

22/11/25



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FUNCIONAL

2025



O LGS Protocol: Curso Introductorio



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Introdução: Em que consiste o Protocolo LGS?

O **Protocolo LGS** é uma metodologia desenvolvida com a finalidade tratar raiz causa de:

**Doenças
Autoimunes**

**Síndrome
Metabólica**

**Câncer
(Cancro)**



**Modulação
do Microbioma**



**Modulação
do Sistema Imune**



**Compensação
Epigenética**

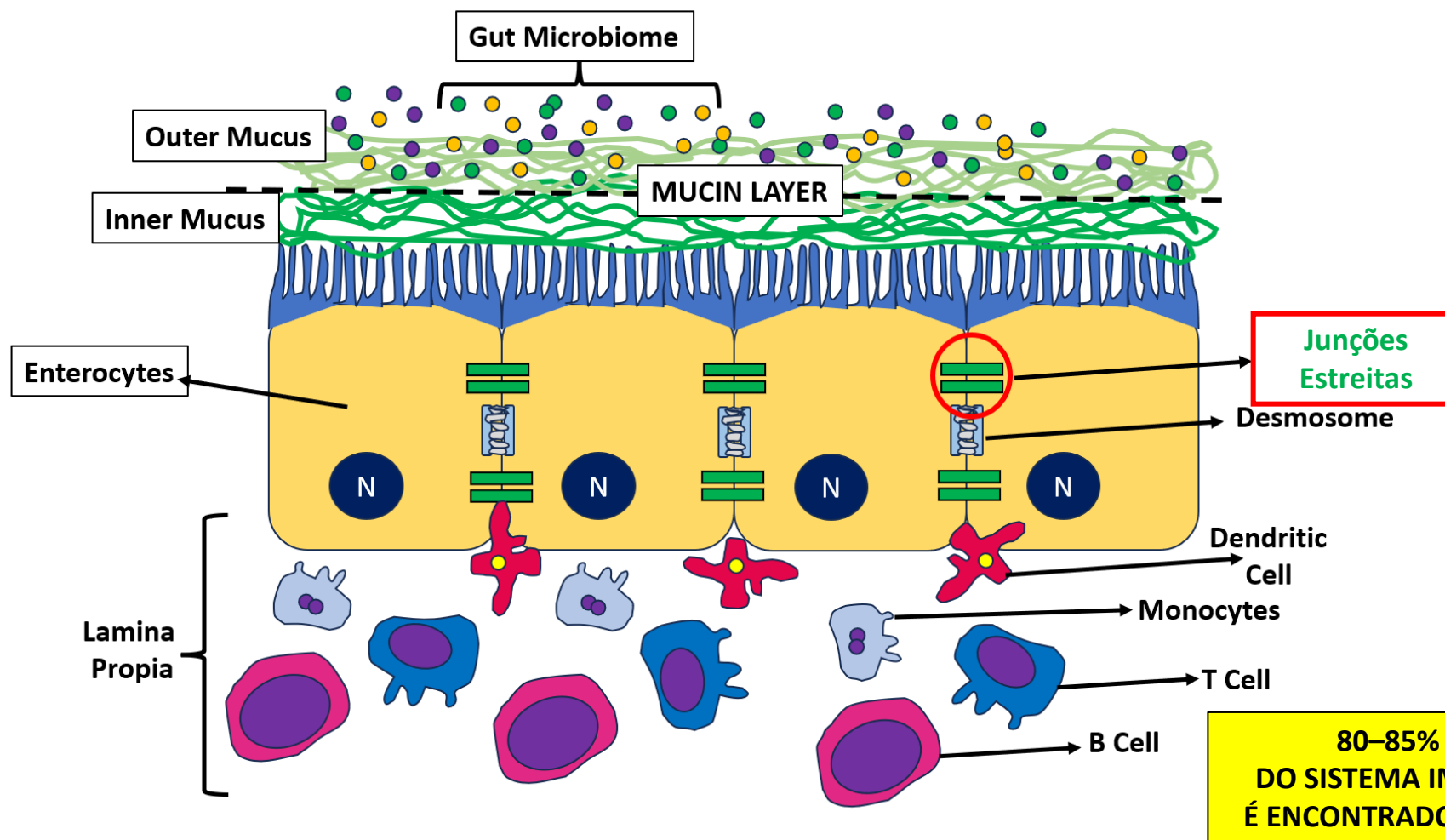
Ja temos profissionais de varios paises atuando com o Protocolo

 Dr. Fabiano Santos Guimarães Occupational Medicine - Brazilian Air Force Integrative Functional Medicine Clinical Researcher - Vitamin D Expert LGS Protocol Provider	 Dr. Oliver Igor Kuljis General Medicine Integrative Functional Medicine Clinical Researcher - Vitamin D Expert LGS Protocol Provider
 Dr. Cassiana Ferian Nutritionist Integrative Functional Nutritionist LGS Protocol Provider	 Dr. Ibrahim Arisoy Integrative Functional Medicine Vitamin D Expert LGS Protocol Provider
 Dr. Carlos Bayma Urologist Integrative Functional Medicine Vitamin D Expert LGS Protocol Provider	 Dr. Obada Yehia Daher Integrative Functional Medicine Vitamin D Expert LGS Protocol Provider
 Dr. Vivian Sarmiento Family Medicine Integrative Functional Medicine Vitamin D Expert LGS Protocol Provider	 Dr. Mounir Mohamed Fayed Barake Integrative Functional Medicine Vitamin D Expert LGS Protocol Provider

 Dr. Côtia Patricia Cruzio Miranda Nutritionist Integrative Functional Nutritionist LGS Protocol Provider	 Dr. Alin-Constantin Stancu Integrative Functional Medicine Vitamin D Expert LGS Protocol Provider
 Dr. Maria Sofia Venâncio de Oliveira Nutritionist Integrative Functional Nutritionist LGS Protocol Provider	 Dr. Selma Maria Melo Nutritionist Integrative Functional Nutritionist LGS Protocol Provider
 Dr. Rui Ribas Integrative Functional Nutritionist Vitamin D Expert LGS Protocol Provider	 Dra. Patricia Soares Fafá Guimarães Nutritionist Integrative Functional Nutritionist LGS Protocol Provider

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A Barreira GI/ Mucina:



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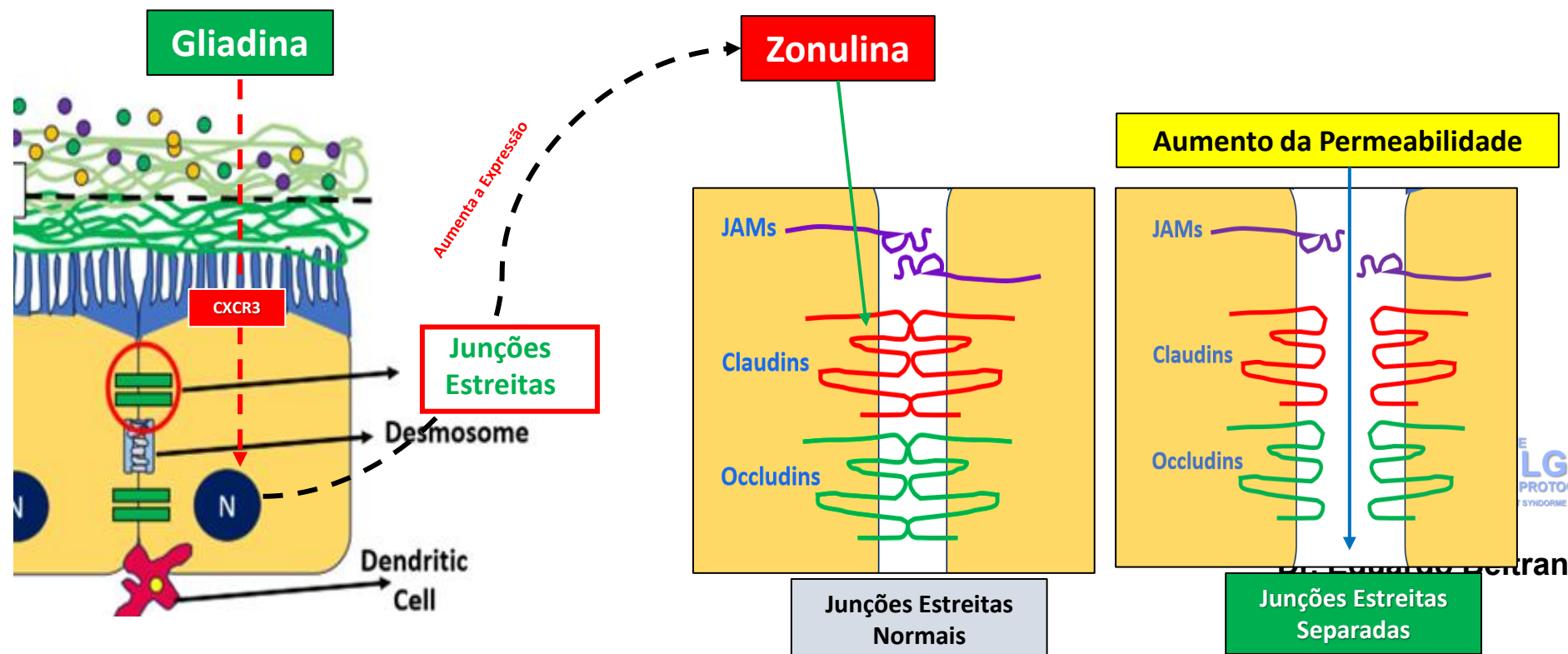
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Glifosato

Lectinas

TRIGO

LPS's



International Journal of
Molecular Sciences



Article

Lifelong Exposure to a Low-Dose of the Glyphosate-Based Herbicide RoundUp® Causes Intestinal Damage, Gut Dysbiosis, and Behavioral Changes in Mice

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Exames usados no Protocolo LGS:



5094 Shiloh Rd, Ste 101 | Alpharetta GA 30005
877.485.5336

Patient: Ima Sample
Collected: 08/02/2022
DOB: 07/11/1981
Gender: M

Accession: 00000000-0001
Received: 08/03/2022
Completed: 8/17/2022
Ordered by: Diane Farhi, MD

DNA STOOL ANALYSIS BY QUANTITATIVE PCR



YOUR PERSONALIZED REPORT

PATHOGENS

The GI-MAP® includes pathogens (bacterial, parasitic and viral) commonly known to cause gastroenteritis. Note that not all individuals with positive findings will present with symptoms. Many factors, including the health of the individual (such as immune health, digestive function, and microbiome balance), the transient nature of most pathogens, and the presence and expression of virulence factors, all contribute to pathogen virulence and individual symptoms.

BACTERIAL PATHOGENS	Result	Reference
<i>Campylobacter</i>	< dl	< 1.00e3
<i>C. difficile</i> Toxin A	1.21e5 High †	< 1.00e3
<i>C. difficile</i> Toxin B	2.27e5 High †	< 1.00e3
<i>Enterohemorrhagic E. coli</i>	< dl	< 1.00e3
<i>E. coli</i> O157	< dl	< 1.00e3
<i>Enteroinvasive E. coli</i> /Shigella	< dl	< 1.00e2
<i>Enterotoxigenic E. coli</i> LT/ST	< dl	< 1.00e3
Shiga-like Toxin <i>E. coli</i> stx1	< dl	< 1.00e3
Shiga-like Toxin <i>E. coli</i> stx2	< dl	< 1.00e3
<i>Salmonella</i>	< dl	< 1.00e4
<i>Vibrio cholerae</i>	< dl	< 1.00e5
<i>Yersinia enterocolitica</i>	4.46e3	< 1.00e5
PARASITIC PATHOGENS		
<i>Cryptosporidium</i>	< dl	< 1.00e6
<i>Entamoeba histolytica</i>	< dl	< 1.00e4
<i>Giardia</i>	< dl	< 5.00e3
VIRAL PATHOGENS		
Adenovirus 40/41	< dl	< 1.00e10
Norovirus GI/II	< dl	< 1.00e7

KEY: Results are reported as genome equivalents per gram of stool, which is a standard method for estimating the number of microbes measured per gram of stool, based on qPCR analysis of DNA samples.

