

Dra. Silvia Ferolla

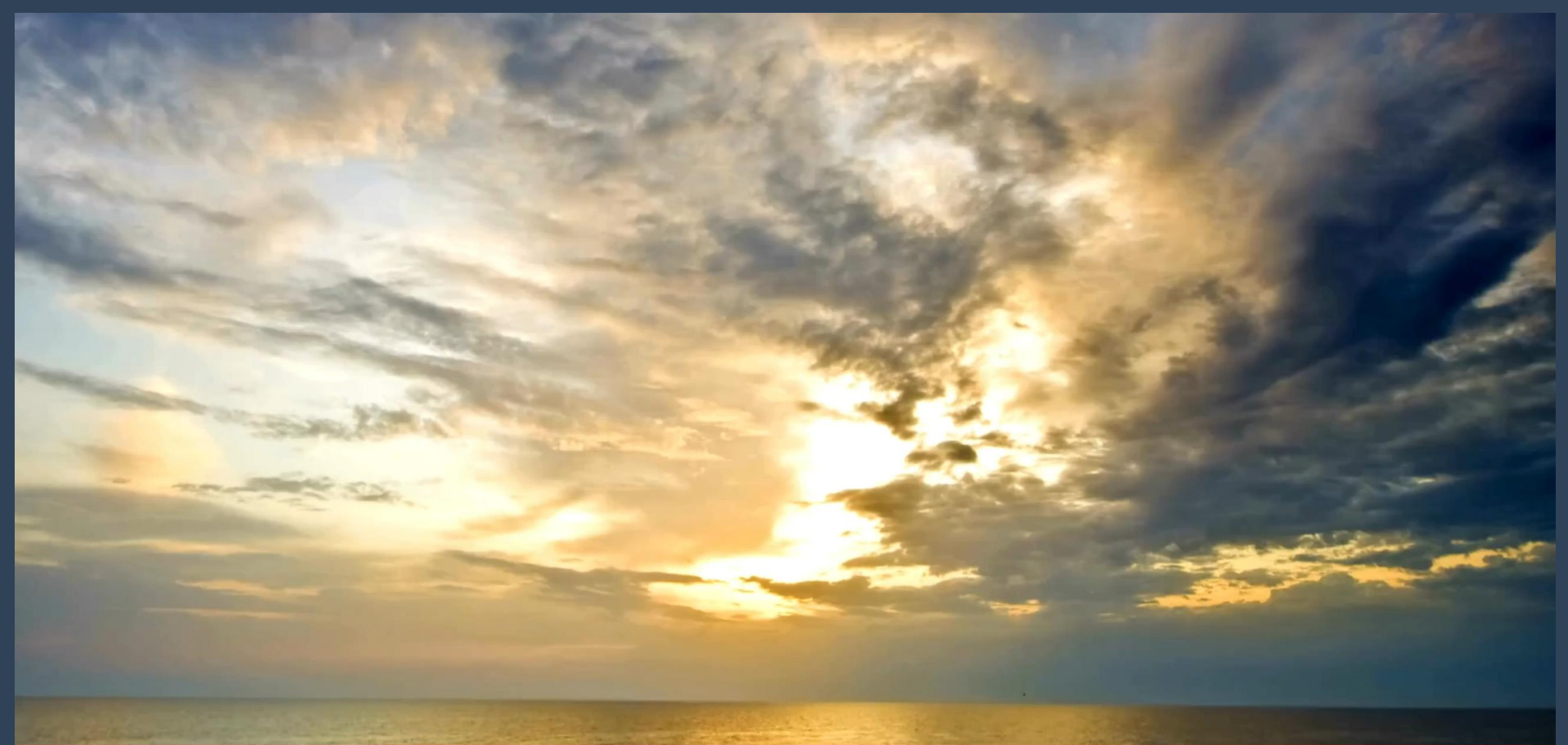


ESTEATO-HEPATITE METABÓLICA:

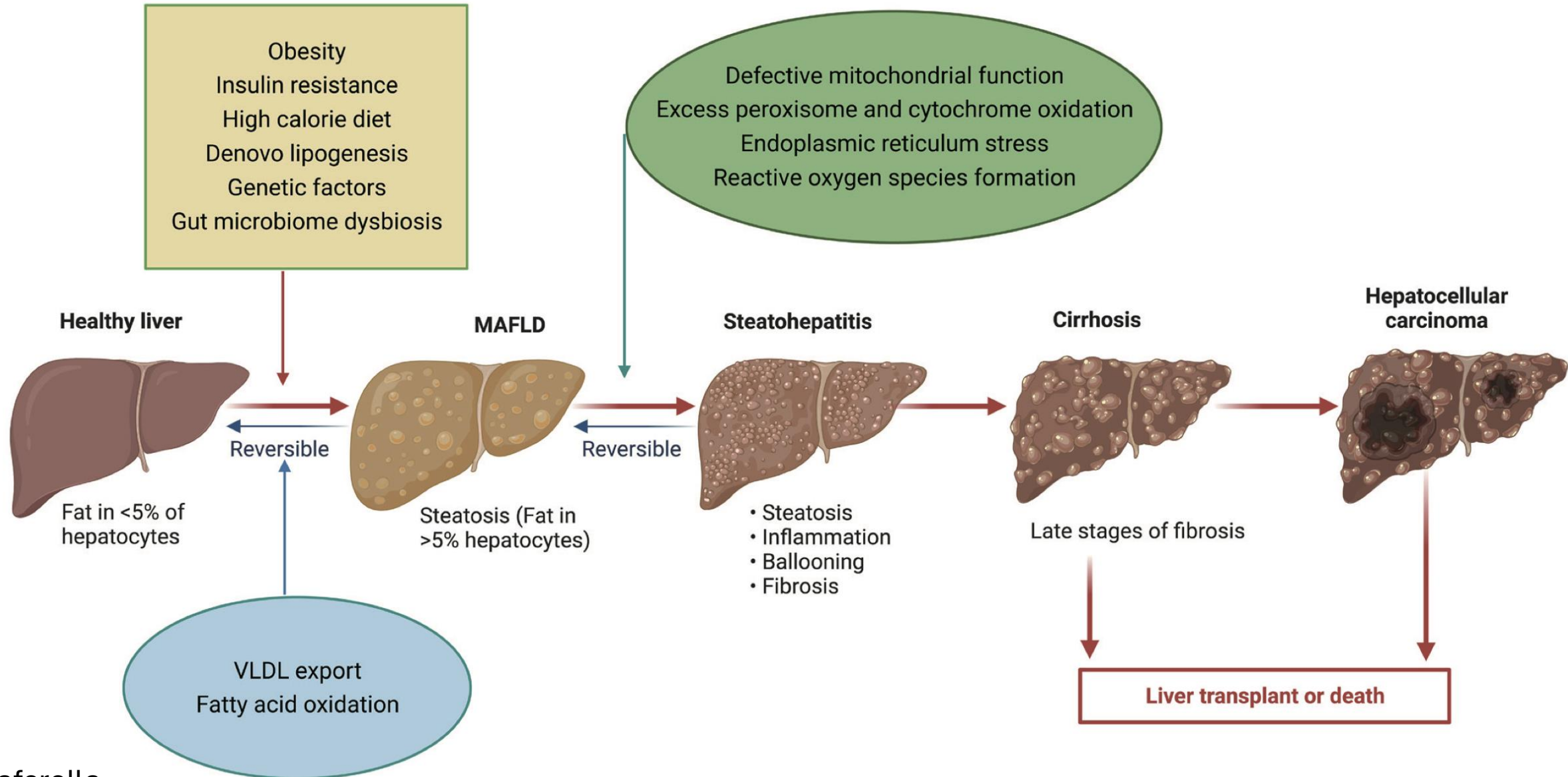
como reduzir inflamação com a
nutrição integrativa ?

PhD e mestre em Saúde do Adulto área de concentração em Aparelho Digestivo UFMG
Autora do Livro Dieta Low LOWFODMAP: fundamentos e aplicações clínicas

@silviaferolla



Espectro da Doença Hepática Gordurosa Metabólica (MALFD)



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MAFLD diagnosis:

Hepatic steatosis plus one of the 3 criteria

- Overweight or obesity
- Type 2 diabetes mellitus
- At least two metabolic risk abnormalities

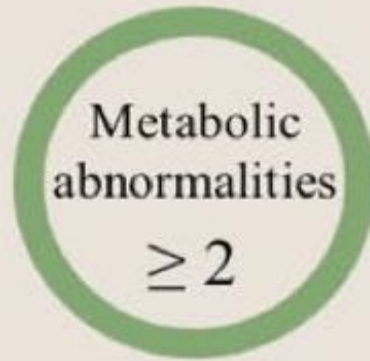


or

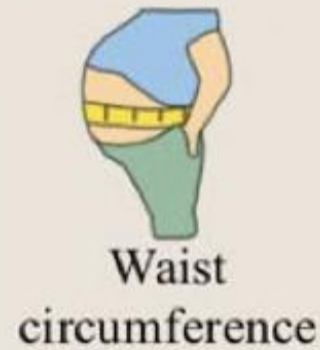


T2DM

or



Metabolic
abnormalities
 ≥ 2



Waist
circumference



Hypertension



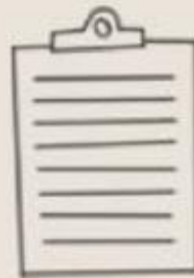
Triglycerides



HDL



Prediabetes



HOMA

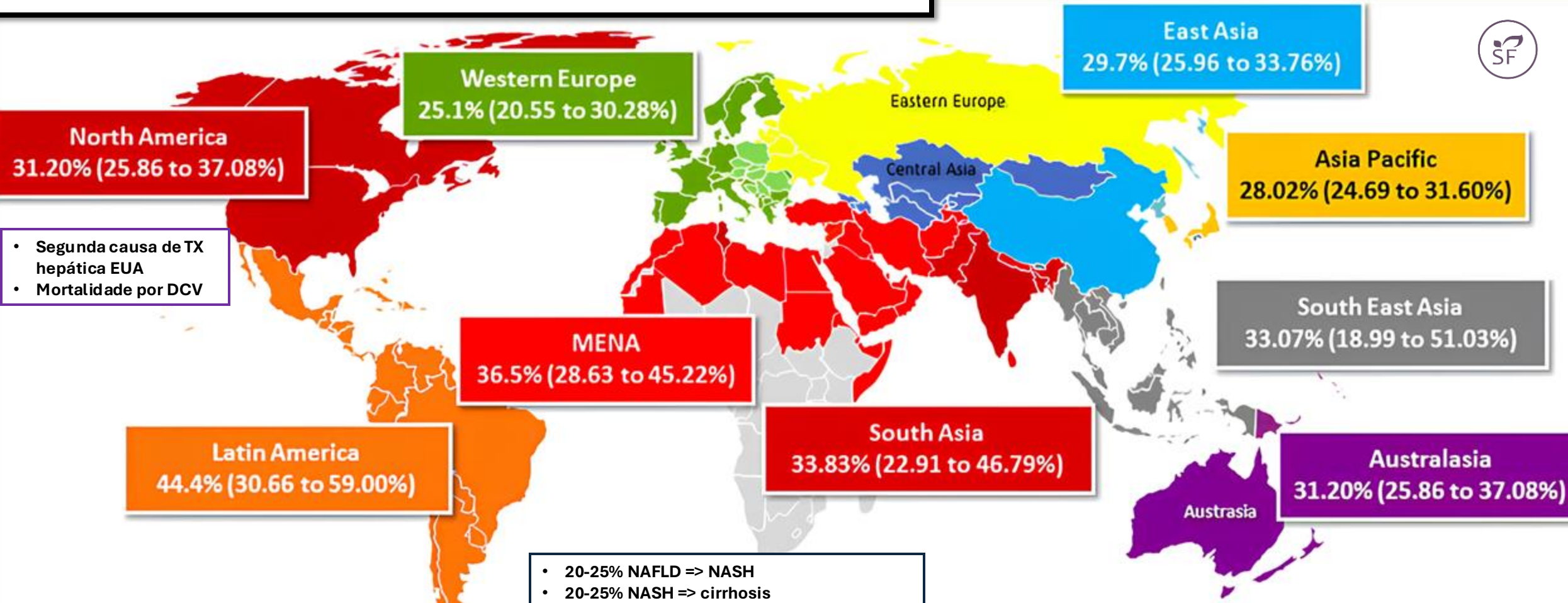


CRP
inflammation

OPEN

The global epidemiology of nonalcoholic fatty liver disease (NAFLD) and nonalcoholic steatohepatitis (NASH): a systematic review

