

CODIGO :
NOMBRE PACIENTE :
FECHA DE NACIMIENTO :
NOMBRE RESPONSABLE :
DOC.IDENTIDAD DE LA MADRE :
FECHA TOMA DE MUESTRA :
FECHA DE IMPRESIÓN :

173866
LIAM EMILIANO MARTINEZ VANEGAS
07/04/2026
LAURA FERNANDA MARTINEZ VANEGAS
1,030,533,731
26/06/2026
22/07/2026

SEXO : MASCULINO
REGISTRO CIVIL: 1,252,958,481
TIPO DE MUESTRA : TALÓN
PESO : 3330



TAMIZAJE NEONATAL

ANÁLISIS MUESTRA DE SANGRE

	RESULTADO	VALORES DE REFERENCIA	INTERPRETACIÓN
T.S.H Neonatal	1.74 µU/mL	>= 6 µU/mL talón en prematuros >= 10 µU/mL talón >= 15 µU/mL cordón	Normal
Deficiencia de G6PDH	6.70 U/gHb	< 2.6 U/gHb	Normal
TÉCNICA: Fluoroimmunoensayo (Delfia).		Procesado en Colombia por PREGEN.	
Hemoglobinopatías	AF	Ausencia de hemoglobinas anormales	Normal
TÉCNICA: Cromatografía Líquida 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, C5OH, 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

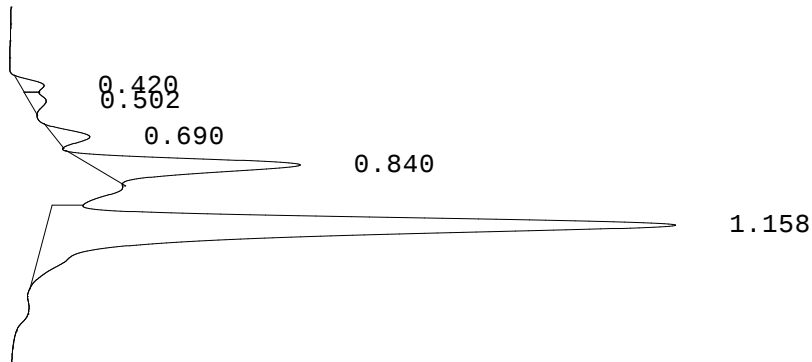
PROCESADO :
MARIA JOSE PINZON GARCIA
Bacterióloga
Reg. 1.015.469.392

FECHA :
22/07/2026

- - - - -
`LABORATORIO PREGEN
Carrera 15a No 106-42
BOGOTA

Batch 2188, Rack A, Plate 1, Well E01, 173866
[8C3CD4C32AF7B13D] Jul 01, 2026 12:31:33 Pressure = 79 bar (79 to 81)

AF



PEAK	RT	REL	RT	% CONC	AREA	COMMENT
1	0.420	F	0.50	1.6%	7679	
2	0.502	F	0.60	1.5%	7422	
3	0.690	F	0.82	2.7%	13369	
4	0.840	F	1.00	18.2%	88762	Consistent with F
5	1.158	A	1.00	76.0%	370301	A peak - REVIEW
Total Area: 487533						

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

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

Date of Report 11.07.2026
Sample Received 07.07.2026
Date of Sampling 26.06.2026
LAB-ID 262030689

Medical Report

Patient name	MARTINEZ VANEGAS LIAM EMILIANO	Sample-ID	A0358539
Date of Birth	07.04.2026	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)	3330	g	-
17-hydroxyprogesterone (17OHP)	<5.0	nmol/L	< 90.0
Thyroid-stimulating hormone (TSH)	1.1	μU/mL	< 15.0
Biotinidase	335.0	U	> 51.0
Galactose-1-P-uridyltransferase (GALT)	6.3	U/g Hb	> 2.5
Immunoreactive trypsinogen (IRT)	<15	ng/mL	< 65.0
Phenylalanine	32.9	μmol/L	< 150.0
Amino acid profile	negative		-
Acylcarnitine profile	negative		-

Interpretation: NEGATIVE RESULT

Contact Details

Assoc.-Prof. Dr. Andrea-Romana KASPER, MD, PhD
E-Mail: info@archimedlife.com

Patient name	MARTINEZ VANEGAS LIAM EMILIANO
Date of Birth	07.04.2026

Sample-ID	A0358539
Gender	M

Results:

Amino Acids

Parameter	Value	Unit	Reference
Phenylalanine (Phe)	32.9	µmol/L	< 150.0
Phenylalanine / Tyrosine ratio (Phe/Tyr)	0.38	µmol/L	< 2.20
Tyrosine (Tyr)	87.6	µmol/L	< 200.0
Leucine (Leu)	105.2	µmol/L	< 270.0
Valine (Val)	53.9	µmol/L	< 200.0
Methionine (MET)	21.6	µmol/L	< 78.0
Methionine / Phenylalanine (Met/Phe)	0.66	µmol/L	< 1.60
Citrulline (Cit)	41.1	µmol/L	< 50.0
Ornithine (Orn)	139.2	µmol/L	< 250.0
Ornithine / Citrulline ratio (Orn/Cit)	3.39	µmol/L	1.50 - 20.00
Proline (Pro)	91.3	µmol/L	< 350.0
Alanine (Ala)	197.9	µmol/L	< 750.0
Arginine (Arg)	60.9	µmol/L	< 100.0
Aspartic acid (Asp)	66.8	µmol/L	< 100.0
Glutamic acid (Glu)	189.1	µmol/L	< 600.0
Glycamine (Gly)	322.9	µmol/L	< 700.0

Acylcarnitines

Free carnitine (C0)	61.31	µmol/L	6.00 - 100.00
acetylcarnitine (C2)	26.08	µmol/L	1.34 - 48.81
propionylcarnitine (C3)	2.99	µmol/L	0.13 - 6.60
butyryl-/isobutyrylcarnitine (C4)	0.17	µmol/L	0.03 - 0.90
isovaleryl-/2-methylbutyrylcarnitine(C5)	0.10	µmol/L	0.02 - 2.00
tiglylcarnitine (C5:1)	0.01	µmol/L	< 0.20
hydroxyvalerylcarnitine (C5OH)	0.40	µmol/L	0.02 - 0.57
glutaryl carnitine (C5DC)	0.04	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)	0.04	µmol/L	0.01 - 0.36
decenoylcarnitine (C10:1)	0.11	µmol/L	< 0.30
dodecanoylcarnitine (C12)	0.08	µmol/L	0.10 - 0.60
myristoylcarnitine (C14)	0.22	µmol/L	0.01 - 0.57
tetradecenoylcarnitine (C14:1)	0.11	µmol/L	0.10 - 0.43
palmitoylcarnitine (C16)	2.22	µmol/L	0.62 - 7.81
3-hydroxypalmitoylcarnitine (C16OH)	0.03	µmol/L	< 0.10
stearoylcarnitine (C18)	1.26	µmol/L	0.30 - 2.40
oleylcarnitine (C18:1)	5.41	µmol/L	0.06 - 3.86
3-hydroxystearoylcarnitine (C18OH)	0.01	µmol/L	< 0.09
malonylcarnitine (C3DC)	0.08	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, methylglutaconiduria, 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.

Contact Details

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