Results are expressed in standard scientific notation. For example, a reported result of 3.5e7 is equivalent to 3.5 x 10⁷ microbes per gram, which equals 35,000,000 (35 million) microbes per gram of stool.

< dl represents results below detectable limit.

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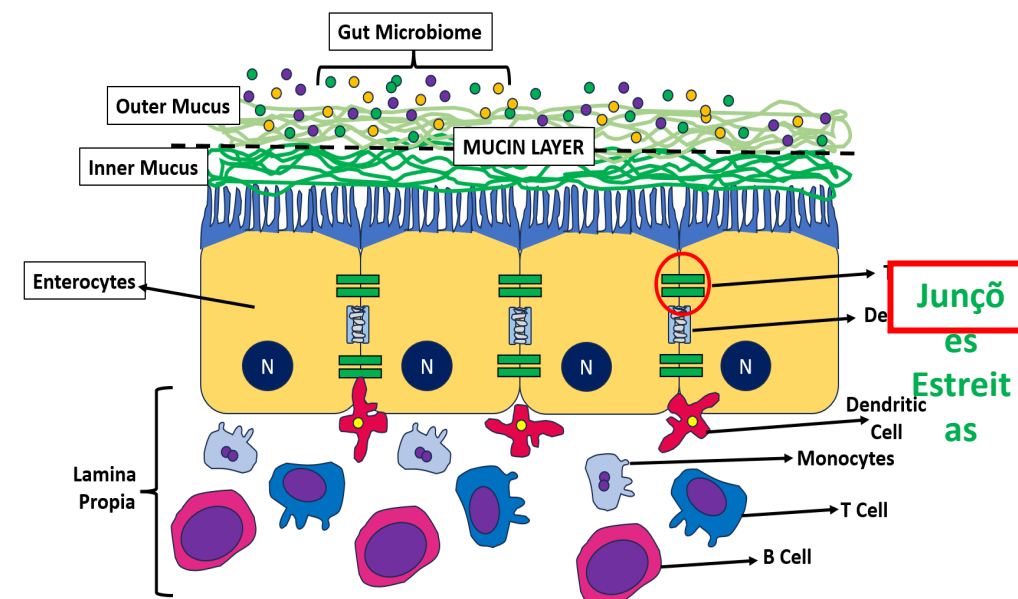
THE LEAKY GUT SYNDROME PROTOCOL

Dr. Eduardo Beltran

Exames usados no Protocolo LGS:



COMMENSAL/KEYSTONE BACTERIA			
COMMENSAL BACTERIA	Result		Reference
<i>Bacteroides fragilis</i>	1.09e11		1.6e9 - 2.5e11
<i>Bifidobacterium</i> spp.	5.49e9		> 6.7e7
<i>Enterococcus</i> spp.	1.09e7		1.9e5 - 2.0e8
<i>Escherichia</i> spp.	5.57e8		3.7e6 - 3.8e9
<i>Lactobacillus</i> spp.	5.69e7		8.6e5 - 6.2e8
<i>Enterobacter</i> spp.	1.33e7		1.0e6 - 5.0e7
<i>Akkermansia muciniphila</i>	<dl L		1.0e1 - 8.2e6
<i>Faecalibacterium prausnitzii</i>	<dl L		1.0e3 - 5.0e8
<i>Roseburia</i> spp.	2.08e9		5.0e7 - 2.0e10
BACTERIAL PHYLA			
<i>Bacteroidetes</i>	2.41e12		8.6e11 - 3.3e12
<i>Firmicutes</i>	1.52e11		5.7e10 - 3.0e11
<i>Firmicutes:Bacteroidetes</i> Ratio	0.06		< 1.0



80–85%
DO SISTEMA IMUNE
É ENCONTRADO AQUI

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GI MAP



Ana L., 51 anos, relata **perda progressiva de pigmentação da pele** nas mãos, ao redor da boca, **nos joelhos, membros inferiores e principalmente na região das costas** ao longo dos últimos dez meses. Ela observa que as manchas são bem delimitadas e aumentam lentamente. Também refere fadiga aumentada e unhas frágeis. Paciente consultou na primeira semana de Setembro, 2024.

Paralelamente às alterações cutâneas, apresenta **sintomas gastrointestinais persistentes**, incluindo distensão abdominal, desconforto, excesso de gases e diarreia intermitente várias vezes por semana. Esses sintomas pioram após ingerir cebola, alho, feijão, pão, maçã ou leite. Ela também relata episódios de névoa mental durante as crises.



GI MAP



ANTES DE INICIAR



Exames usados no Protocolo LGS:



5894 Shiloh Rd, Ste 101 | Alpharetta GA 30005
877.485.5336

Patient: [REDACTED]

Collected: 9/6/2024

DOB: 6/18/1974

Accession: [REDACTED]

Received: 9/10/2024

Completed: 10/1/2024

Ordered by: [REDACTED]

DNA STOOL ANALYSIS BY QUANTITATIVE PCR



YOUR PERSONALIZED REPORT

PATHOGENS

The testing includes pathogens (bacterial, parasitic and viral) commonly known to cause gastroenteritis. Note that not all individuals with positive findings will present with symptoms. Many factors, including the health of the individual (such as immune health, digestive function, and microbiome balance), the transient nature of most pathogens, and the presence and expression of virulence factors, all contribute to pathogen virulence and individual symptoms.

BACTERIAL PATHOGENS	Result	Reference
<i>Campylobacter</i>	<dl	< 1.00e3
<i>C. difficile</i> Toxin A	<dl	< 1.00e3
<i>C. difficile</i> Toxin B	<dl	< 1.00e3
Enterohemorrhagic <i>E. coli</i>	4.41e6 High ↑	< 1.00e3
<i>E. coli</i> O157	<dl	< 1.00e3
Enteroinvasive <i>E. coli/Shigella</i>	<dl	< 1.00e3
Enterotoxigenic <i>E. coli</i> LT/ST	<dl	< 1.00e3
Shiga-like Toxin <i>E. coli</i> stx1	<dl	< 1.00e3
Shiga-like Toxin <i>E. coli</i> stx2	<dl	< 1.00e3
<i>Salmonella</i>	<dl	< 1.00e4
<i>Vibrio cholerae</i>	<dl	< 1.00e5
<i>Yersinia enterocolitica</i>	<dl	< 1.00e5
PARASITIC PATHOGENS		
<i>Cryptosporidium</i>	<dl	< 1.00e6
<i>Entamoeba histolytica</i>	<dl	< 1.00e4
<i>Giardia</i>	<dl	< 5.00e3
VIRAL PATHOGENS		
Adenovirus 40/41	<dl	< 1.00e10
Norovirus GI/II	<dl	< 1.00e7

HELICOBACTER PYLORI

H. PYLORI & VIRULENCE FACTORS

	Result	Reference
<i>Helicobacter pylori</i>	4.38e3 High ↑	< 1.00e3
Virulence Factor, babA	Negative	Negative
Virulence Factor, cagA	Negative	Negative
Virulence Factor, dupA	Negative	Negative
Virulence Factor, iceA	Negative	Negative
Virulence Factor, oipA	Negative	Negative
Virulence Factor, vacA	Negative	Negative
Virulence Factor, virB	Negative	Negative
Virulence Factor, virD	Negative	Negative

H. PYLORI ANTIBIOTIC RESISTANCE GENES

	Result	Reference
Amoxicillin	Negative	Negative
Genes associated with amoxicillin resistance		
PBP1A S414R	Absent	
PBP1A T556S	Absent	
PBP1A N562Y	Absent	
Clarithromycin	Negative	Negative
Genes associated with clarithromycin resistance		
A2142C	Absent	
A2142G	Absent	
A2143G	Absent	
	Result	Reference
Fluoroquinolones	Negative	Negative
Genes associated with fluoroquinolone resistance		
gyrA N87K	Absent	
gyrA D91N	Absent	
gyrA D91G	Absent	
gyrB S479N	Absent	
gyrB R484K	Absent	
Tetracycline	Negative	Negative
Genes associated with tetracycline resistance		
A926G	Absent	
AGA926-928TTC	Absent	

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OPPORTUNISTIC/OVERGROWTH MICROBES

DYSBIOTIC & OVERGROWTH BACTERIA	Result	Reference
<i>Bacillus</i> spp.	4.48e5	< 1.76e6
<i>Enterococcus faecalis</i>	<dl	< 1.00e4
<i>Enterococcus faecium</i>	<dl	< 1.00e4
<i>Morganella</i> spp.	<dl	< 1.00e3
<i>Pseudomonas</i> spp.	1.79e2	< 1.00e4
<i>Pseudomonas aeruginosa</i>	<dl	< 5.00e2
<i>Staphylococcus</i> spp.	<dl	< 1.00e4
<i>Staphylococcus aureus</i>	3.21e3 High ↑	< 5.00e2
<i>Streptococcus</i> spp.	8.57e3 High ↑	< 1.00e3
COMMENSAL OVERGROWTH MICROBES		
<i>Desulfovibrio</i> spp.	1.73e8	< 7.98e8
<i>Methanobacteriaceae</i> (family)	1.10e8	< 3.38e8
INFLAMMATORY & AUTOIMMUNE-RELATED BACTERIA		
<i>Citrobacter</i> spp.	<dl	< 5.00e6
<i>Citrobacter freundii</i>	<dl	< 5.00e5
<i>Klebsiella</i> spp.	<dl	< 5.00e3
<i>Klebsiella pneumoniae</i>	2.43e2	< 5.00e4
<i>M. avium</i> subsp. <i>paratuberculosis</i>	<dl	< 5.00e3
<i>Proteus</i> spp.	<dl	< 5.00e4
<i>Proteus mirabilis</i>	<dl	< 1.00e3
COMMENSAL INFLAMMATORY & AUTOIMMUNE-RELATED BACTERIA		
<i>Enterobacter</i> spp.	1.33e7	< 5.00e7
<i>Escherichia</i> spp.	5.57e8	< 3.80e9
<i>Fusobacterium</i> spp.	1.13e8 High ↑	< 1.00e8
<i>Prevotella</i> spp.	2.49e7	< 1.00e8

FUNGI/YEAST

FUNGI/YEAST	Result	Reference
<i>Candida</i> spp.	<dl	< 5.00e3
<i>Candida albicans</i>	<dl	< 5.00e2
<i>Geotrichum</i> spp.	<dl	< 3.00e2
<i>Microsporidium</i> spp.	<dl	< 5.00e3
<i>Rhodotorula</i> spp.	<dl	< 1.00e3

VIRUSES

VIRUSES	Result	Reference
Cytomegalovirus	<dl	< 1.00e5
Epstein-Barr Virus	<dl	< 1.00e7

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GI Map



PARASITES

PROTOZOA	Result	Reference
<i>Blastocystis hominis</i>	<dl	< 2.00e3
<i>Chilomastix mesnili</i>	<dl	< 1.00e5
<i>Cyclospora</i> spp.	<dl	< 5.00e4
<i>Dientamoeba fragilis</i>	<dl	< 1.00e5
<i>Endolimax nana</i>	<dl	< 1.00e4
<i>Entamoeba coli</i>	<dl	< 5.00e6
<i>Pentatrichomonas hominis</i>	<dl	< 1.00e2
WORMS		
<i>Ancylostoma duodenale</i>	Not Detected	Not Detected
<i>Ascaris lumbricoides</i>	Not Detected	Not Detected
<i>Necator americanus</i>	Not Detected	Not Detected
<i>Trichuris trichiura</i>	Not Detected	Not Detected
<i>Taenia</i> spp.	Not Detected	Not Detected

INTESTINAL HEALTH MARKERS

DIGESTION	Result	Reference
Steatocrit	<dl	< 15 %
Elastase-1	485	> 200 ug/g
GI MARKERS		
β-Glucuronidase	1029	< 2486 U/mL
Occult Blood - FIT	47 H	< 10 ug/g
IMMUNE RESPONSE		
Secretory IgA	>6000 H	510 - 2010 ug/g
Anti-gliadin IgA	295 H	< 175 U/L
Eosinophil Activation Protein (EDN, EPX)	0.19	< 2.34 ug/g
INFLAMMATION		
Calprotectin	1748 H	< 173 ug/g
ADD-ON TESTS		
Gluten Peptide	<dl	< 5.0 ng/g
Zonulin	351.6 H	< 175 ng/g