PREVALÊNCIA

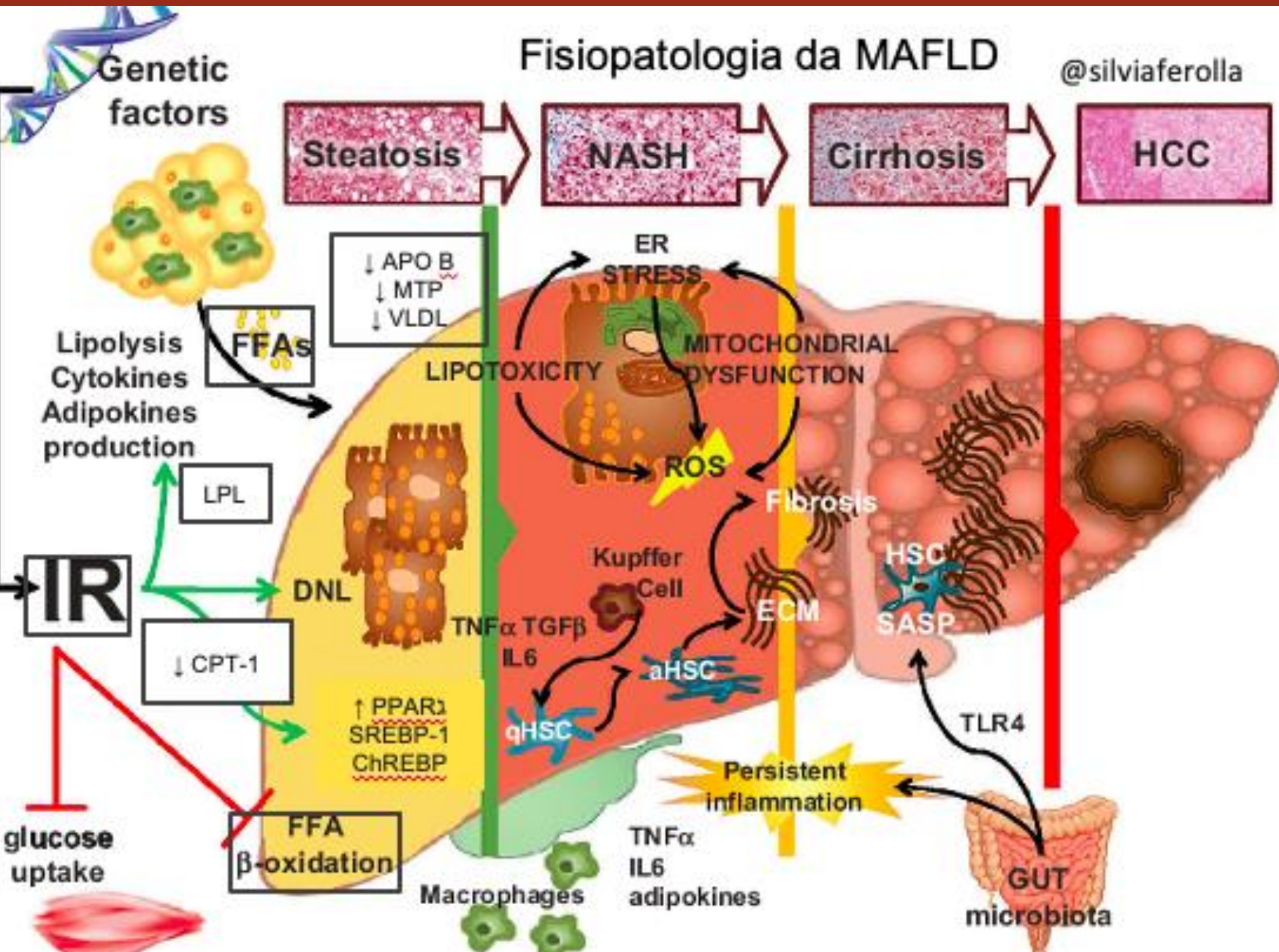


- Segunda causa de TX hepática EUA
- Mortalidade por DCV

- 20-25% NAFLD => NASH
- 20-25% NASH => cirrhosis
- 2,4 -12,8% => HCC risk (>7 years)

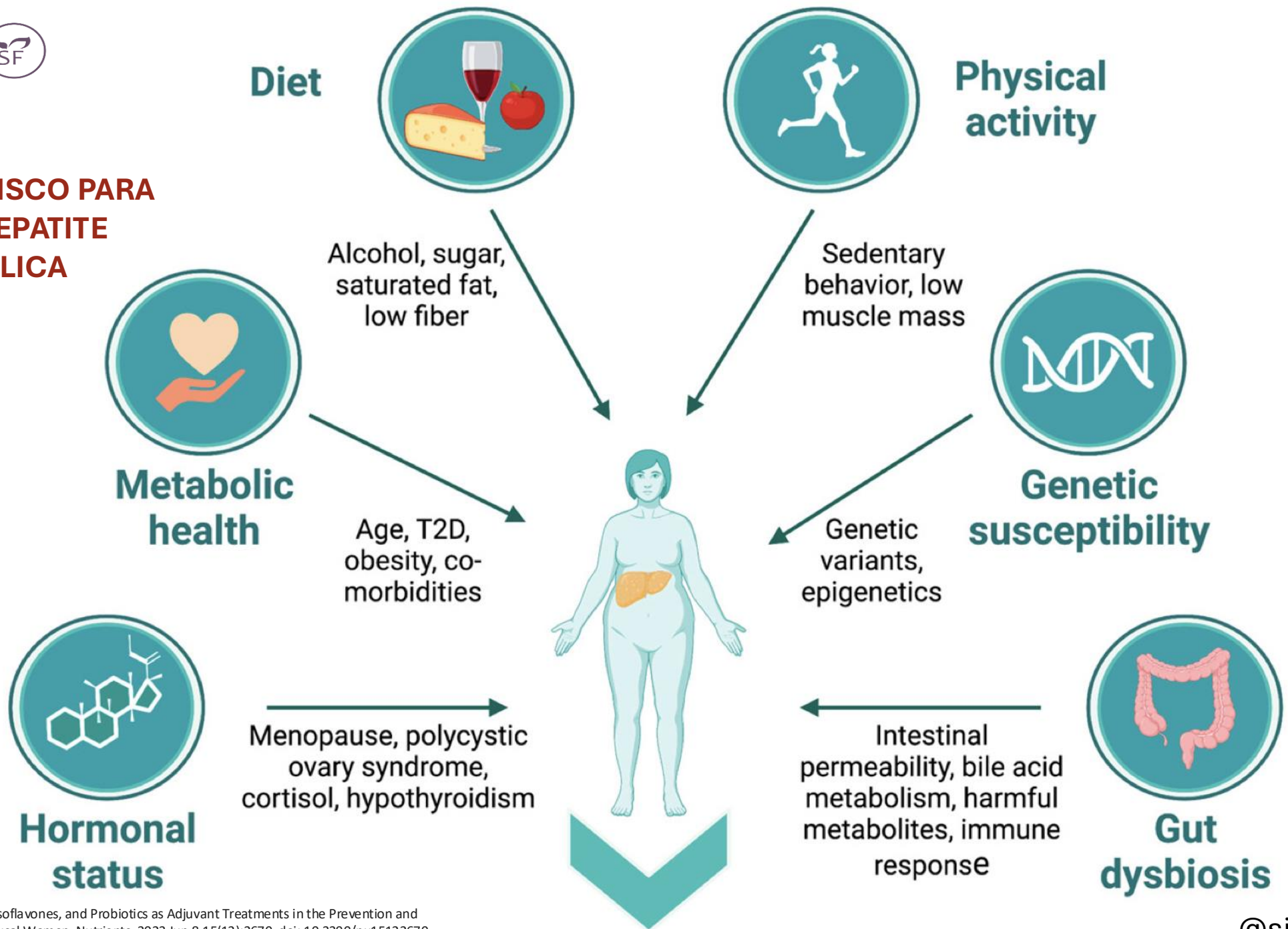
Fisiopatologia da MAFLD

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FATORES DE RISCO PARA ESTEATO-HEPATITE METABÓLICA



EIXO INTESTINO-FÍGADO

OPEN ACCESS
nutrients
ISSN 2072-6643
www.mdpi.com/journal/nutrients

Review

doi: 10.3390/nu6125583.

The Role of Intestinal Bacteria Overgrowth in Obesity-Related Nonalcoholic Fatty Liver Disease

Silvia M. Ferolla *, Geyza N. A. Armiliato †, Cláudia A. Couto † and Teresa C. A. Ferrari *

 nutrients



DOI: 10.3390/nu8070397



Article

Beneficial Effect of Synbiotic Supplementation on Hepatic Steatosis and Anthropometric Parameters, But Not on Gut Permeability in a Population with Nonalcoholic Steatohepatitis

Silvia M. Ferolla ^{1,*}, Cláudia A. Couto ¹, Luciana Costa-Silva ², Geyza N. A. Armiliato ¹, ~~Cláudia A. Couto~~, Flaviano S. Martins ³, Maria de Lourdes A. Ferrari ¹, Eduardo G. Vilela ¹, Henrique O. G. Torres ¹, Aloísio S. Cunha ¹ and Teresa C. A. Ferrari ^{1,*}

JOEL FAINTUCH
(Editor)
**Microbioma,
disbiose,
probióticos e
bacterioterapia**

<https://doi.org/10.1590/1516-3180.2021.0015.R1.14062021>

ORIGINAL ARTICLE

High prevalence of functional dyspepsia in nonalcoholic fatty liver disease: a cross-sectional study

Érika Cristina Lima^I, Maria do Carmo Friche Passos^{II}, Sílvia Marinho Ferolla^{III}, Raissa Soares Neves da Costa^{IV}, Quelson Coelho Lisboa^V, Lucas Ismael Dias Pereira^{VI}, Mateus Jorge Nogueira^{VII}, ~~Cláudia A. Couto~~, Claudia Alves Couto^X

W J H World Journal of
Hepatology

Submit a Manuscript: <http://www.wjgnet.com/esps/>
Help Desk: <http://www.wjgnet.com/esps/helpdesk.aspx>
DOI: 10.4254/wjh.v7.i3.559

World J Hepatol 2015 March 27; 7(3): 559-565

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MINIREVIEW

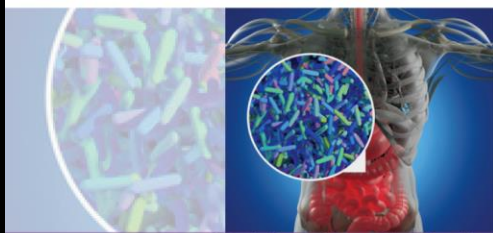
Probiotics as a complementary therapeutic approach in nonalcoholic fatty liver disease

doi: 10.4254/wjh.v7.i3.559

Silvia Marinho Ferolla, Geyza Nogueira de Almeida Armiliato, Cláudia Alves Couto, Teresa Cristina Abreu Ferrari

Silvia Marinho Ferolla

Microbiota intestinal e doença gordurosa hepática



Intestinal microbiota and fatty liver disease

W J H World Journal of
Hepatology

Submit a Manuscript: <http://www.wjgnet.com/esps/>
Help Desk: <http://www.wjgnet.com/esps/helpdesk.aspx>
DOI: 10.4254/wjh.v7.i24.2522

