CODIGO : 173447

NOMBRE PACIENTE : JAIDER SMITH NIÑO RODRIGUEZ **SEXO :** MASCULINO

FECHA DE NACIMIENTO : 23/03/2026 **REGISTRO CIVIL:** 1,070,998,533

NOMBRE RESPONSABLE : SHARIT YULIANA NIÑO RODRIGUEZ

DOC.IDENTIDAD DE LA MADRE : 1,104,070,007

FECHA TOMA DE MUESTRA : 16/05/2026 **TIPO DE MUESTRA :** TALÓN

FECHA DE IMPRESIÓN : 01/06/2026 **PESO :** 2830

TAMIZAJE NEONATAL

ANÁLISIS MUESTRA DE SANGRE

	RESULTADO	VALORES DE REFERENCIA	INTERPRETACIÓN
T.S.H Neonatal	0.98 µl/mL	>= 6 µl/mL talón en prematuros >= 10 µl/mL talón >= 15 µl/mL cordón	Normal
Deficiencia de G6PDH	5.70 U/gHb	< 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 : 01/06/2026

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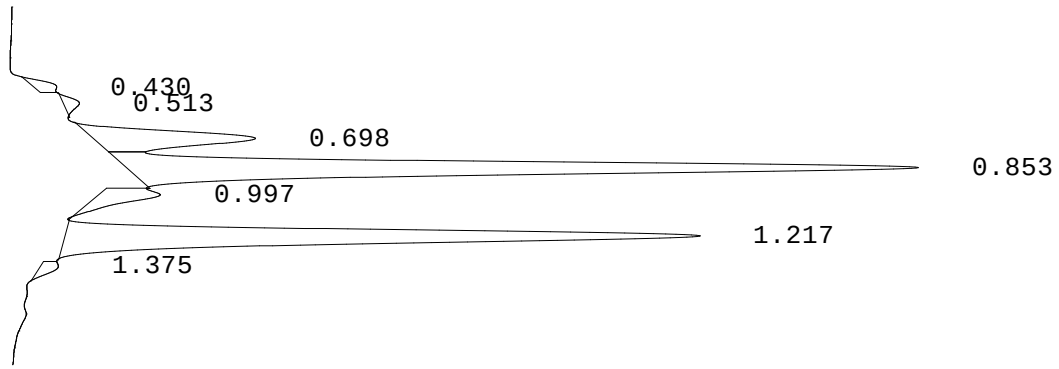
`LABORATORIO PREGEN
Carrera 15a No 106-42
BOGOTA

Batch 2149, Rack A, Plate 1, Well G12, 173447 FA
[9869A54949073315] May 20, 2026 14:58:27 Pressure = 64 bar (64 to 66)



PEAK	RT	REL RT	% CONC	AREA	COMMENT
1	0.430	F 0.50	1.1%	7777	
2	0.513	F 0.59	0.9%	6242	
3	0.698	F 0.81	10.8%	74780	Acetylated F peak
4	0.853	F 0.98	44.4%	309074	Consistent with F
5	0.997	F 1.15	4.1%	28407	
6	1.217	A 1.01	37.7%	262119	A peak - REVIEW
7	1.375	A 1.14	1.0%	6954	3
			Total Area:	695353	

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	29.05.2026
Sample Received	26.05.2026
Date of Sampling	16.05.2026
LAB-ID	262023700

Medical Report

Patient name	**NIÑO RODRIGUEZ JAIDER SMITH**
Date of Birth	**23.03.2026**

Sample-ID	A0341548
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)	2830	g	-
17-hydroxyprogesterone (17OHP)	<5.0	nmol/L	< 90.0
Thyroid-stimulating hormone (TSH)	<0.7	µU/mL	< 15.0
Biotinidase	183.5	U	> 51.0
Galactose-1-P-uridyltransferase (GALT)	6.5	U/g Hb	> 2.5
Immunoreactive trypsinogen (IRT)	<15	ng/mL	< 65.0
Phenylalanine	27.2	µmol/L	< 150.0
Amino acid profile	negative		-
Acylcarnitine profile	negative		-

Interpretation: NEGATIVE RESULT

Contact Details
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Patient name	**NIÑO RODRIGUEZ JAIDER SMITH**	Sample-ID	A0341548
Date of Birth	**23.03.2026**	Gender	M

Results:

Amino Acids

Parameter	Value	Unit	Reference
Phenylalanine (Phe)	27.2	µmol/L	< 150.0
Phenylalanine / Tyrosine ratio (Phe/Tyr)	0.42	µmol/L	< 2.20
Tyrosine (Tyr)	64.7	µmol/L	< 200.0
Leucine (Leu)	87.6	µmol/L	< 270.0
Valine (Val)	42.2	µmol/L	< 200.0
Methionine (MET)	15.1	µmol/L	< 78.0
Methionine / Phenylalanine (Met/Phe)	0.56	µmol/L	< 1.60
Citrulline (Cit)	7.5	µmol/L	< 50.0
Ornithine (Orn)	77.7	µmol/L	< 250.0
Ornithine / Citrulline ratio (Orn/Cit)	10.36	µmol/L	1.50 - 20.00
Proline (Pro)	93.9	µmol/L	< 350.0
Alanine (Ala)	114.5	µmol/L	< 750.0
Arginine (Arg)	13.1	µmol/L	< 100.0
Aspartic acid (Asp)	64.4	µmol/L	< 100.0
Glutamic acid (Glu)	209.5	µmol/L	< 600.0
Glycamine (Gly)	128.4	µmol/L	< 700.0

Acylcarnitines

Free carnitine (C0)	24.05	µmol/L	6.00 - 100.00
acetylcarnitine (C2)	12.63	µmol/L	1.34 - 48.81
propionylcarnitine (C3)	1.70	µmol/L	0.13 - 6.60
butyryl-/isobutyrylcarnitine (C4)	0.11	µmol/L	0.03 - 0.90
isovaleryl-/2-methylbutyrylcarnitine(C5)	0.08	µmol/L	0.02 - 2.00
tiglylcarnitine (C5:1)	0.01	µmol/L	< 0.20
hydroxyvalerylcarnitine (C5OH)	0.39	µmol/L	0.02 - 0.57
glutarylcarnitine (C5DC)	0.04	µmol/L	< 0.30
hexanoylcarnitine (C6)	0.03	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)	0.09	µmol/L	< 0.30
dodecanoylcarnitine (C12)	0.04	µmol/L	0.10 - 0.60
myristoylcarnitine (C14)	0.09	µmol/L	0.01 - 0.57
tetradecenoylcarnitine (C14:1)	0.07	µmol/L	0.10 - 0.38
palmitoylcarnitine (C16)	0.85	µmol/L	0.62 - 7.81
3-hydroxypalmitoylcarnitine (C16OH)	0.02	µmol/L	< 0.10
stearoylcarnitine (C18)	0.42	µmol/L	0.30 - 2.40
oleylcarnitine (C18:1)	2.22	µmol/L	0.06 - 3.86
3-hydroxystearoylcarnitine (C18OH)	0.01	µmol/L	< 0.09
malonylcarnitine (C3DC)	0.03	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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