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GI Map



UNIVERSAL ANTIBIOTIC RESISTANCE GENES

	Result	Reference		Result	Reference		Result	Reference
b-Lactams	Positive	Negative	Macrolides	Positive	Negative	Trimethoprim	Positive	Negative
blaNDM-1	Absent		acrA	Absent		dfrA1	Absent	
CTX-M 1	Absent		acrB	Present		dfrA12	Absent	
CTX-M 2	Absent		emrE	Present		dfrA14	Present	
CTX-M 8/25	Absent		ermA	Absent		dfrA15	Absent	
CTX-M 9	Absent		ermB	Present		dfrA17	Absent	
GES	Absent		ermC	Absent		dfrA5	Absent	
OXA-1	Absent		macA	Absent		dfrA7	Absent	
PER-1	Absent		macB	Absent		dfrB1	Absent	
PER-2	Absent		mefA	Present		dfrB2	Absent	
SHV	Absent		mphA	Absent		dfrB3	Absent	
TEM	Present		msrA	Absent				
VEB	Absent		tolC	Present				
Fluoroquinolones	Positive	Negative	Ciprofloxacin	Negative	Negative	Sulfonamides	Positive	Negative
qnrA	Absent		emeA	Absent		sul1	Absent	
qnrB	Present		pmrA	Absent		sul2	Present	
qnrS1	Present					sul3	Absent	
qnrS2	Present							
Vancomycin	Negative	Negative	Nitroimidazoles	Positive	Negative	Methacillin	Negative	Negative
vanA	Absent		nimA	Absent		mecA	Absent	
vanB	Absent		nimB	Absent		Chloramphenicol	Positive	Negative
vanC1	Absent		nimC	Absent				
vanC2-1	Absent		nimD	Absent		catA13	Present	
vanC2-2	Absent		nimE	Present				

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Exames usados no Protocolo LGS:



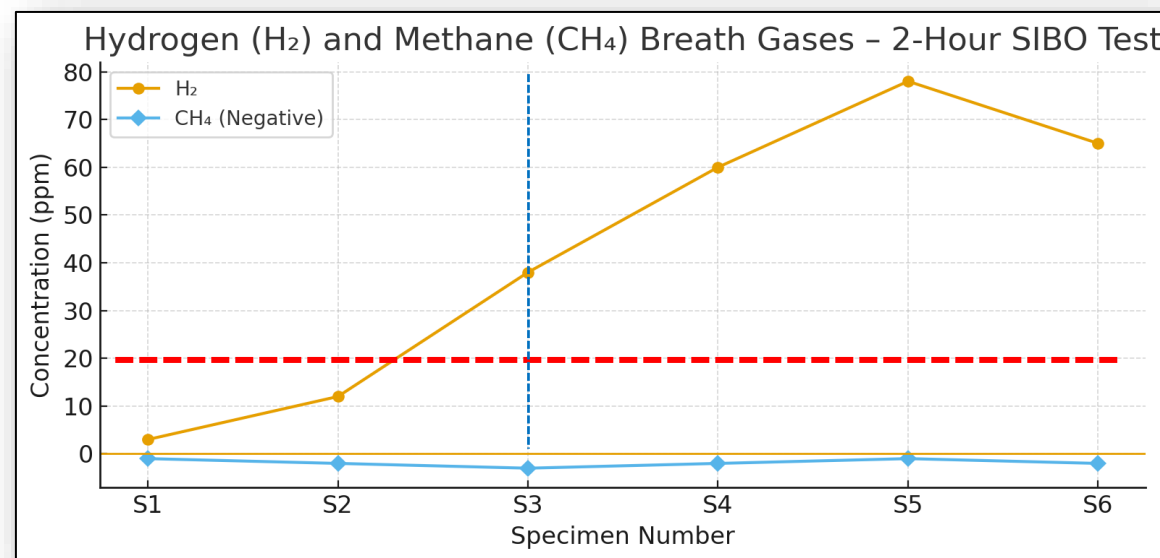
Teste Respiratorio SIBO



Perfil das Vias de Metilação do DNA

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Exames usados no Protocolo LGS:



Gas (ppm)	Baseline 0 min (S1)	20 min (S2)	40 min (S3)	60 min (S4)	90 min (S5)	120 min (S6)
H ₂	2	9	38	60	78	64
CH ₄	-2	-5	-11	-9	-4	-6

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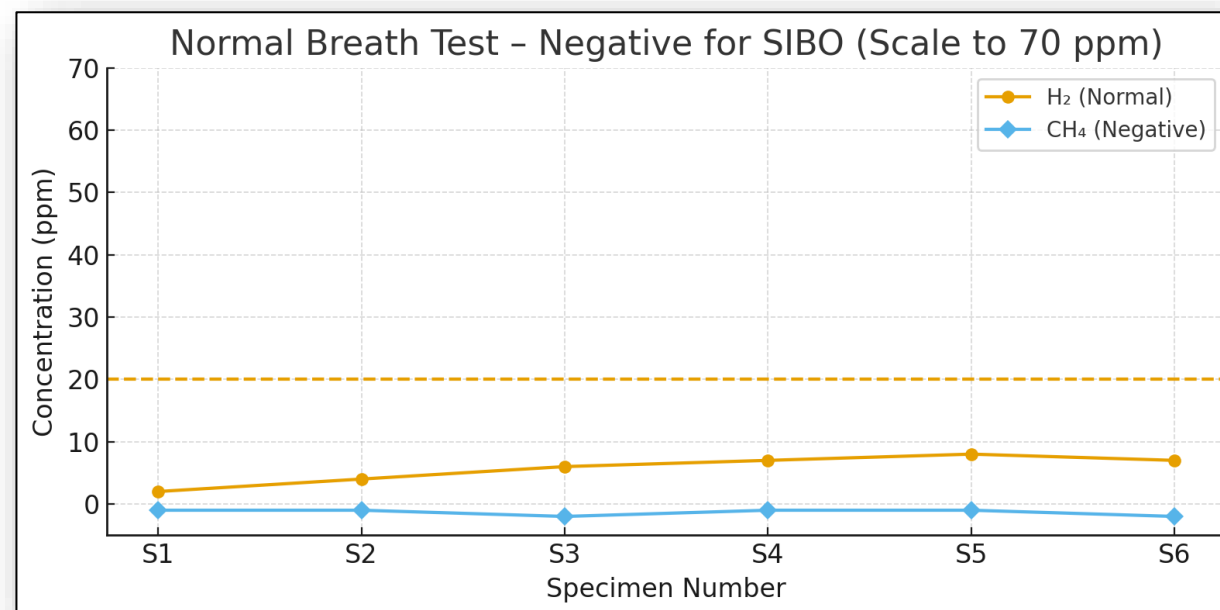
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Teste Respiratorio SIBO

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Exames usados no Protocolo LGS:



Gas (ppm)	Baseline 0 min (S1)	20 min (S2)	40 min (S3)	60 min (S4)	90 min (S5)	120 min (S6)
H ₂	3	6	8	9	10	9
CH ₄	0	0	-3	1	1	-2

Teste Respiratorio SIBO

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THE LEAKY GUT SYNDROME PROTOCOL

Exames usados no Protocolo LGS:



O Perfil de Metilação do DNA identifica 30 SNPs que podem influenciar a saúde e o risco de doenças:

ACAT – 1-02
 AHCY – 19
 AHCY – 1
 AHCY – 2
 BHMT – 1
 BHMT – 2
 BHMT – 4
 BHMT – 8
 CBS – A360A
 CBS – C699T
 CBS – N212N

COMT – 61
 COMT – H62H
 COMT – V158M
 MAO A – R297R
 MTHFR – 3
 MTHFR – A1298C
 MTHFR – C677T
 MTR – A2756G
 MTRR – 11
 MTRR – A66G
 MTRR – H595Y

MTRR – K350A
 MTRR – R415T
 MTRR – S257T
 NOS – D298E
 SHMT – C1420T
 SUOX – S370S
 VDR – Fok1
 VDR – Taq1

Test requisition #
 ORDER:
 CLIENT REF:
 PATIENT ID:
 SEX:
 AGE: DOB:

CLIENT #:
 DNA Methylation Pathway Profile; Buccal Swab

GENE NAME / VARIATION	MUTATION NOT PRESENT	MUTATION(S) PRESENT	CALL	
SHMT/C1420T		+/-	Hetero	
AHCY1	-/-		A	
AHCY2	-/-		T	
AHCY19	-/-		A	
MTHFR/C677T		+/-	Hetero	
MTHFR/A1298C	-/-		A	
MTHFR/3	-/-		C	
MTR/A2756G		+/-	Hetero	
MTRR/A66G		+/+	G	
MTRR/H595Y	-/-		C	
MTRR/K350A	-/-		A	
MTRR/R415T	-/-		C	
MTRR/S257T	-/-		T	
MTRR/11	-/-		G	
BHMT/1		+/-	Hetero	
BHMT/2	-/-		C	
BHMT/4	-/-		A	
BHMT/8	-/-		C	
CBS/C699T		+/+	T	
CBS/A360A		+/-	Hetero	
CBS/N212N	-/-		C	
COMT/V158M	-/-		G	
COMT/H62H	-/-		C	
COMT/61	-/-		G	
SUOX/S370S	-/-		C	
VDR/Taq1	-/-		C	
VDR/Fok1	-/-		C	
MAOA		+/+	T	
NOS/D298E	-/-		G	
ACAT/1-02	-/-		G	

Minus "-" represents no mutation
 Plus "+" represents a mutation
 "-/-" indicates there is no mutation
 "+/-" indicates there is one mutation
 "+/+" indicates there is a double mutation

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Exames usados no Protocolo LGS:



Gene & Variation	rsID	Alleles	Result
CYP1A1 C2453A	rs1799814	GG	-/-
CYP1A2 164A>C	rs762551	AC	+/-
CYP1B1 L432V	rs1056836	CG	+/-
CYP2A6*2 1799T>A	rs1801272	AA	-/-
CYP2C9*2 C430T	rs1799853	CC	-/-
CYP2C9*3 A1075C	rs1057910	AA	-/-
CYP2C19*17	rs12248560	CC	-/-
CYP2D6 S486T	rs1135840	CC	+/+
CYP2D6 2850C>T	rs16947	--	No call
CYP2E1*1B 9896C>G	rs2070676	CG	+/-
CYP2E1*4 4768G>A	rs6413419	GG	-/-
CYP3A4*1B	rs2740574	TT	-/-
CYP3A4*2 S222P	rs55785340	AA	-/-
CYP3A4*3 M445T	rs4986910	AA	-/-
CYP3A4*16 T185S	rs12721627	GG	-/-
GSTP1 I105V	rs1695	AG	+/-
GSTP1 A114V	rs1138272	CC	-/-
SOD2 A16V	rs4880	GG	+/+
NAT1 R64W	rs1805158	CC	-/-
NAT2 I114T	rs1801280	CT	+/-
NAT2 R197Q	rs1799930	AG	+/-
NAT2 G286E	rs1799931	GG	-/-
NAT2 R64Q	rs1801279	GG	-/-
NAT2 K268R	rs1208	AG	+/-

Gene & Variation	rsID	Alleles	Result
COMT V158M	rs4680	AG	+/-
COMT H62H	rs4633	CT	+/-
COMT P199P	rs769224	AG	+/-
VDR Bsm	rs1544410	CC	-/-
VDR Taq	rs731236	AA	+/+
MTHFR C677T	rs1801133	GG	-/-
MTHFR A1298C	rs1801131	TT	-/-
MTR A2756G	rs1805087	AA	-/-
MTRR A66G	rs1801394	AA	-/-
MTRR H595Y	rs10380	CC	-/-
MTRR K350A	rs162036	AA	-/-
MTRR R415T	rs2287780	CC	-/-
BHMT-02	rs567754	TT	+/+
CBS C699T	rs234706	GG	-/-
SHMT1 C1420T	rs1979277	AG	+/-

geneticgenie

Homozygous	+/+
Heterozygous	+/-
Normal	-/-

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Detoxification Superoxide Dismutase (SOD2 A16V) – Homozygous Variant (+/+)



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Exames usados no Protocolo LGS:



Receptor de Vitamina D (VDR) – Importante

☐ VDR Taq: AA (+/+) → variante homozigota

☐ VDR Bsm: CC (-/-) → normal

Interpretação:

O VDR Taq (+/+) está associado a:

☐ Sinalização reduzida do receptor de vitamina D

☐ Maior necessidade de vitamina D

☐ Menor modulação imune

☐ Metabolismo mais lento de dopamina (quando combinado com variantes COMT)

→ **Clinicamente:** Este paciente costuma responder melhor a doses mais altas de vitamina D.