World J Hepatol 2015 October 28; 7(24): 2522-2534

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REVIEW

Dietary approach in the treatment of nonalcoholic fatty liver disease

Silvia Marinho Ferolla, Luciana Costa Silva, Maria de Lourdes Abreu Ferrari, Aloísio Sales da Cunha, Flaviano ~~Cláudia A. Couto~~, Teresa Cristina Abreu Ferrari

Centro de
**Nutrição
Funcional**



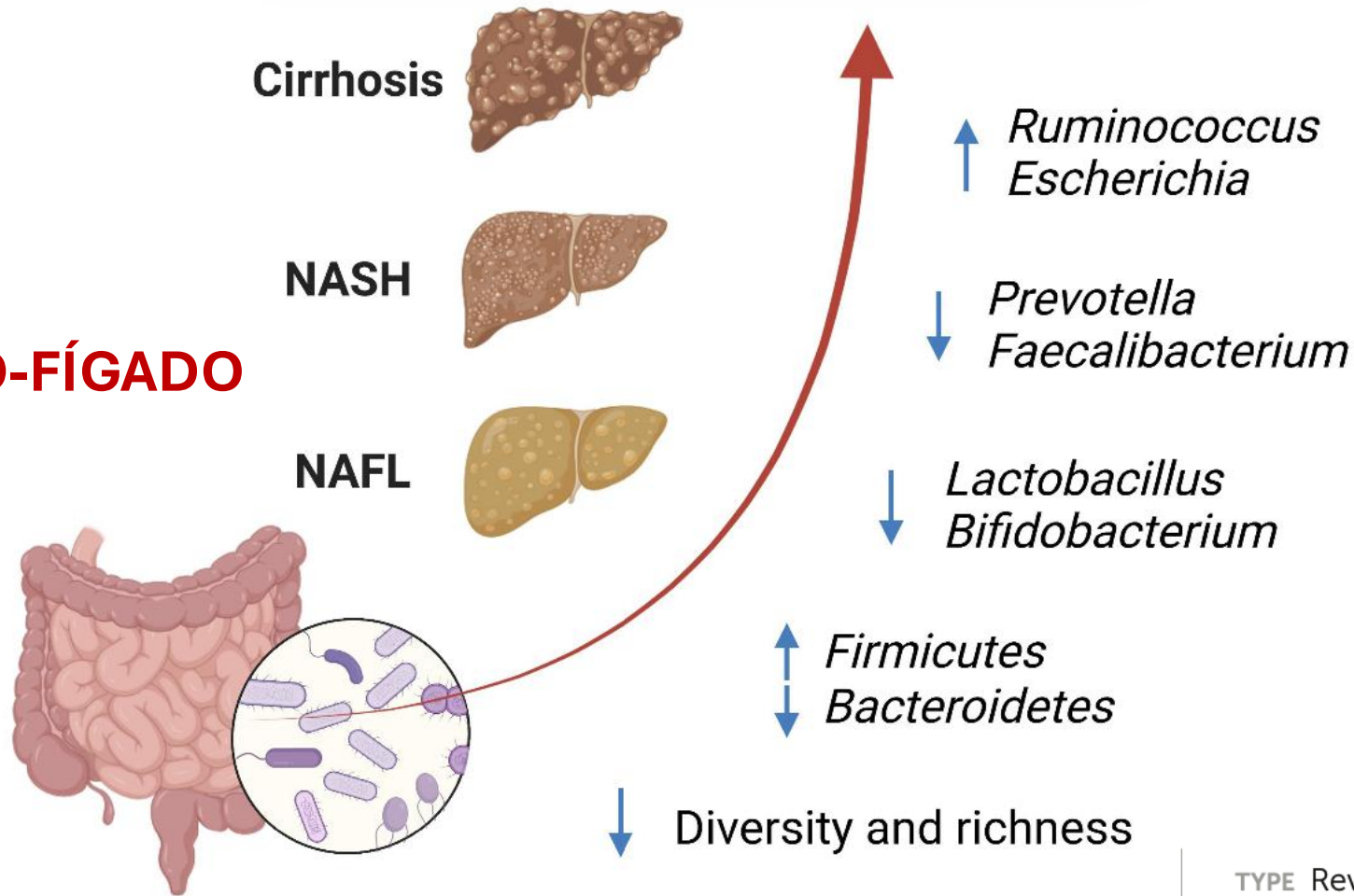
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Gut microbiota modulation in patients with non-alcoholic fatty liver disease: Effects of current treatments and future strategies

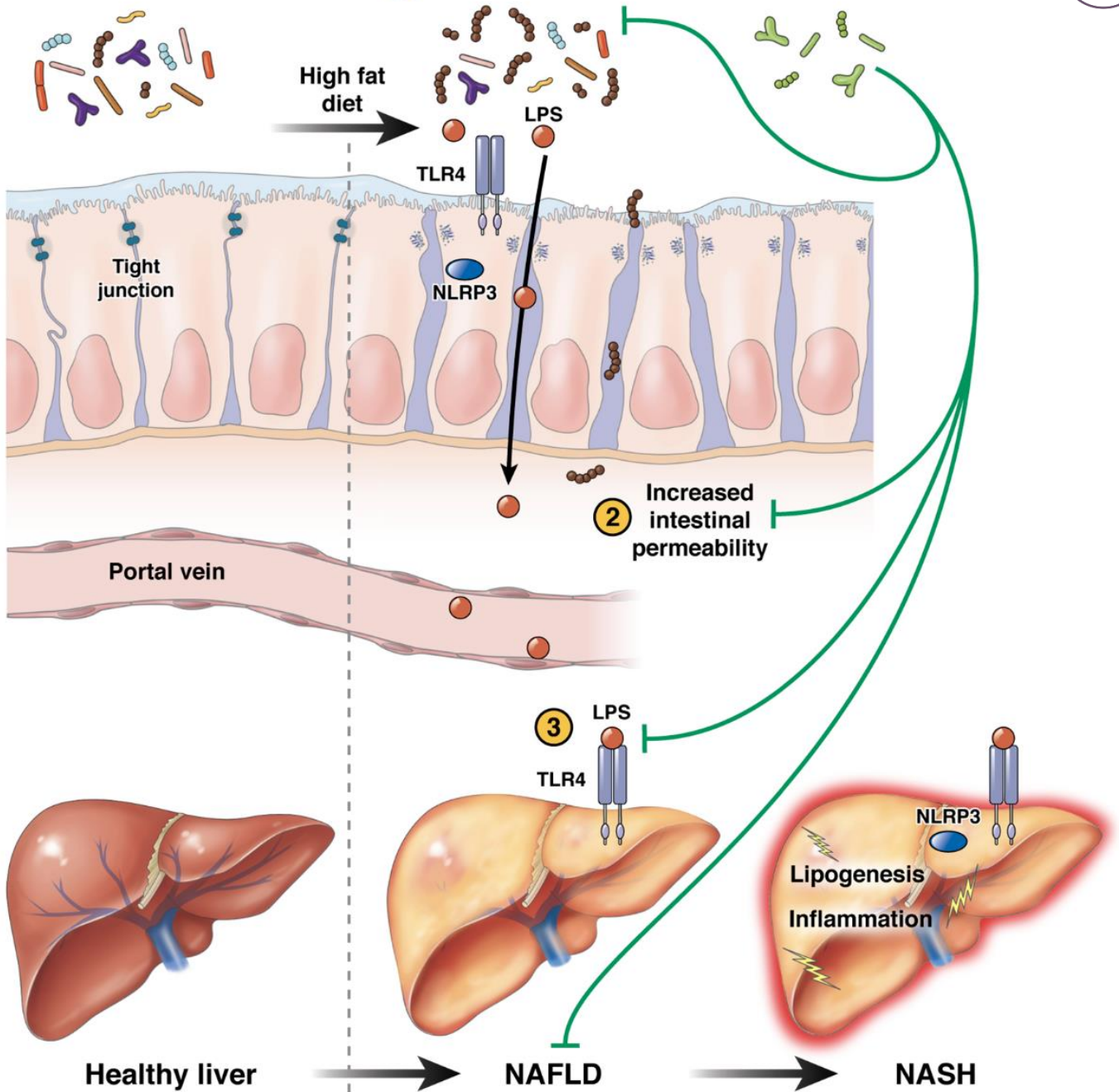
EIXO INTESTINO-FÍGADO



Gut microbiota

1 Gut dysbiosis

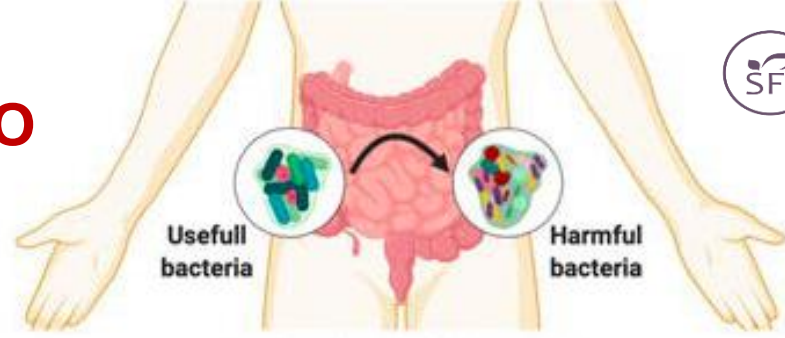
Probiotics



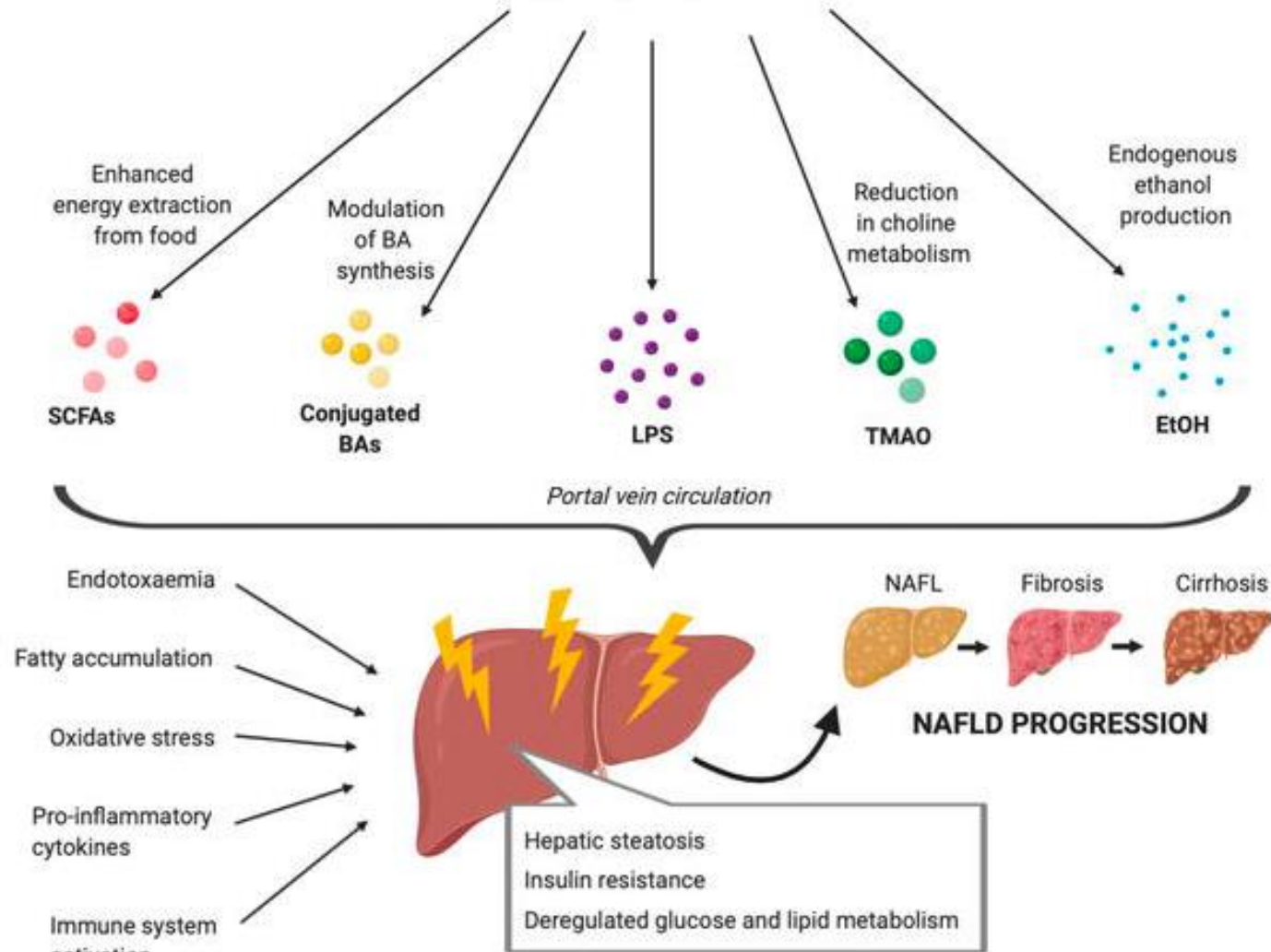
EIXO INTESTINO-FÍGADO

Aron-Wisniewsky J, Wambrunn MV, Nieuwdorp M, Clément K. Nonalcoholic Fatty Liver Disease: Modulating Gut Microbiota to Improve Severity? *Gastroenterology*. 2020 May;158(7):1881-1898. doi: 10.1053/j.gastro.2020.01.049. Epub 2020 Feb 8. PMID: 32044317.

