.10
stearoylcarnitine (C18)	0.38	µmol/L	0.30 - 2.40
oleylcarnitine (C18:1)	3.74	µmol/L	0.06 - 3.86
3-hydroxystearoylcarnitine (C18OH)	0.01	µmol/L	< 0.09
malonylcarnitine (C3DC)	0.03	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.

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