SOD2 A16V – Variante Homozigota (+/+)

Esse é um dos achados mais importantes do painel.

SOD2 +/+ causa:

• Redução da capacidade antioxidante mitocondrial

• Maior estresse oxidativo

• Risco aumentado de inflamação crônica

• Relação forte com autoimunidade cutânea (como vitiligo)

→ **Clinicamente:** focar em proteção mitocondrial

— NAC, ALA, CoQ10, Glutathione, Vitamina C, Ômega-3, polifenóis.



Estudo Genetico:



Resumo Funcional no Estilo

✓ Metilação intacta (sem defeitos de MTHFR)

✓ **Alto risco de estresse oxidativo (SOD2 +/+)** → muito relevante para vitiligo

✓ Metabolismo lento de catecolaminas (COMT variantes)

→ Sensível a metil-B12/metilfolato

✓ **VDR Taq +/+**

→ Necessidade aumentada de vitamina D

✓ **Detox fase I e II com variantes relevantes (CYP2D6, NAT2)**

→ Mais lento para metabolizar drogas e toxinas

Exames usados no Protocolo LGS:



CLIENT #:

DNA Methylation Profile; Buccal Swab

GENE NAME / VARIATION	MUTATION NOT PRESENT	MUTATION(S) PRESENT	CALL
SHMT/C1420T	-/-	+/-	Hetero
AHCY1	-/-	-/-	A
AHCY2	-/-	-/-	T
AHCY19	-/-	-/-	A
MTHFR/C677T	-/-	+/-	Hetero
MTHFR/A1298C	-/-	-/-	A
MTHFR/3	-/-	-/-	C
MTR/A2758G	-/-	+/-	Hetero
MTRR/A68G	-/-	+/+	G
MTRR/H595Y	-/-	-/-	C
MTRR/K350A	-/-	-/-	A
MTRR/R415T	-/-	-/-	C
MTRR/S257T	-/-	-/-	T
MTRR/11	-/-	-/-	G
BHMT/1	-/-	+/-	Hetero
BHMT/2	-/-	-/-	C
BHMT/4	-/-	-/-	A
BHMT/8	-/-	-/-	C
CBS/C690T	-/-	+/+	T
CBS/A380A	-/-	+/-	Hetero
CBS/N212N	-/-	-/-	C
COMT/V158M	-/-	-/-	G
COMT/H52H	-/-	-/-	C
COMT/I61	-/-	-/-	G
SUOX/S370S	-/-	-/-	C
VDR/Taq1	-/-	-/-	C
VDR/Fok1	-/-	-/-	C
MAOA	-/-	+/+	T
NOS/D298E	-/-	-/-	G
ACAT1-02	-/-	-/-	G

Test requisition #

ORDER:

CLIENT REF:

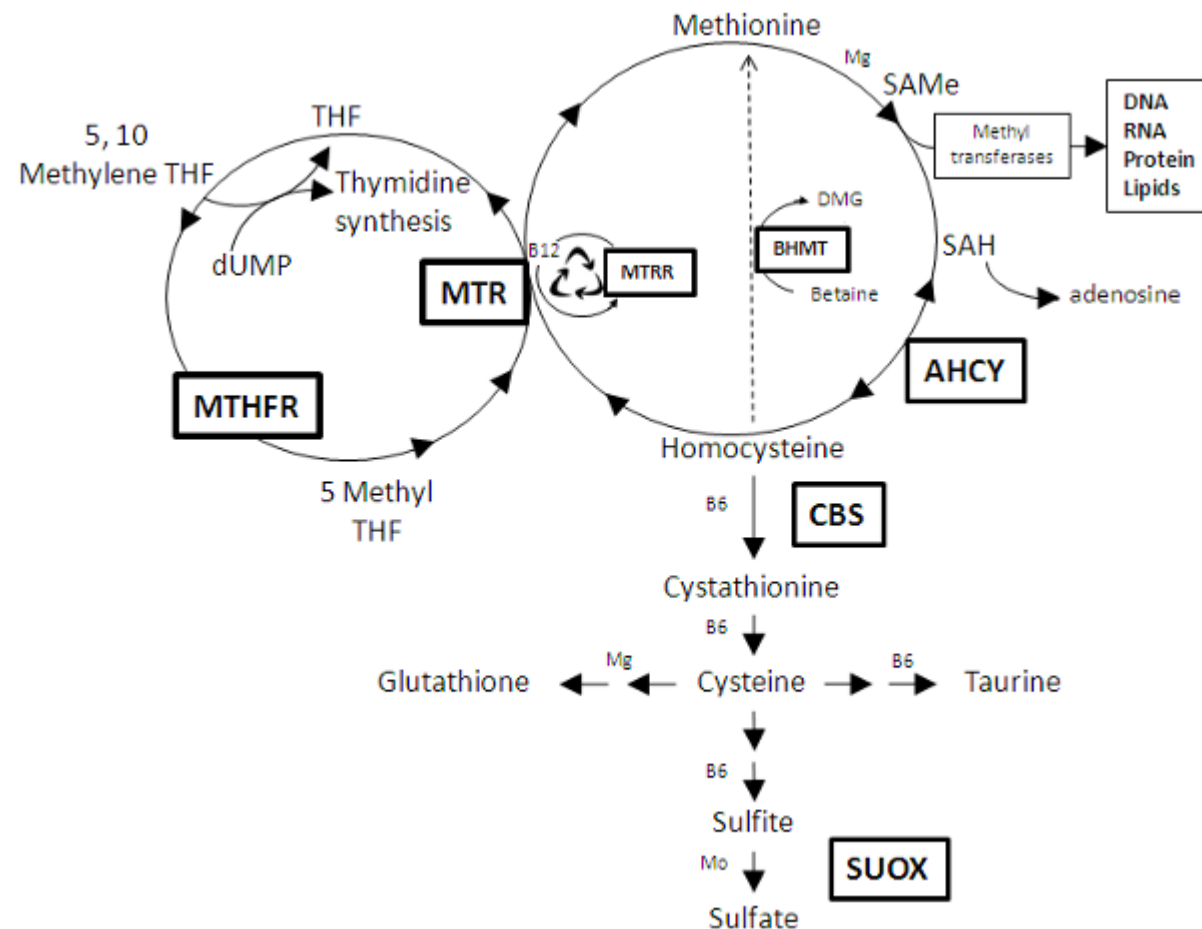
PATIENT:

ID:

SEX:

AGE: DOB:

Minus "-" represents no mutation
Plus "+" represents a mutation
"/-" indicates there is no mutation
"+/-" indicates there is one mutation
"+/+" indicates there is a double mutation



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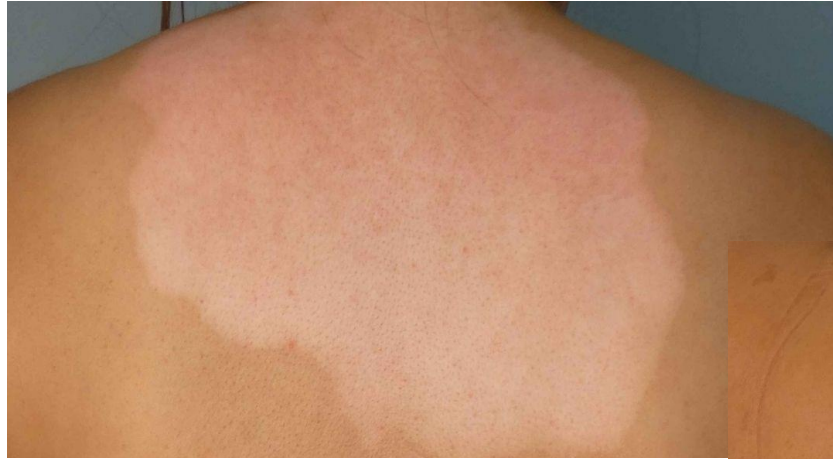
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FUNCIONAL



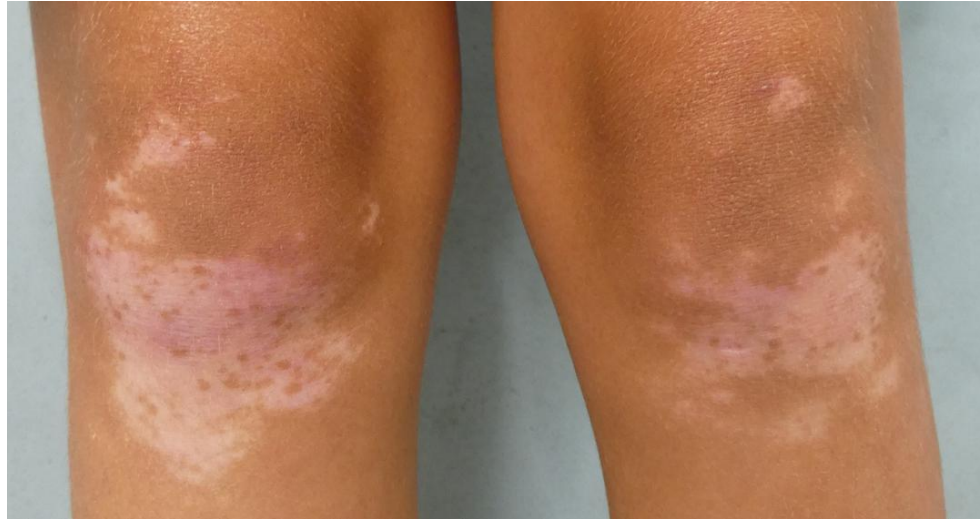
THE LEAKY GUT SYNDROME PROTOCOL

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GI MAP



GI MAP



Os 3 Pilares do LGS Protocol

Modulação do Microbioma



Restaurar a Diversidade Microbioma

- ☐ Estratégias dietéticas
- ☐ Prebióticos e fibras específicas
- ☐ Probióticos direcionados
- ☐ Pós-bióticos e SCFAs (butirato, propionato, acetato)
- ☐ Redução de inflamação e reparo da barreira mucosa.

Modulação do Sistema Imune



Altas Doses de Vitamina D

- ☐ Aplicação clínica da terapia de imunomodulação com vitamina D.
- ☐ Uso criterioso de cofatores essenciais: K2, magnésio, ômega-3, vitaminas B, zinco, selênio, entre outros.
- ☐ Redução de inflamação crônica e autorregulação imunológica.

Compensação Epigenética



Compensação Polimorfismos (SNPs)

- ☐ Identificação dos polimorfismos envolvidos.
- ☐ Correção desses polimorfismos, como no ciclo de metilação, vias de detoxificação, estresse oxidativo, entre outros.
- ☐ Uso de suplementos (vitaminas, minerais, aminoácidos, enzimas, hormônios etc.) que auxiliam na ativação de vias epigenéticas de reparação tecidual.