EIXO INTESTINO-FÍGADO



INTESTINAL DYSBIOSIS



Silvia Ferolla



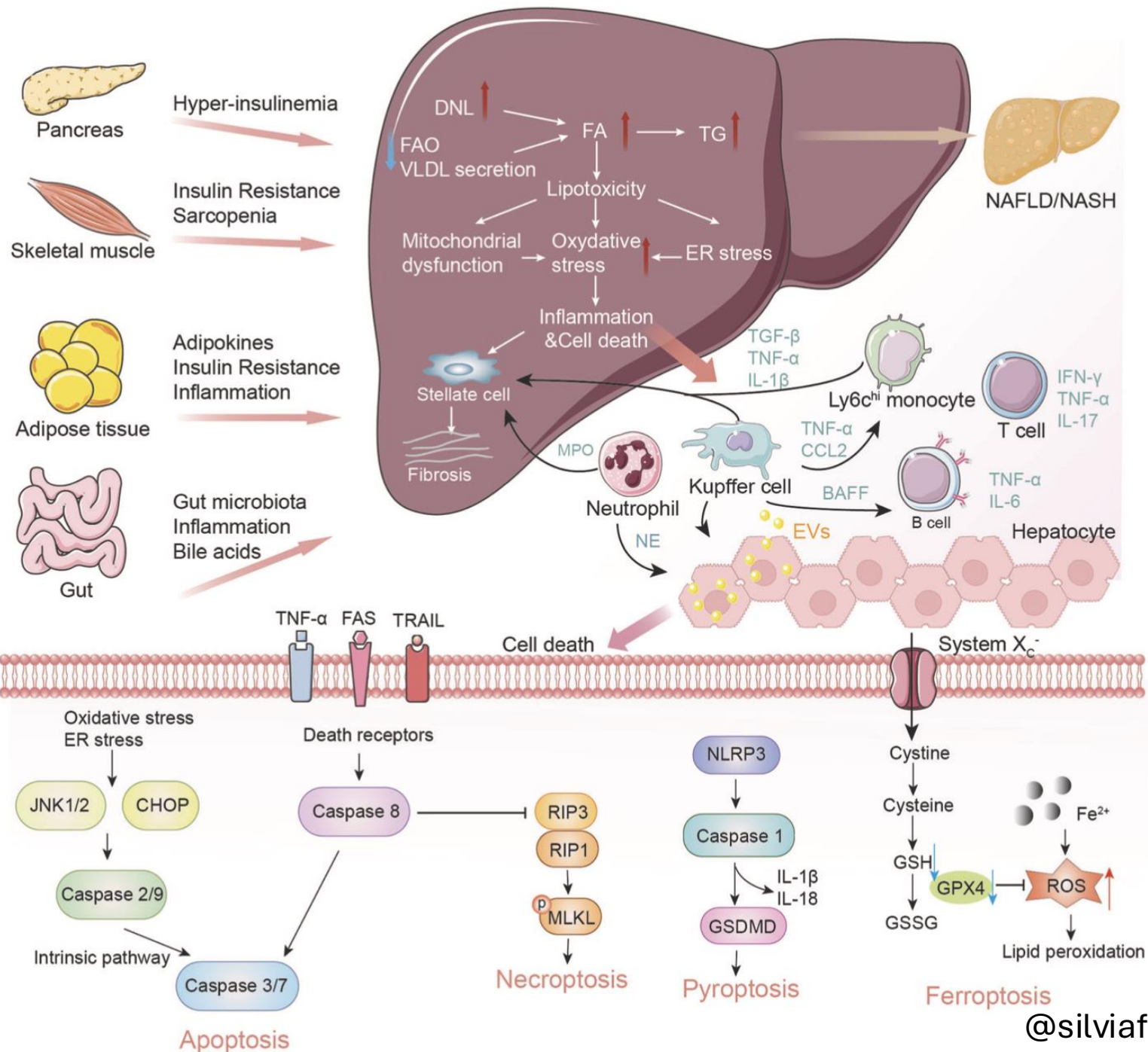
ESTEATO-HEPATITE METABÓLICA:

como reduzir inflamação com a
nutrição integrativa ?

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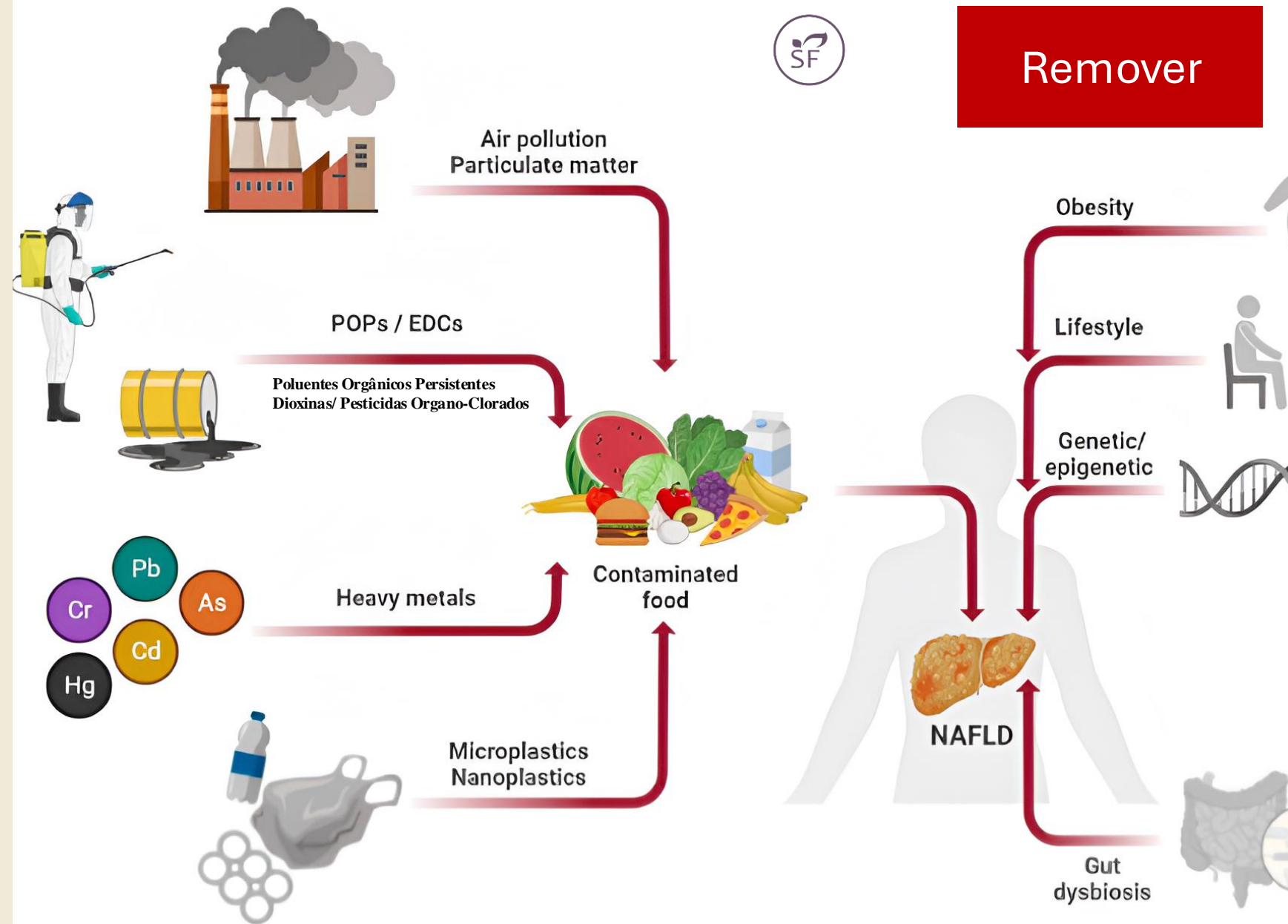


INFLAMAÇÃO NA ESTEATO-HEPATITE METABÓLICA



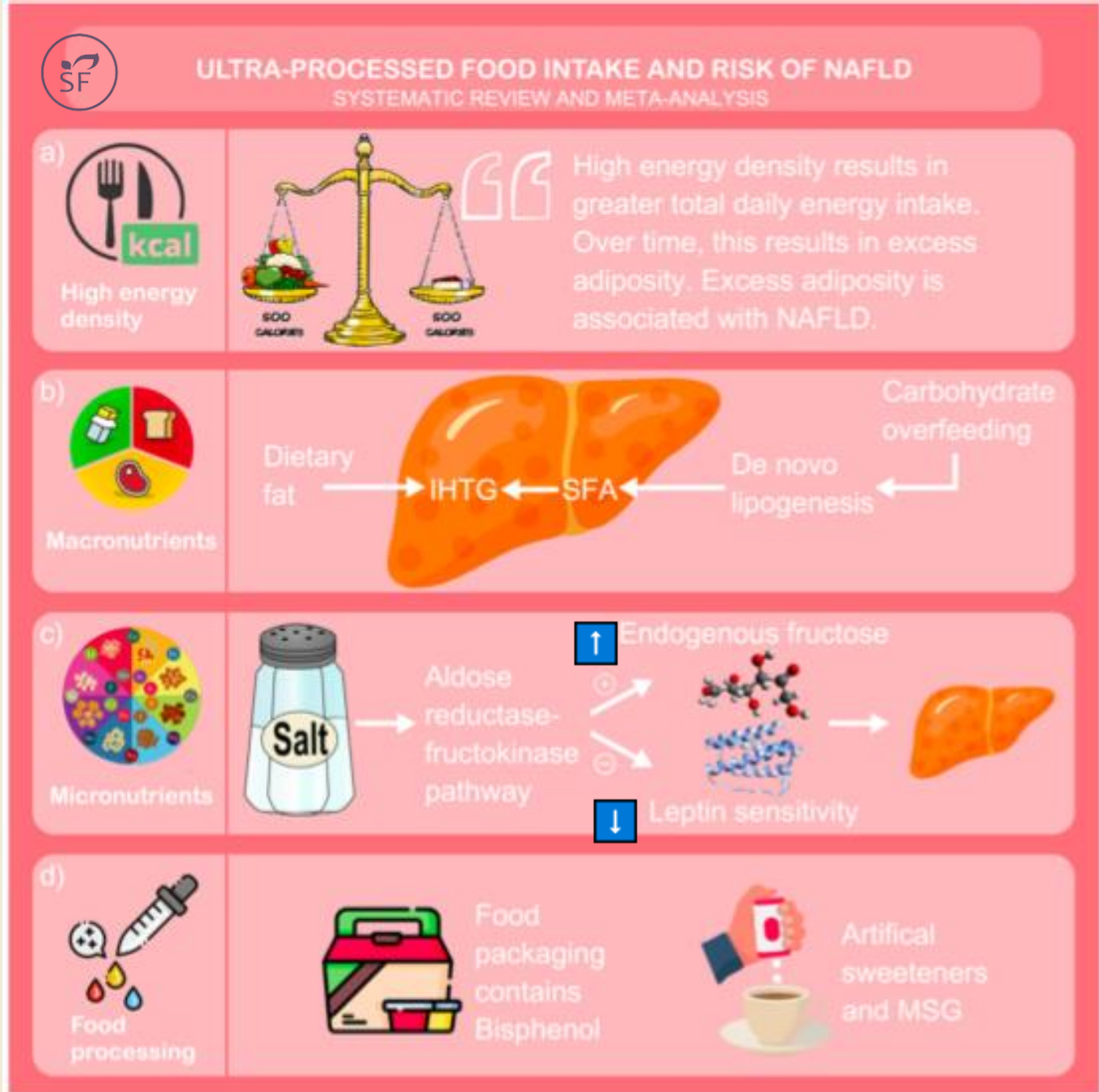
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A ingestão de poluentes ambientais, incluindo poluentes orgânicos persistentes (POPs), produtos químicos desreguladores endócrinos (EDCs), metais pesados, micro e nanoplásticos, promove o desenvolvimento e progressão da DHGM



ULTRAPROCESSADOS E RISCO DE ESTEATO-HEPATITE METABÓLICA

Remover



American Society for Nutrition
Excellence in Nutrition Research and PracticeThe American Journal of
CLINICAL NUTRITIONjournal homepage: <https://ajcn.nutrition.org/>

Original Research Article

Higher ultra-processed food intake is associated with adverse liver outcomes: a prospective cohort study of UK Biobank participants

Longgang Zhao^{1,2}, Alyssa Clay-Gilmour¹, Jiajia Zhang¹, Xuehong Zhang^{2,3,†}, Susan E. Steck^{1,*,†}¹ Department of Epidemiology and Biostatistics, Arnold School of Public Health, University of South Carolina, Columbia, SC, United States;² Department of Medicine, Brigham and Women's Hospital and Harvard Medical School, Boston, MA, United States; ³ Department of Nutrition, Harvard T.H. Chan School of Public Health, Boston, MA, United States

ABSTRACT

Background: Ultra-processed food (UPF) intake has been positively associated with obesity and diabetes. The relationship between UPF intake and liver health has been scarcely studied.

Objectives: We aimed to evaluate the association of UPF intake with risk of adverse liver outcomes including nonalcoholic fatty liver disease (NAFLD), liver fibrosis/cirrhosis, liver cancer, severe liver disease, and serum biomarkers of liver health.

Methods: A total of 173,889 participants aged 40 to 69 y from the UK Biobank were included. UPF intake was defined using 24-h dietary recalls and NOVA classification. Liver outcome data were obtained from cancer registry, in-hospital records, and death registries. Serum biomarkers were measured at baseline. We used Cox proportional hazards models to estimate hazard ratios (HRs) and 95% confidence intervals (CIs) for associations between UPF and adverse liver outcomes adjusting for demographics, lifestyle factors, body mass index, and diabetes. We used multinomial logistic regression to evaluate associations between UPF and liver function biomarkers.

Results: After a median follow-up of 8.9 y, we documented 1108 NAFLD, 350 liver fibrosis/cirrhosis, 134 liver cancer, and 550 severe liver disease cases. Higher UPF intake was associated with increased risk of NAFLD (HR_{Quartile 4 vs. Quartile 1}: 1.43; 95% CI: 1.21, 1.70; $P_{\text{trend}} < 0.001$), liver fibrosis/cirrhosis (HR: 1.18; 95% CI: 0.87, 1.59; $P_{\text{trend}} = 0.009$), and severe liver disease (HR: 1.50; 95% CI: 1.19, 1.90; $P_{\text{trend}} < 0.001$) but not with liver cancer (HR: 1.00; 95% CI: 0.63, 1.58; $P_{\text{trend}} = 0.88$). Higher UPF intake was associated with elevated levels of C-reactive protein, alkaline phosphatase, aspartate aminotransferase, γ -glutamyltransferase, and triglycerides and lower cholesterols (all $P_{\text{trend}} < 0.001$).



Conclusions: Higher UPF intake is associated with an increased risk of NAFLD, liver fibrosis and cirrhosis, and severe liver disease and adverse levels of multiple clinical biomarkers, suggesting the potential importance of reducing UPF intake to improve liver health.

Keywords: liver health, cirrhosis, liver cancer, NAFLD, ultra-processed food, prospective, cohort, biomarker

Duração do Estudo: 8,9**Casos Observados:**

- 1108 casos de DHGNA
- 350 casos de fibrose/cirrose hepática
- 134 casos de câncer de fígado
- 550 casos de doença hepática avançada

• Associações de risco > consumo UP e:

- **Esteatose:**  43% no risco (quartil mais alto vs. mais baixo)
- **Doença Hepática Avançada:**  50% no risco

Marcadores Inflamatórios:

- níveis elevados de: PCR, FA, AST, GGT, TG

Perspective

“Sweet death”: Fructose as a metabolic toxin that targets the gut-liver axis

Mark A. Febbraio^{1,*} and Michael Karin^{2,*}

¹Monash Institute of Pharmaceutical Sciences, Monash University, Parkville, VIC, Australia

²Department of Pharmacology, School of Medicine, University of California, San Diego, San Diego, CA, USA

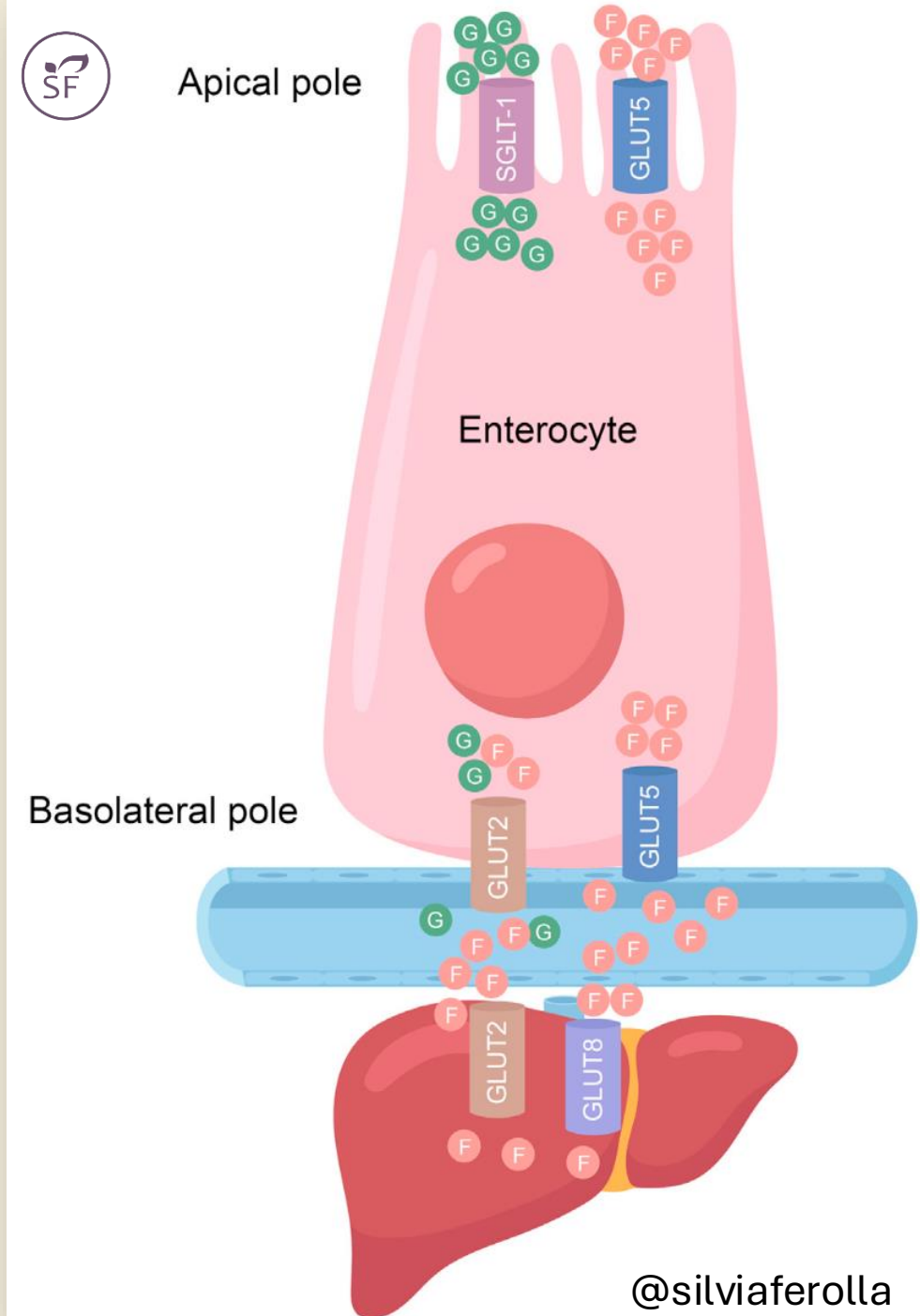
*Correspondence: mark.febbraio@monash.edu (M.A.F.), mkarin@health.ucsd.edu (M.K.)

<https://doi.org/10.1016/j.cmet.2021.09.004>

“Morte doce”: A frutose como uma toxina metabólica que afeta o eixo intestino-fígado

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FRUTOSE E ESTEATO-HEPATITE METABÓLICA



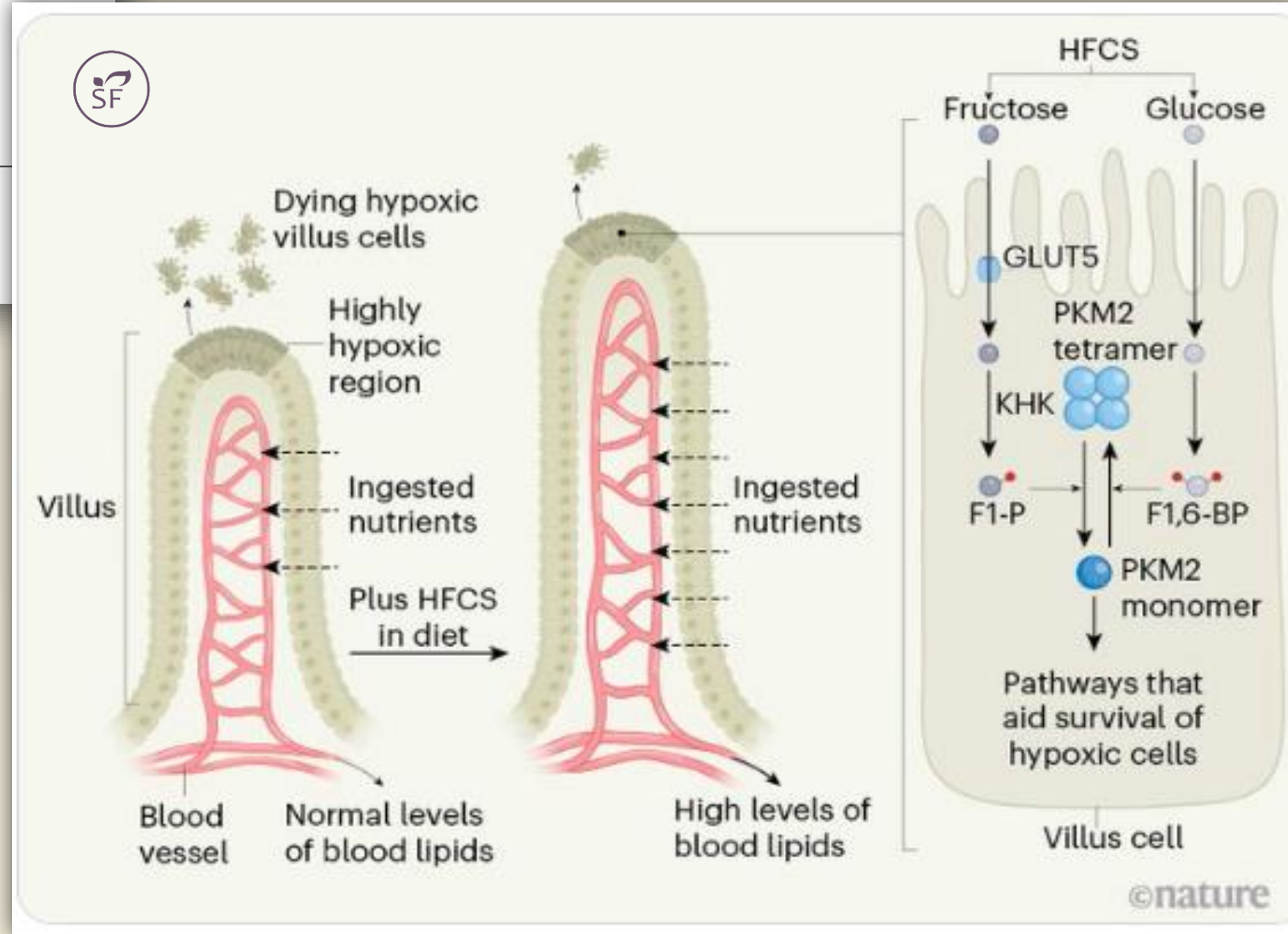
Dietary fructose expands the gut and aids fat uptake