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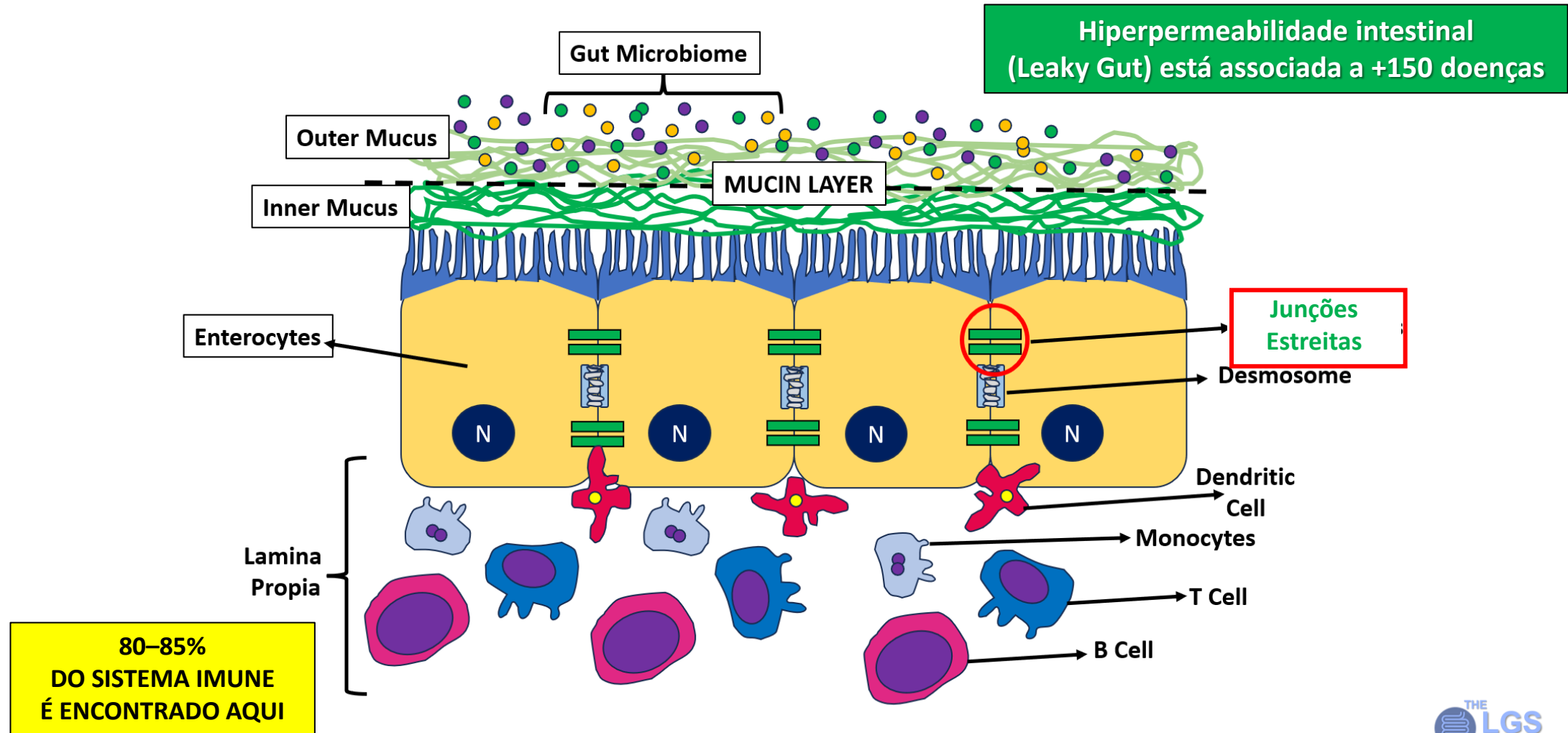


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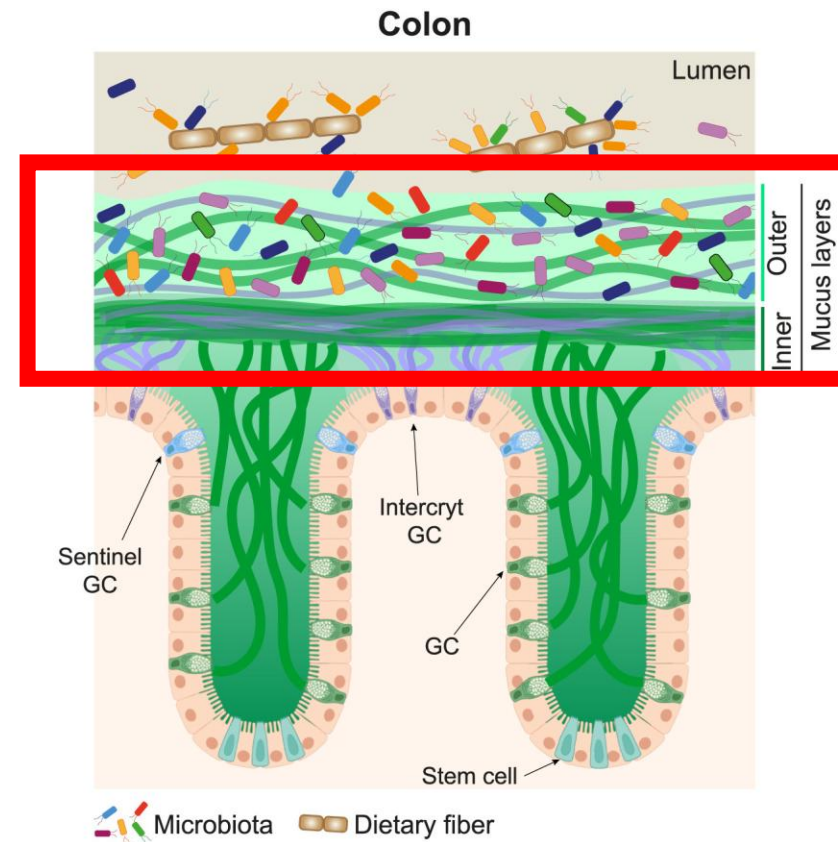
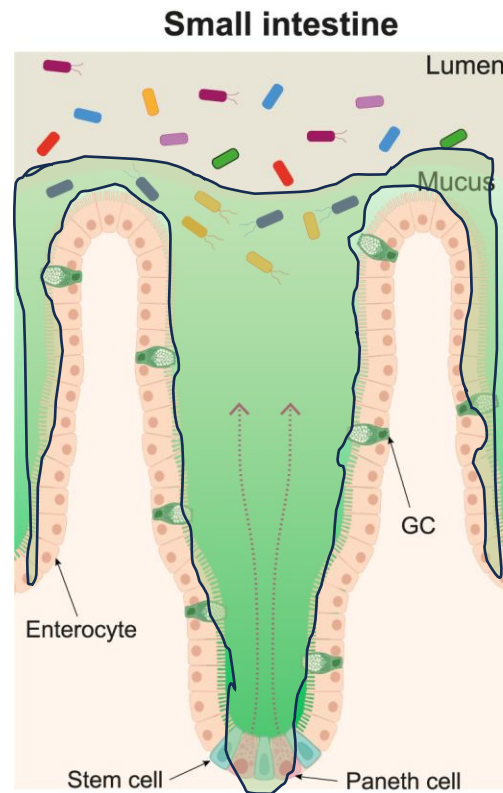
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A Barreira Intestinal: O Verdadeiro Órgão da Imunidade



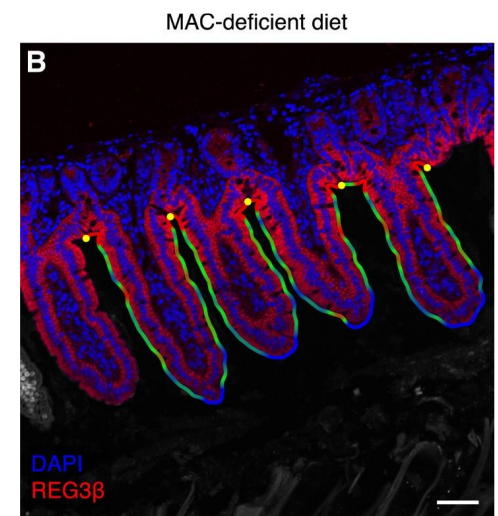
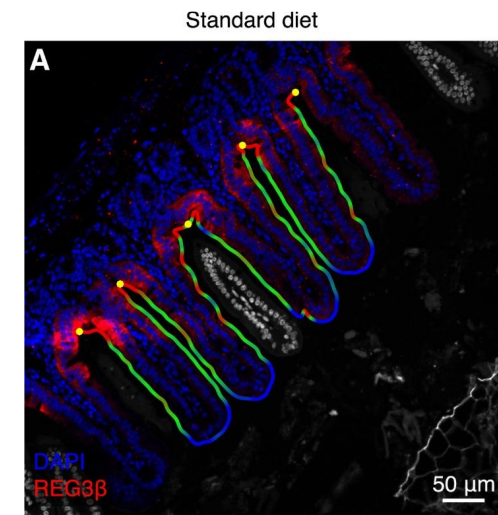
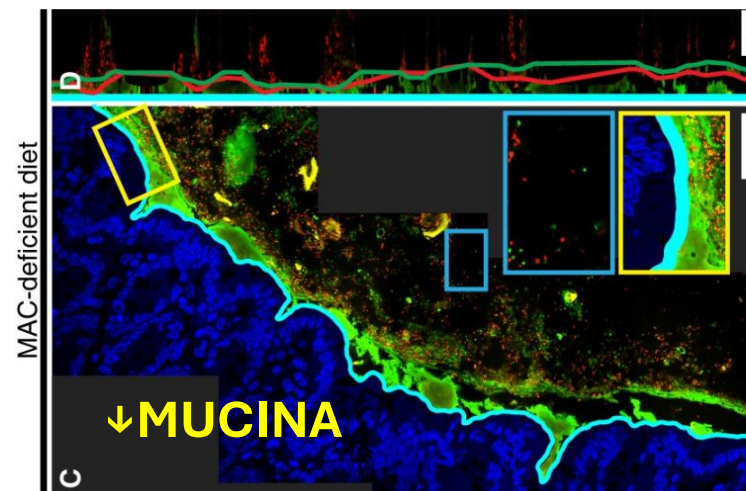
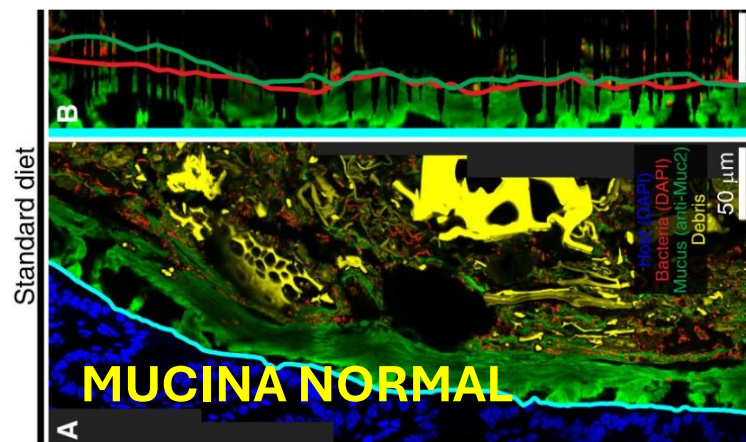
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A Barreira Intestinal: O Verdadeiro Órgão da Imunidade