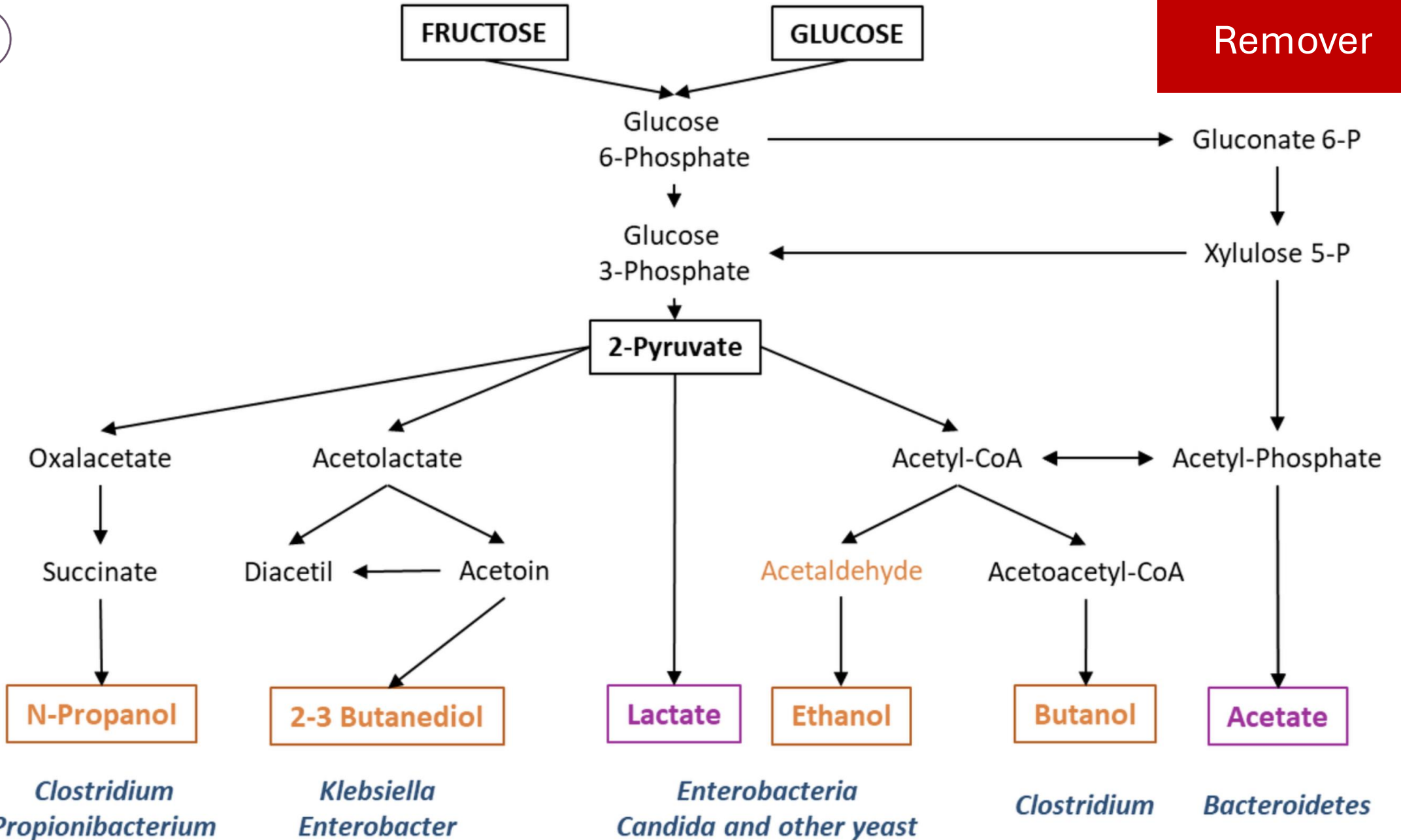
Patrícia M. Nunes & Dimitrios Anastasiou

Feeding mice high-fructose corn syrup, a widely used sweetener in human diets, has been found to drive an increase in the surface area of the gut that is associated with enhanced absorption of dietary nutrients and weight gain. See p.263

nature

Remove





Remover

Perspective

“Sweet death”: Fructose as a metabolic toxin that targets the gut-liver axis

Mark A. Febbraio^{1,*} and Michael Karin^{2,*}

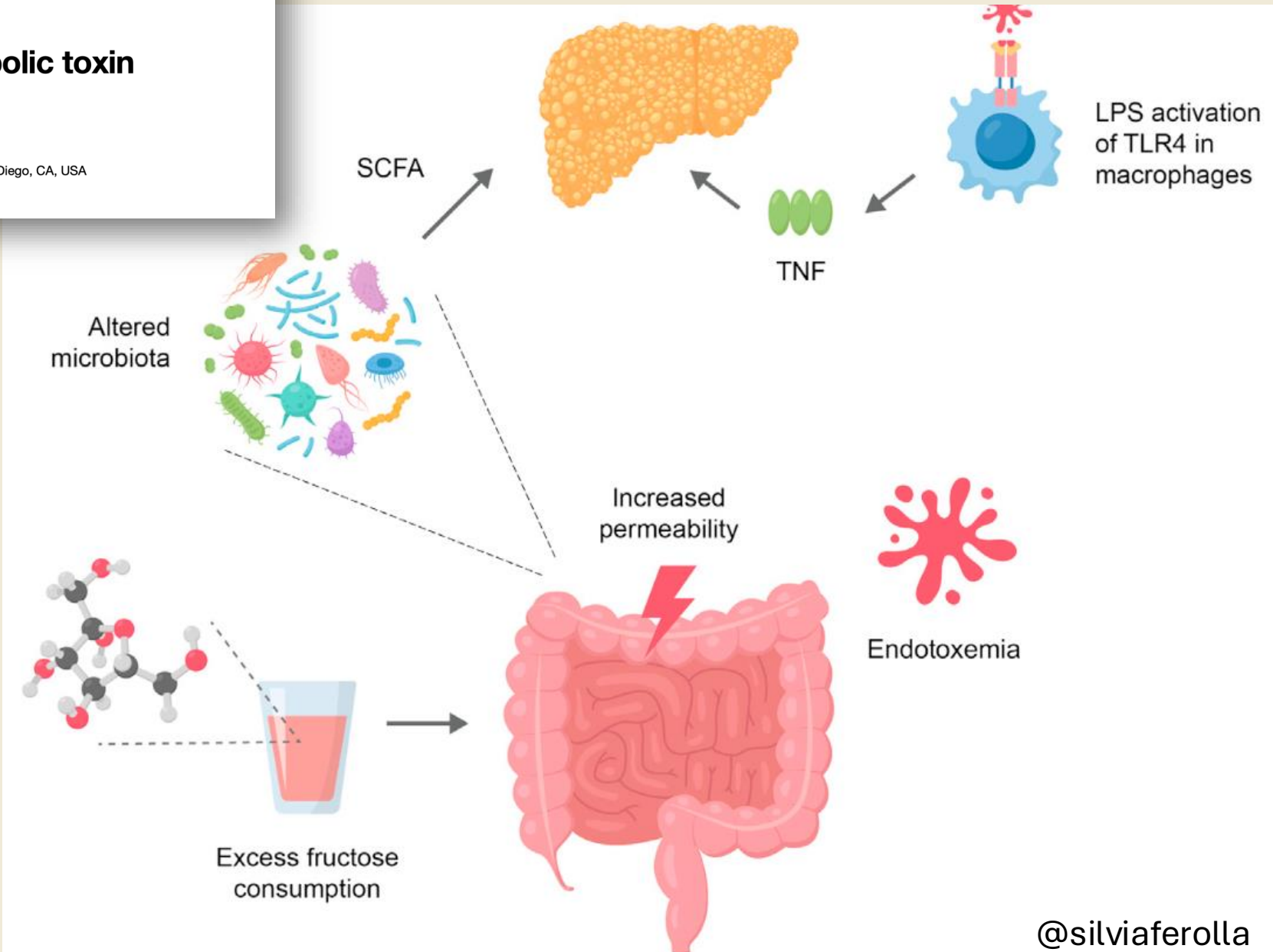
¹Monash Institute of Pharmaceutical Sciences, Monash University, Parkville, VIC, Australia