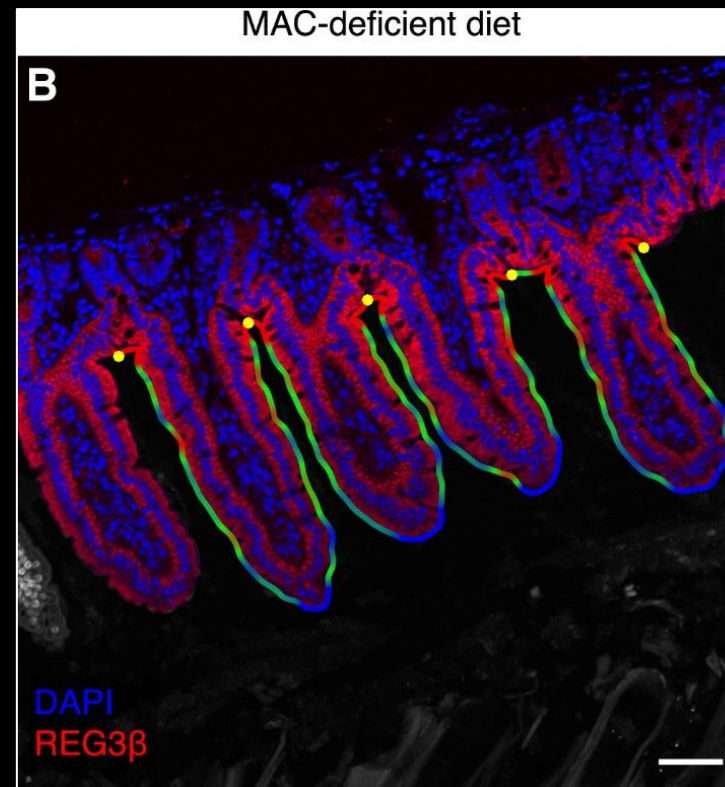
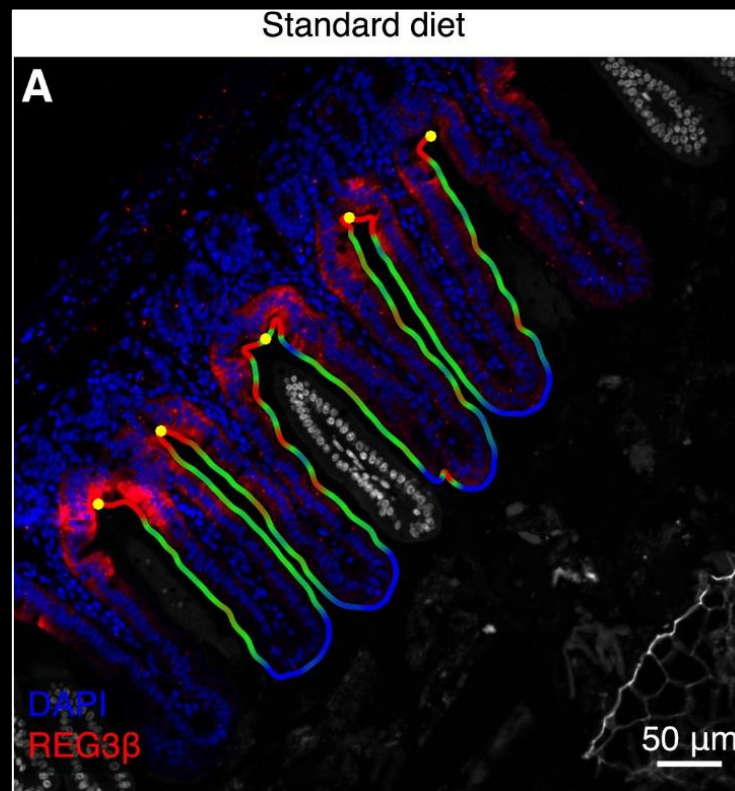


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CAMADA DE MUCINA



CAMADA DE MUCINA



As imagens mostram cortes de tecido intestinal vistos em microscopia de fluorescência, onde os núcleos aparecem em azul e a proteína REG3β em vermelho. Na primeira imagem observa-se uma camada de mucina preservada, funcionando como barreira protetora e mantendo as células de defesa concentradas na base das criptas, sem contato direto com a microbiota. Já na segunda imagem há deficiência de mucina, o que permite maior aproximação das bactérias às células epiteliais e leva a uma interação mais intensa entre o sistema imune e a microbiota, cenário que pode favorecer processos inflamatórios.

doi: 10.1016/j.chom.2015.09.002.

VITAMINA D NO MUNDO



47,9 % da população mundial tem deficiência de vitamina D
(**25(OH)D < 20 ng/mL**).

76,6 % tem níveis de 25(OH)D abaixo de **30 ng/mL**
(insuficiência + deficiência)

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Vitamina D → Hormônio

+80 Funções Metabólicas

Regula + 3500 genes (fetal)

+3000 genes no adulto

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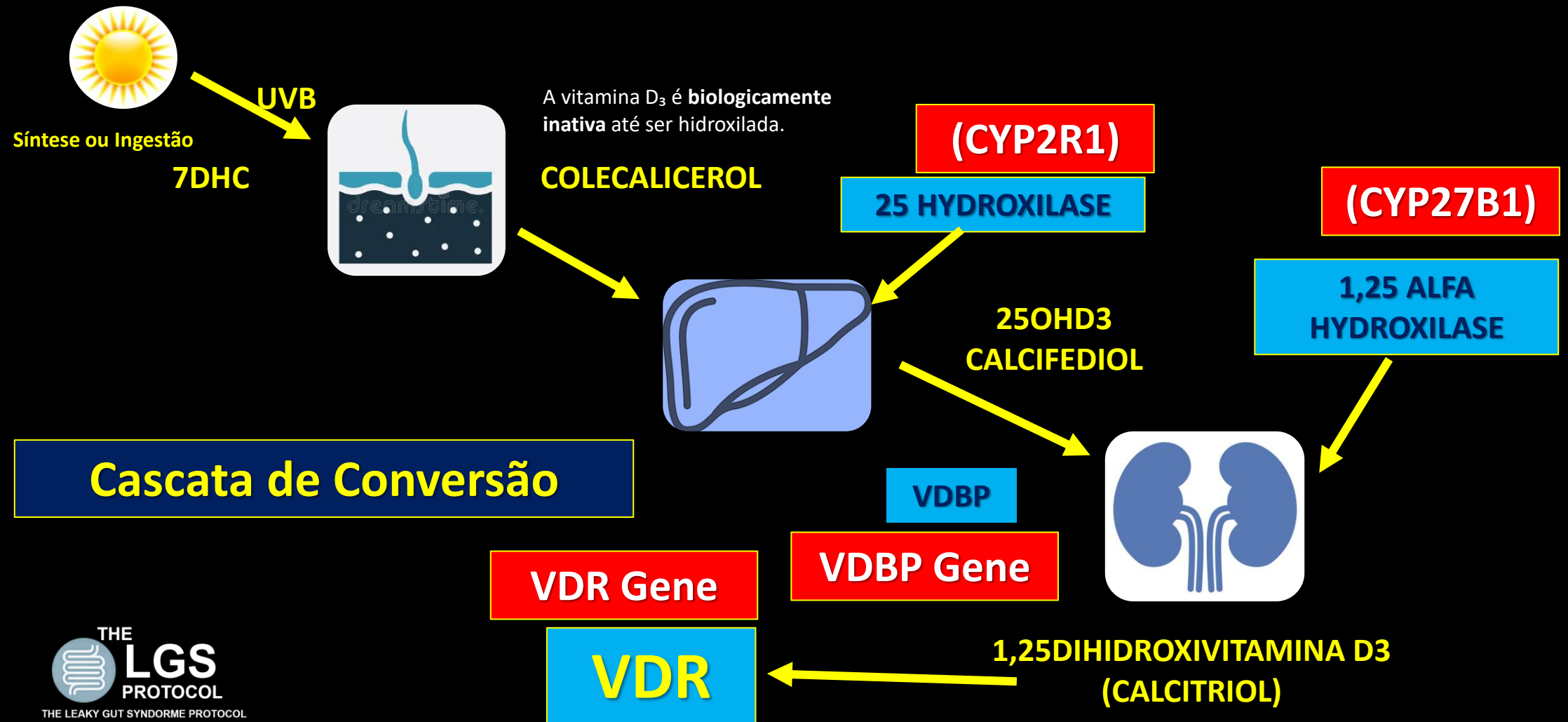


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FISIOLOGIA - VITAMINA D



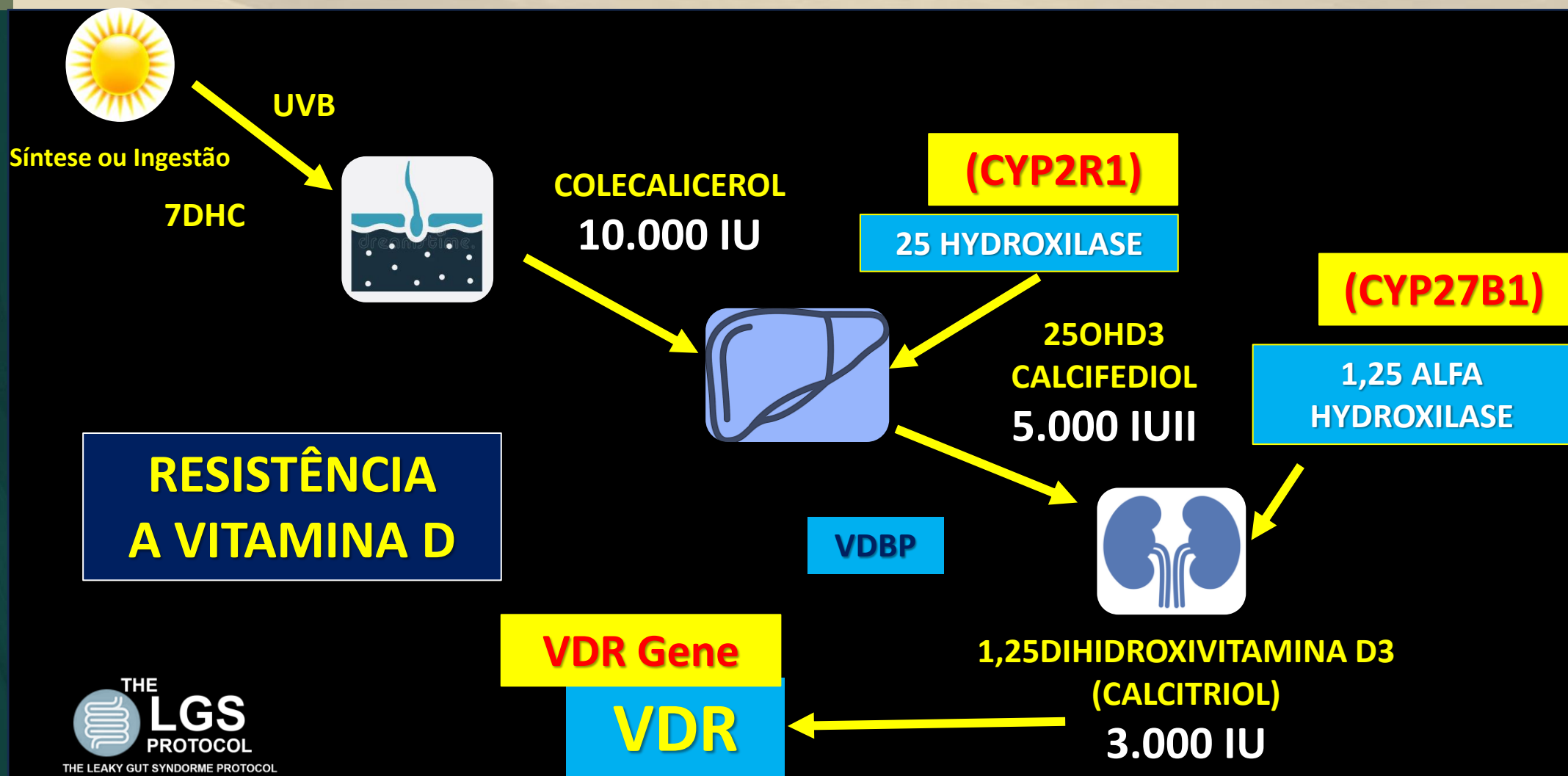
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THE
LGS
PROTOCOL
THE LEAKY GUT SYNDROME PROTOCOL

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POLIMORFISMOS (SNP's)



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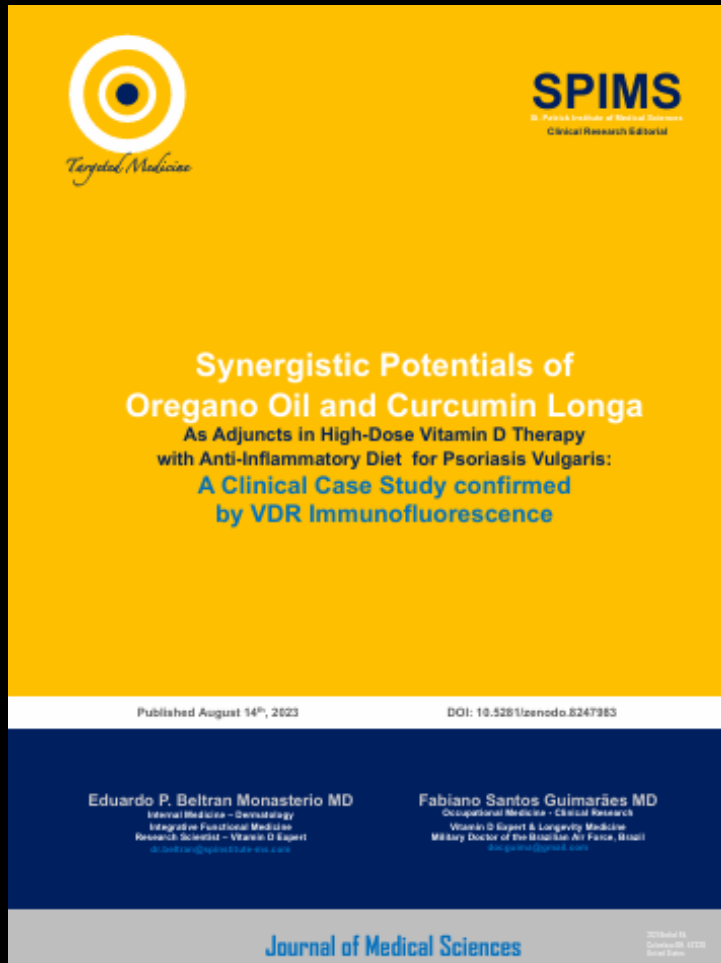
THE
LGS
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THE LEAKY GUT SYNDROME PROTOCOL