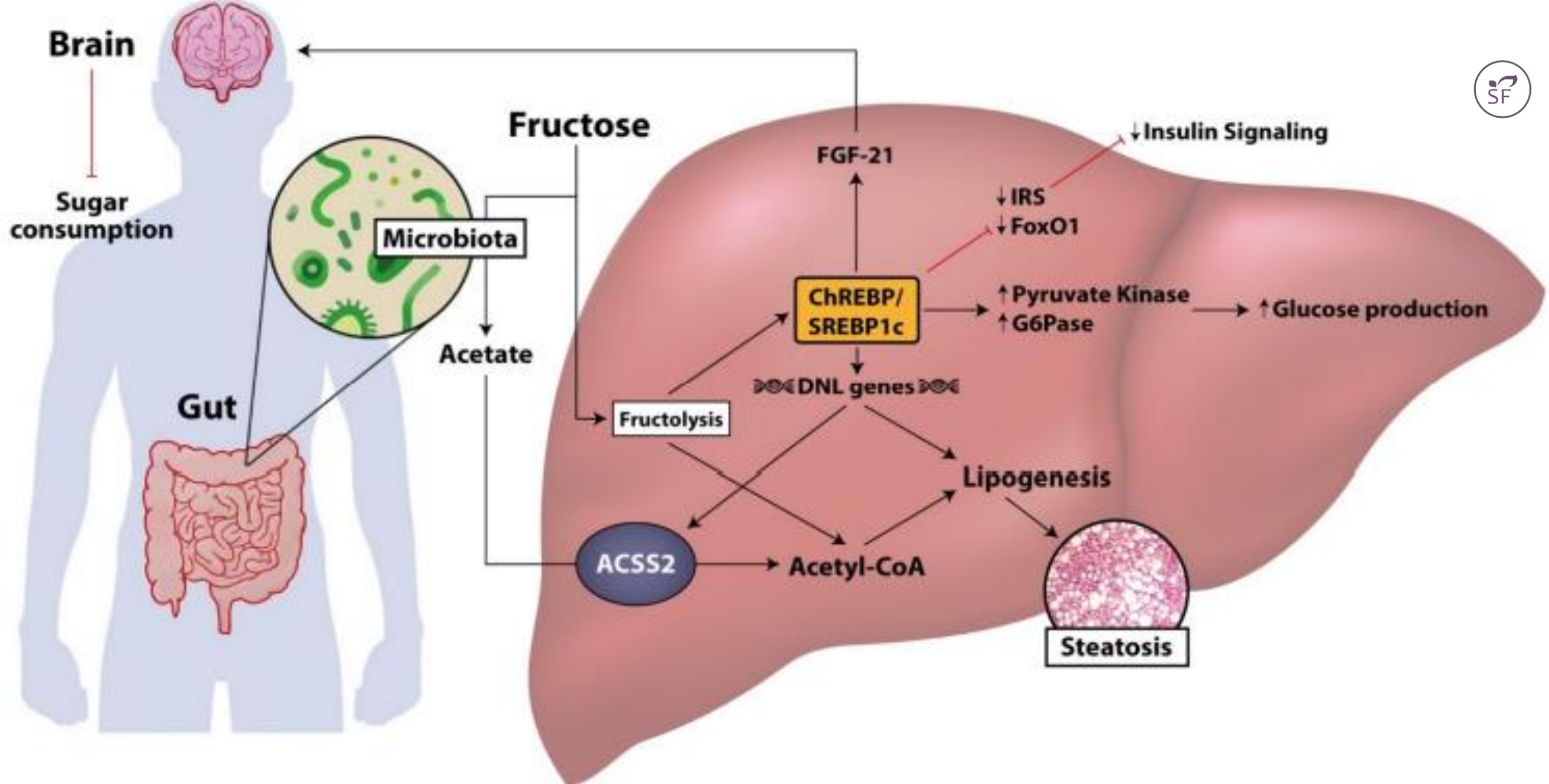
²Department of Pharmacology, School of Medicine, University of California, San Diego, San Diego, CA, USA

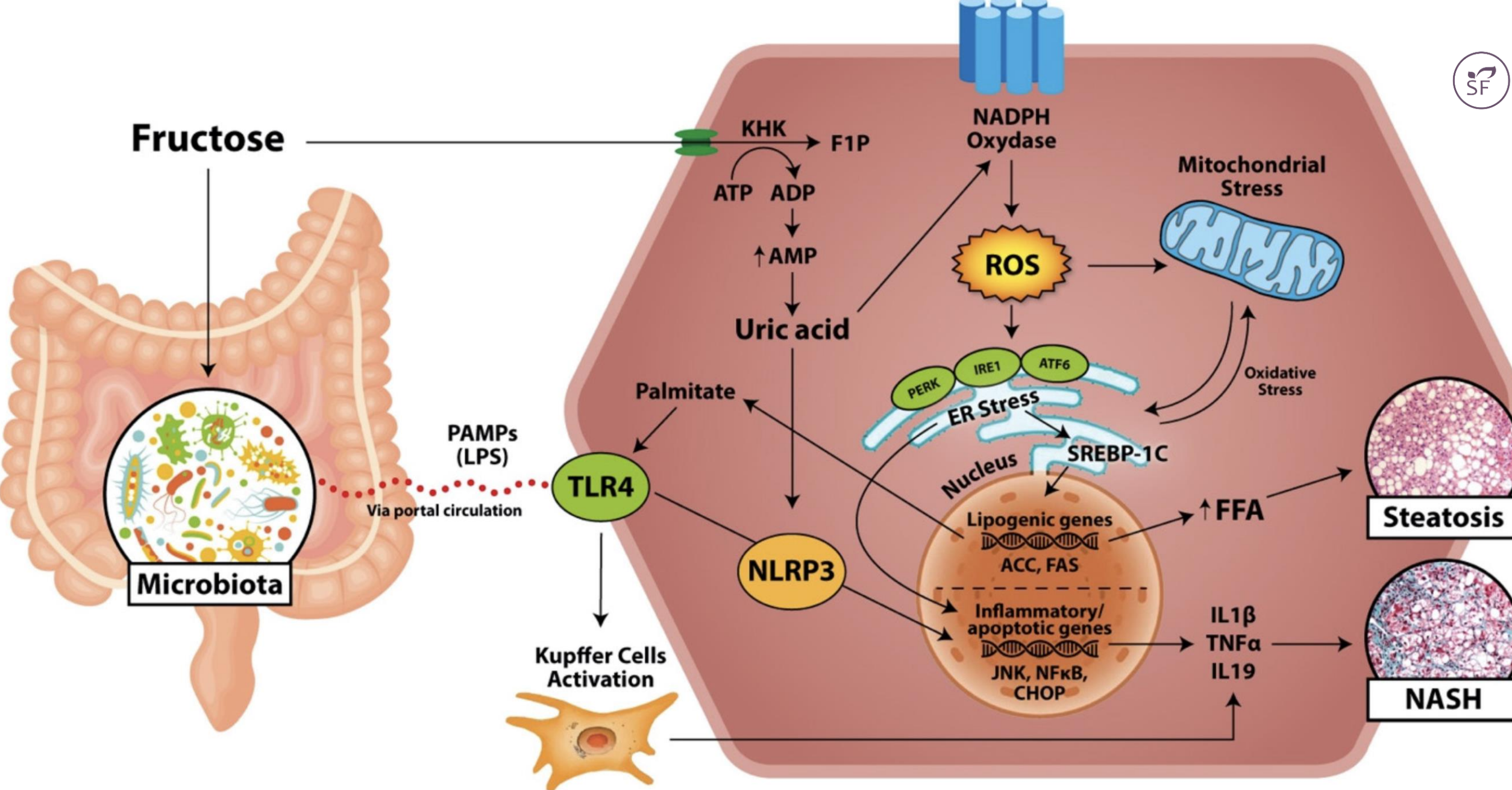
*Correspondence: mark.febrario@monash.edu (M.A.F.), mkarin@health.ucsd.edu (M.K.)

<https://doi.org/10.1016/j.cmet.2021.09.004>

Remover







Remover



Article

Consumption of Sugar-Sweetened Beverages Has a Dose-Dependent Effect on the Risk of Non-Alcoholic Fatty Liver Disease: An Updated Systematic Review and Dose-Response Meta-Analysis

Abstract: *Background:* Non-alcoholic fatty liver disease (NAFLD) is a serious health problem, but the dose-response relationship between sugar-sweetened beverages (SSBs) and NAFLD remains uncertain. *Methods:* A systematic review and dose-response meta-analysis were conducted following the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) guidelines. Review Manager 5.3 and Stata 14.0 were used to combine trials and analyze data. The dose-response meta-analysis was performed by non-linear trend regression. *Results:* Twelve studies recruiting a total of 35,705 participants were included. The results showed that the consumption of SSBs was associated with 1.39-fold increased odds of NAFLD (95% CI: 1.29–1.50, $p < 0.00001$). The risk of NAFLD rose with an increased consumption of SSBs, while the consumptions of low doses (<1 cup/week), middle dose (1–6 cups/week) and high doses (≥ 7 cups/week) of SSBs increased the relative risk of NAFLD by 14%, 26% and 53%, respectively ($p = 0.01$, $p < 0.00001$, $p = 0.03$, respectively). *Conclusions:* This study demonstrates that consumers of SSBs are at significantly increased risk of NAFLD and the consumption of SSBs has a dose-dependent effect on the risk of NAFLD. The findings of this study strengthen the evidence base for healthy dietary patterns and are meaningful for the primary prevention of NAFLD.

Metanálise com amostra de 35.705

Dose “pequena” < 1 copo/ sem.
aumenta 14% RR de DHGM

Dose “moderada” 1 a 6 copos/ sem.
aumenta 26% RR de DHGM

Dose “alta” ≥ 7 copos/ sem. aumenta
53% RR de DHGM

JOURNAL ARTICLE

Associations of Macronutrients Intake With MRI-determined Hepatic Fat Content, Hepatic Fibroinflammation, and NAFLD

AGS associada ao **↑** da esteatose, fibroinflamação e prevalência da DHGM



AMOSTRA: 12.620

HEALTHY LIVER



FATTY LIVER



Maior ingestão de fibras associada a **↓** esteatose e fibroinflamação



Maior ingestão de MUFA associada a **↓** fibroinflamação



Maior ingestão de proteínas associada a **↓** esteatose e fibroinflamação

Remover

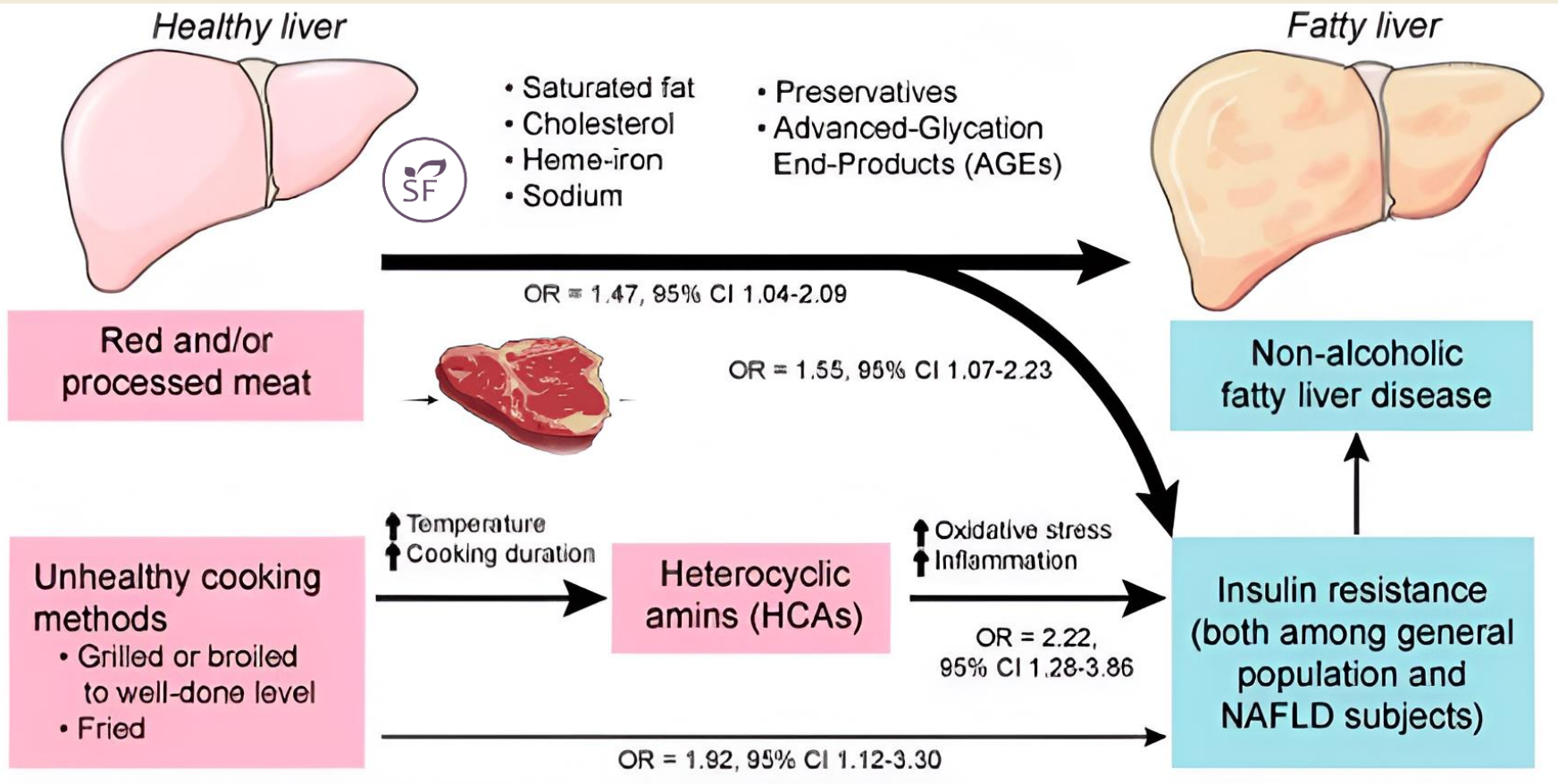
Maior ingestão de amido e açúcar associada a **↑** fibroinflamação