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Caso Clínico



PSORIASSE VULGAR



Dr. Eduardo Beltran



Dr. Fabiano Guimarães



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Lab Results	03/18/2019	07/16/2019	10/23/2019	Normal Reference Values
25OHD3 - Calcifediol	46 ng/ml	123 ng/ml	148 ng/ml	20-100 ng/ml
1,25OHD3 - Calcitriol	72 pg/ml	78 ng/ml	76 ng/ml	19.9-79 pg/ml
PTH	81 pg/ml	41 pg/ml	24 pg/ml	13.6 – 85.8 pg.ml
Serum Ca ⁺	1.24 mmol/L	1.20 mmol/L	1.23 mmol/L	1,00 a 1,35 mmol/L
Ionized Ca ⁺	8.9 mg/dL	9.3 mg/dL	9.6 mg/dL	8,4 a 10,2 mg/dL
Urea	23 mg/dL	28 mg/dL	26 mg/dL	15,0 a 36,0 mg/dL
Creatinine	1.0 mg/dL	0.9 mg/dL	1.0 mg/dL	0,7 a 1,2 mg/dL
ALT (Alanine Aminotransferase)	44 U/L	24 U/L	17 U/L	<35 U/L (women)
AST (Aspartate Aminotransferase)	41 U/L	26 U/L	15U/L	14,0 a 36,0 U/L
Insulin	28.2 µU/mL	10.4 µU/mL	7.7 µU/mL	2,0 a 25,0 µU/mL


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GI MAP Comensal Bacteria/Keystone Bacteria	Result	Normal Reference Range
Bacteroides fragilis:	3.42e11 High	1.6e9 to 2.5e11
Bifidumbacterium:	5.28e7 Low	> 6.0e7
Escherichia spp:	1.54e10 High	3.7e6 to 3.8e8
Lactobacillus:	9.08e2 Low	8.6e5 to 6.2e8
Akkermansia muciniphila:	Undetectable	1.01e3 to 8.2e6
Roseburia spp	1.56e8	5.0e7 to 2.0e10
Bacterial Phyla	Result	Normal Reference Range
Bacteroidetes:	7.18e12 High	8.6e11 to 3.3e12
Firmicutes:	2.09e10 Low	5.7e10 to 3.0e11
Firmicutes:Bacteroidetes Ratio	0.00	< 1.0
Opportunistic/Overgrowth Microbes - Dysbiotic & Overgrowth Bacteria	Results	Normal Reference Range
Staphylococcus spp:	1.11e4 High	< 1.00e4
Inflammatory & Autoimmune-Related Bacteria:	Result	Normal Reference Range
Klebsiella spp:	7.23e4 High	< 5.00e3
Commensal Inflammatory & Autoimmune-Related Bacteria:	Result	Normal Reference Range
Escherichia spp:	4.23e9 High	< 3.80e9
Fungi/Yeast:	Result	Normal Reference Range
Candida spp.:	1.98e6 High	< 5.00e3
Candida Albicans:	1.83e5 High	< 5.00e2

INTESTINAL HEALTH MARKERS			
DIGESTION	Result		Reference
Steatocrit	< dL ▼		< 15 %
Elastase-1	>750		> 200 ug/g
GI MARKERS			
β-Glucuronidase	639 ▼		< 2486 U/mL
Occult Blood - FIT	3 ▼		< 10 ug/g
IMMUNE RESPONSE			
Secretory IgA	<210 L ▼		510 - 2010 ug/g
Anti-gliadin IgA	21 ▼		< 175 U/L
Eosinophil Activation Protein (EON, EPX)	0.40 ▼		< 2.34 ug/g
INFLAMMATION			
Calprotectin	429 H		< 173 ug/g
ADD-ON TESTS			
Zonulin	358 H		< 175 ng/g

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Caso Clínico – SNPs found in Patient



Genes:	CYP27B1	VDR	MTHFR	MTR
SNP	rs10877012 rs4646536	rs2228570 (FokI) rs731236 (TaqI) rs7975232 (ApaI)	rs1801133 (C677T) rs1801131 (A1298C)	rs1805087 (A2756G)
Function	Involved in Vitamin D activation from 25OHD3 (calcifediol) to 1,25OHD3 (calcitriol).	Vitamin D Receptor responsible for responding to vitamin D levels.	Methylenetetrahydro- folate Reductase enzyme responsible for converting 5,10- methylenetetrahydro- folate to 5- methyltetrahydrofolate	Methionine Synthase Reductase enzyme responsible for converting homocysteine to methionine
Health Implication	May influence vitamin D metabolism conversion and overall vitamin D status of active vitamin D3 (Calcitriol).	Impact on Calcium absorption, bone health, and immune system function.	Associated with altered folate metabolism, potentially affecting methylation processes.	Involved in homocysteine and methionine metabolism, impacting DNA methylation and protein synthesis.

BIOPSIA DE PELE

(Punch Biopsy)



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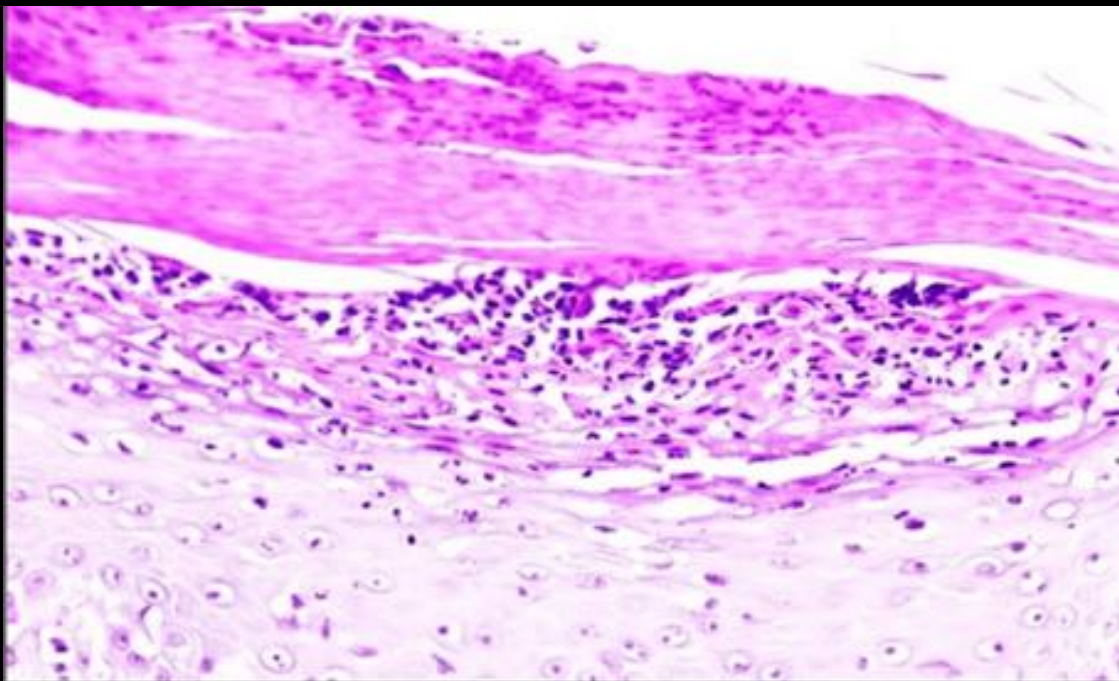
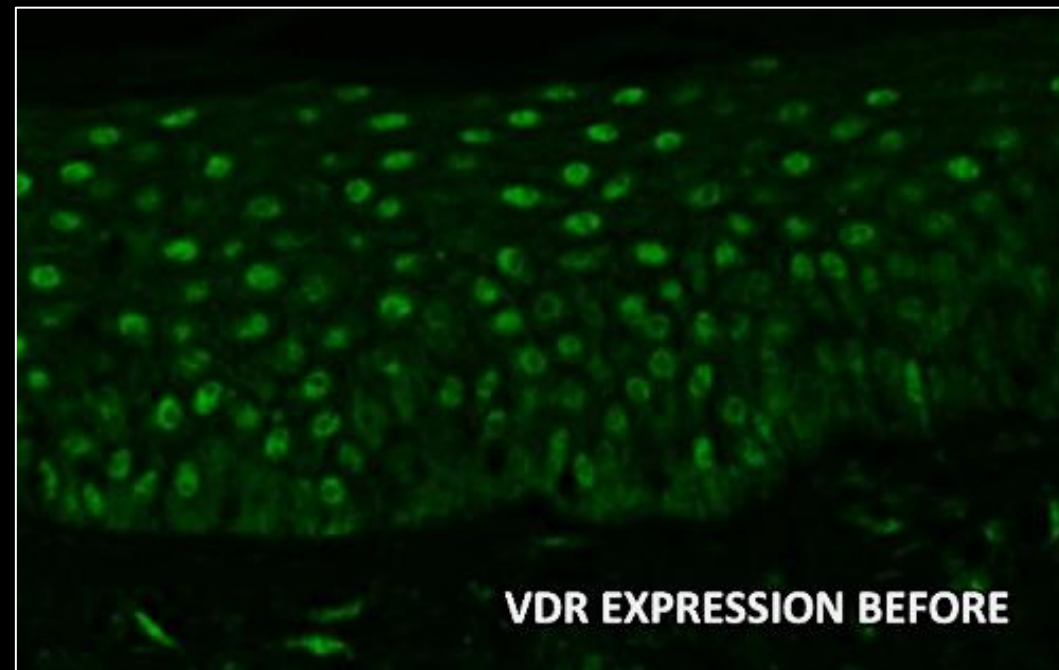


FIGURE 1: Skin biopsy Anatomic Location: Left forearm

The skin biopsy under microscopic examination displays distinctive features consistent with psoriasis. The epidermis exhibits significant hyperplasia, characterized by elongation and thickening of the rete ridges. Notable findings include focal parakeratosis, uniform acanthosis, and elongated epidermal projections known as "regular acanthosis." Additionally, Munro microabscesses, aggregations of neutrophils within the epidermis, are evident. In the underlying dermis, there is a mild perivascular lymphocytic infiltrate.



VDR EXPRESSION BEFORE

IMMUNOFLOURESENCIA DOS VDR

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3 Meses Depois

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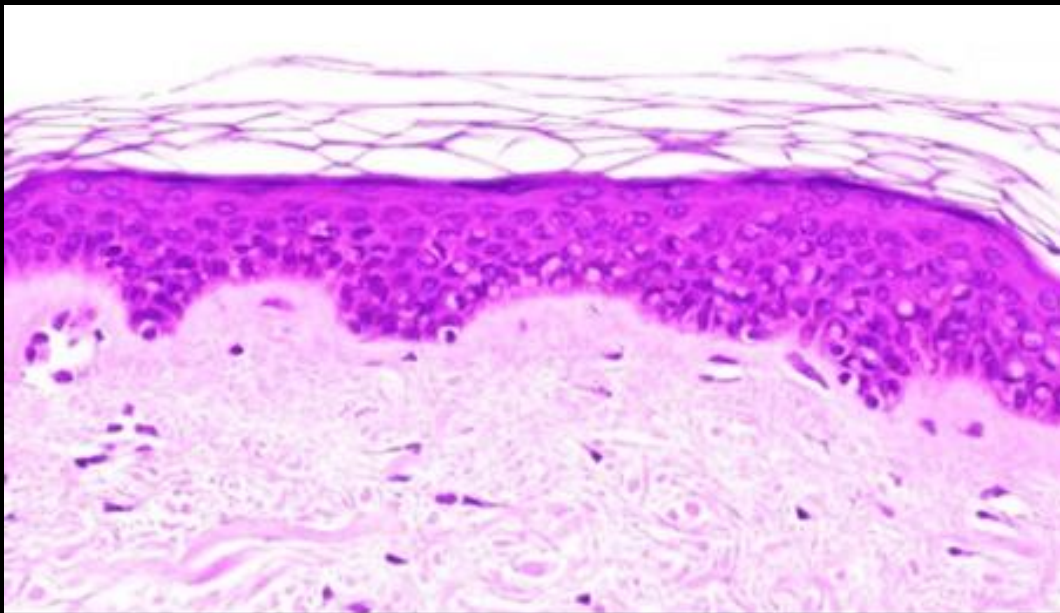
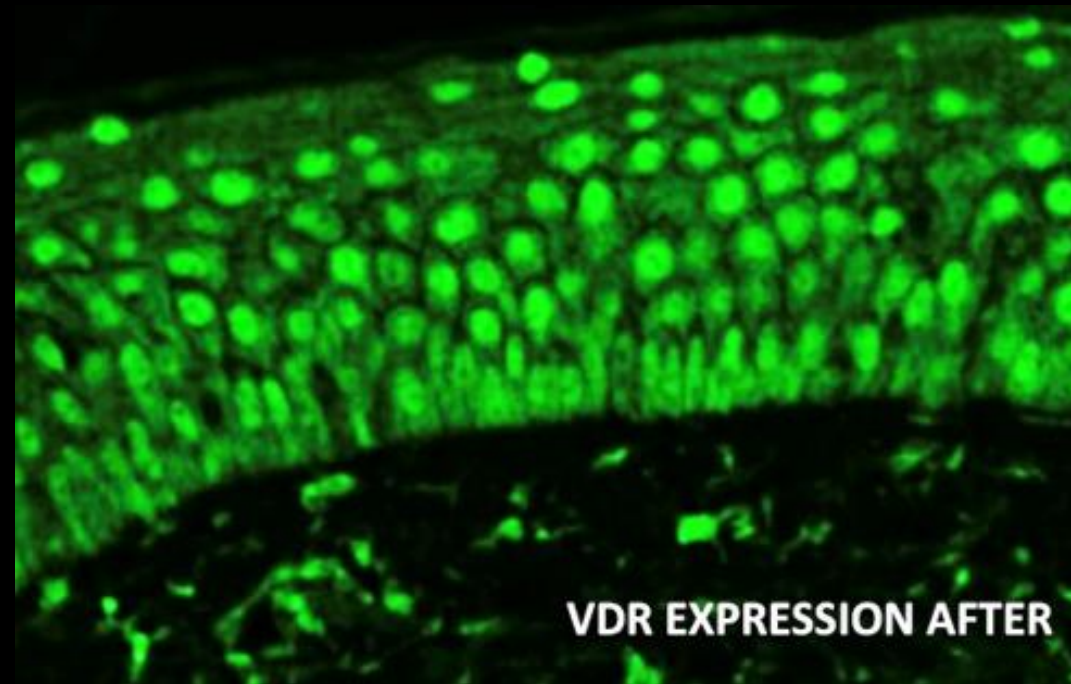


FIGURE 3: Second Skin Biopsy - Anatomic Location: Left forearm (Near original site)

Microscopic Description: Upon examining the skin biopsy, the findings indicate substantial improvement in the skin lesions, now displaying an almost normal appearance. The epidermis exhibits regular thickness and architecture, with no signs of hyperplasia or elongation of rete ridges. Parakeratosis and acanthosis are not present. The absence of regular acanthosis and previously noted Munro microabscesses suggests near-complete resolution. The underlying dermis appears unremarkable, showing no significant inflammation or infiltrates.



VDR EXPRESSION AFTER

IMMUNOFLOURESENCIA DOS VDR



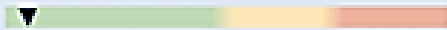



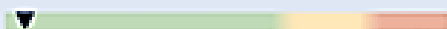

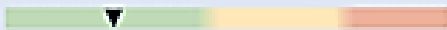
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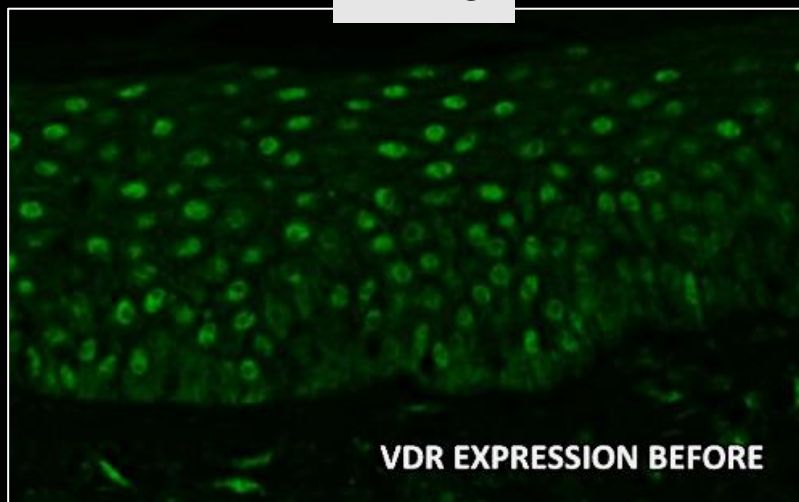
INTESTINAL HEALTH MARKERS			
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INFLAMMATION			
Calprotectin	87		< 173 ug/g
ADD-ON TESTS			
Zonulin	54		< 175 ng/g

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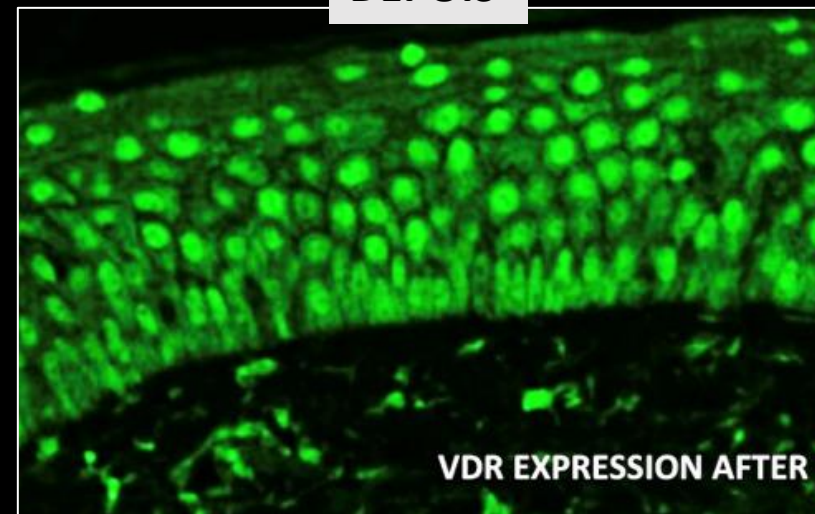
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ANTES



DEPOIS



“Renovação do VDR via Dieta Anti-inflamatória”


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Vitamin D Receptor Renewal Through Anti-inflammatory Diet: (Another Contributing Factor for Vitamin D Resistance)

Version 1.0



Dr. Eduardo Beltran, 2024, "Vitamin D Receptor Renewal Through Anti-inflammatory Diet: (Another Contributing Factor for Vitamin D Resistance)", <https://doi.org/10.7910/DVN/LC0RXQ>, Harvard Dataverse, V1

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Description ⓘ

Vitamin D is an essential nutrient that plays a crucial role in maintaining bone health, immune function, and overall health. Adequate levels of vitamin D are important for proper calcium absorption, regulation of gene expression, and modulation of immune responses. However, patients with autoimmune diseases may have genetic polymorphisms that contribute to vitamin D resistance, such as single nucleotide polymorphisms (SNPs) affecting the genes CYP2R1, CYP27B1, VDR, and VDBP. In addition, these patients may have ongoing leaky gut syndrome due to inflammatory foods, such as gluten, dairy, lectins, and sugars, which can lead to molecular mimicry and the production of autoantibodies. This condition sets the stage for foreign proteins and toxins, including lipopolysaccharides, endotoxins, exotoxins, and mycotoxins, to enter the bloodstream, further inducing

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The LGS Protocol, validated through VDR Immunofluorescence and Microbiome Test Correlation, supports Vitamin D Receptor Renewal via an Anti-inflammatory Diet in a clinical case of Psoriasis Vulgaris.

Eduardo P. Beltran Monasterio ^{1, 2}, Fabiano Santos Guimaraes ³

1. Dermatology - Integrative Functional Medicine, Universidade Gama Filho (Institute of Research and Medical Education), Florianopolis, BRA 2. Research and Development, St. Patrick Institute of Medical Sciences, Columbus, OH, USA 3. Occupational Medicine - Integrative Functional Medicine, Brazilian Air Force, Sao Paulo, BRA

Corresponding author: Eduardo P. Beltran Monasterio, dreduardobeltran@yahoo.com

Abstract

This clinical case study investigates the relationship between the Vitamin D receptor (VDR) and psoriasis using Dr. Eduardo Beltran's 'Leaky Gut Syndrome (LGS) Protocol.' The intervention, combining an anti-inflammatory diet, high-dose vitamin D supplementation, and antimicrobial herbs (particularly oregano oil and turmeric), targeted small intestinal fungal overgrowth induced by Candida. Immunofluorescence analysis of VDR expression in skin biopsies was conducted before and after the intervention. Initial biopsy revealed reduced VDR expression correlating with persistent psoriasis. After four months, significant clinical improvement and increased VDR expression were observed in the follow-up biopsy. Significant improvements were also observed in a follow-up microbiome test. The study explores turmeric's integration for its anti-inflammatory and anti-microbial properties, emphasizing implications for managing chronic inflammatory disorders like psoriasis. The findings support the efficacy of an anti-inflammatory diet in reducing intestinal hyperpermeability, increasing VDR expression, and offer innovative insights for therapeutic strategies in resistant psoriasis.

O Futuro da Medicina Integrativa: Imunofluorescência, Microbioma e Vitamina D

1. Imunofluorescência: um marco inovador

- ❑ Evidencia que **LPS bacteriano** (disbiose → hiperpermeabilidade) **inibe a expressão dos receptores de Vitamina D (VDR)**.
- ❑ Explica o fenômeno da **resistência à Vitamina D**: ↓ VDR = ↓ resposta ao calcitriol, mesmo com doses elevadas.

2. Modulação Intestinal reduz a resistência à Vitamina D

- ❑ Correção da disbiose → **menos LPS** → **mais VDR**.
- ❑ Renovação da mucina (Akkermansia, Roseburia, Faecalibacterium).
- ❑ **Redução natural da necessidade de altas doses de Vitamina D**.
- ❑ Conceito introduzido:
“VDR Renewal through Anti-inflammatory Diet”.

3. Por que isso representa inovação mundial

- ❑ Integra **microbioma + imunomodulação + epigenética**. (**LGS Protocol**)
- ❑ Demonstra que **não existe modulação imunológica eficaz sem modulação intestinal**.
- ❑ Define novo paradigma clínico: **tratar a barreira intestinal é tratar o sistema imune**.

4. Sequenciamento genômico do microbioma: estar 15–20 anos à frente.

- ❑ **Profissionais que usam genômica e Vit. D estão décadas à frente**.
- ❑ Quem não utiliza → **perde 90% dos dados clínicos essenciais**.

O Futuro da Medicina Integrativa: Imunofluorescência, Microbioma e Vitamina D

5. Direções futuras da Medicina Integrativa

- ❑ Genômica + epigenômica + microbioma + IA clínica.
- ❑ Medicina de precisão aplicada ao intestino, imunidade como abordagem terapêutica.

**Modulação
do Microbioma**



**Modulação
do Sistema Imune**



**Compensação
Epigenética**



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O Futuro da Medicina Integrativa: Imunofluorescência, Microbioma e Vitamina D



“O futuro da medicina pertence a quem entende o microbioma, a epigenética e o papel central da vitamina D.”

Dr. Eduardo Beltran

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Muito Obrigado!



Dr. Eduardo Patrick Beltran Monasterio

Internal Medicine – Dermatology

Director of St. Patrick Institute of Medical Sciences

Columbus OH - USA



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