Reparar



High red and processed meat consumption is associated with non-alcoholic fatty liver disease and insulin resistance

Remove



Carne Total:

- ↑ 49% nas chances de DHGM (porções/dia acima da mediana)
- ↑ 63% nas chances de RI

Carne Vermelha e/ou

Processada:

- ↑ 47% nas chances de DHGNA
- ↑ 55% nas chances de RI

Métodos de Cozimento Não Saudáveis:

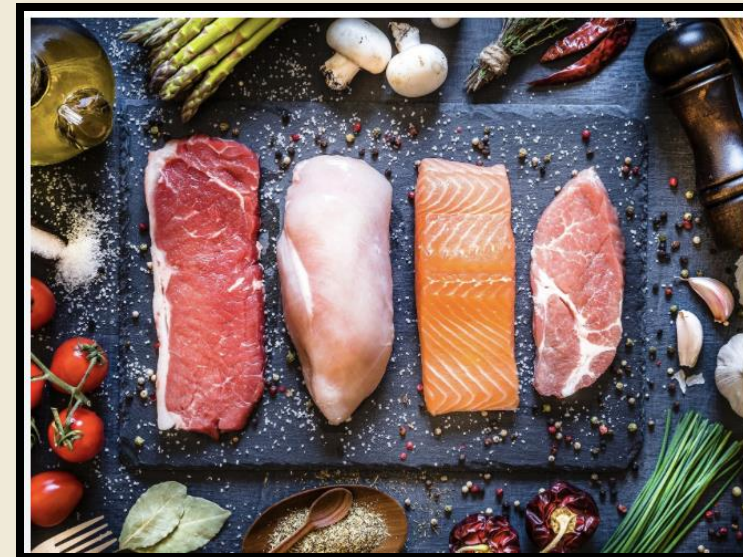
- ↑ 92% nas chances de RI

Aminas Heterocíclicas (HCAs):

- ↑ 122% nas chances de RI

Research Article

Differential Effects of Dietary White Meat and Red Meat on NAFLD Progression by Modulating Gut Microbiota and Metabolites in Rats



Reparar

Carne Branca (carpa e frango):

- A dieta baseada em carpa mostrou melhorias significativas nas alterações patológicas hepáticas e no metabolismo glicolipídico.
- A dieta baseada em frango mostrou melhorias parciais nos parâmetros metabólicos.
- Observou-se enriquecimento de bactérias benéficas como *Lactobacillus* ou *Akkermansia*, aumento de AGCC s e redução de bactérias patogênicas.

Remover

Carne Vermelha (porco e boi):

- Progressão da DHGNA foi observada nos grupos alimentados com carne de porco e boi.
- Encontradas mudanças na microbiota que incluíam o enriquecimento de bactérias patogênicas (como *Prevotella_9* ou *Lachnospiraceae_UCG-010*), e redução de AGCC

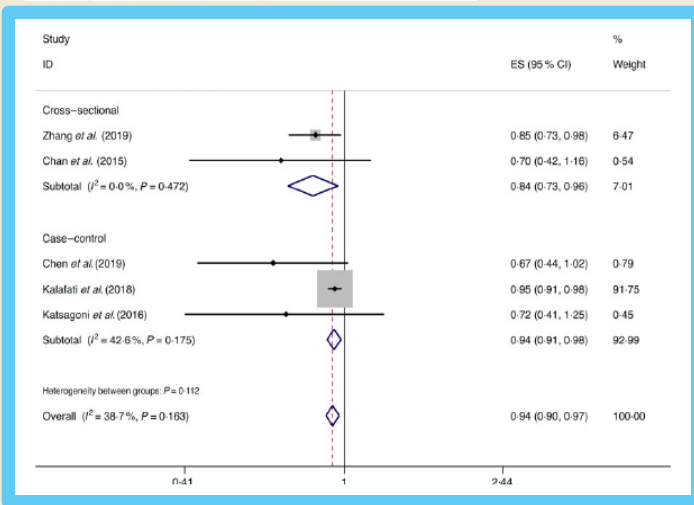
Food groups and the likelihood of non-alcoholic fatty liver disease: a systematic review and meta-analysis

Reparar

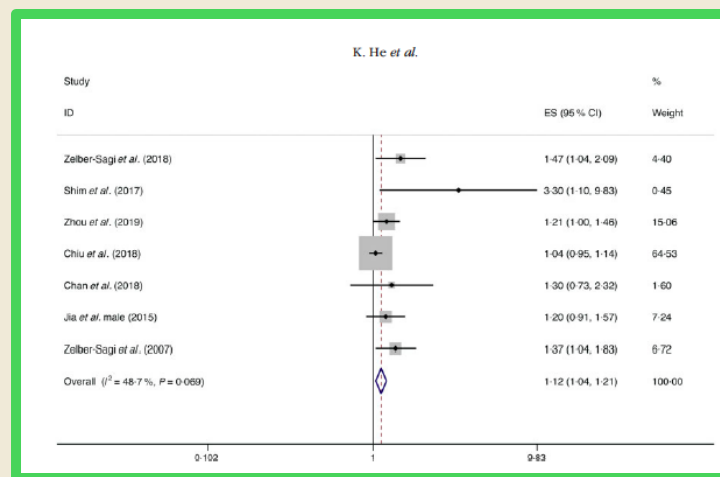
Remover



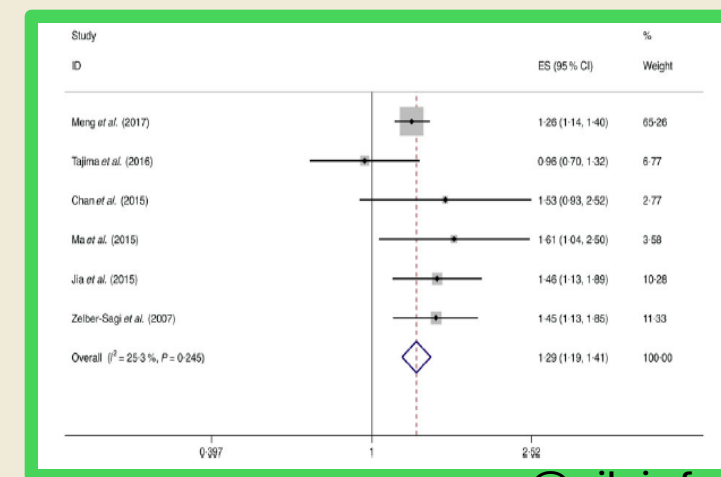
Consumo de oleaginosas e o risco de DHGNA



Consumo de carne vermelha e o risco de DHGNA



Consumo de refrigerante e o risco de DHGNA



Guidance statements:

23. There are currently no FDA-approved medications for the treatment of NAFLD, but drugs approved to treat associated comorbidities with potential benefit in NAFLD may be considered in the appropriate clinical setting.

24. Semaglutide can be considered for its approved indications (T2DM/obesity) in patients with NASH, as it confers a cardiovascular benefit and improves NASH.

25. Pioglitazone improves NASH and can be considered for patients with NASH in the context of patients with T2DM .

26. Vitamin E can be considered in select individuals as it improves NASH in some patients without diabetes.

27. Available data on semaglutide, pioglitazone, and vitamin E do not demonstrate an antifibrotic benefit, and none has been carefully studied in patients with cirrhosis.

HEPATOLOGY

Received: 18 January 2023 | Accepted: 18 January 2023

DOI: 10.1097/HEP.0000000000000323



PRACTICE GUIDANCE

AASLD Practice Guidance on the clinical assessment and management of nonalcoholic fatty liver disease

Mary E. Rinella¹ | Brent A. Neuschwander-Tetri² |
Mohammad Shadab Siddiqui³ | Manal F. Abdelmalek⁴ | Stephen Caldwell⁵ |
Diana Barb⁶ | David E. Kleiner⁷ | Rohit Loomba⁸



Reparar

Rinella ME, Neuschwander-Tetri BA, Siddiqui MS, Abdelmalek MF, Caldwell S, Barb D, Kleiner DE, Loomba R. AASLD Practice Guidance on the clinical assessment and management of nonalcoholic fatty liver disease. *Hepatology*. 2023 May 1;77(5):1797-1835. doi: 10.1097/HEP.0000000000000323. Epub 2023 Mar 17. PMID: 36727674; PMCID: PMC10735173.

@silviaferolla

Reparar



Castanha do pará



5,73mg



Nozes

27mg

Amêndoa



24mg

Avelã



23mg

ALIMENTOS FONTES DE VITAMINA E

Semente de girassol



35 mg

Azeite



14,42mg

Azeitona



20mg



Abacate

21mg

800 UI de vitamina E equivalem a 536 mg.

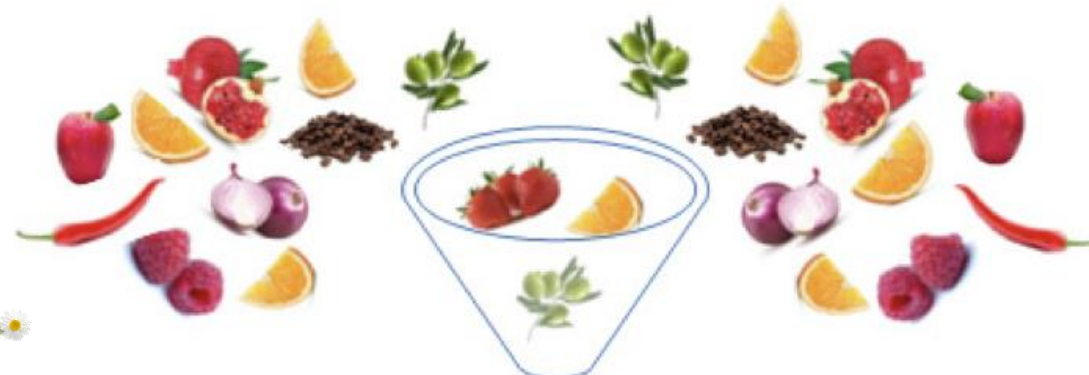
UL para adultos de vitamina E é de 1.000 mg por dia

RDA para vitamina E em adultos é de 15 mg por dia (ou 22,4 UI) de alfa-tocoferol

Reparar



Salsa, alecrim, hortelã,
tomilho, sálvia



Polyphenols



Lignans

Non-Flavonoids

Hydroxybenzoic acids

Hydroxycinnamic acids

Curcuminoids

Stilbenes

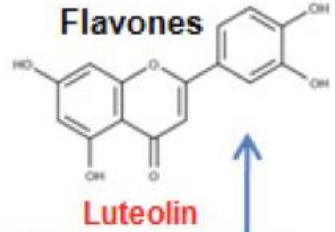
Resveratrol

Curcumin

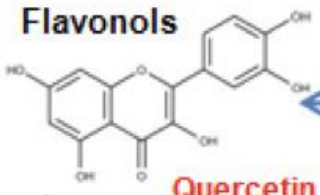
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Flavonoids

Flavones



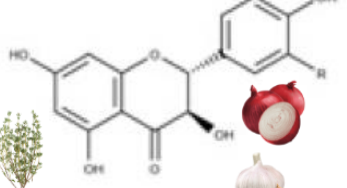
Flavonols



Flavanones

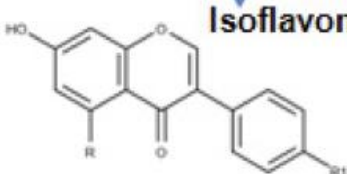
Hesperidina,
naringenina

Flavanonols



Anthocyanidins

Isoflavones



catechins

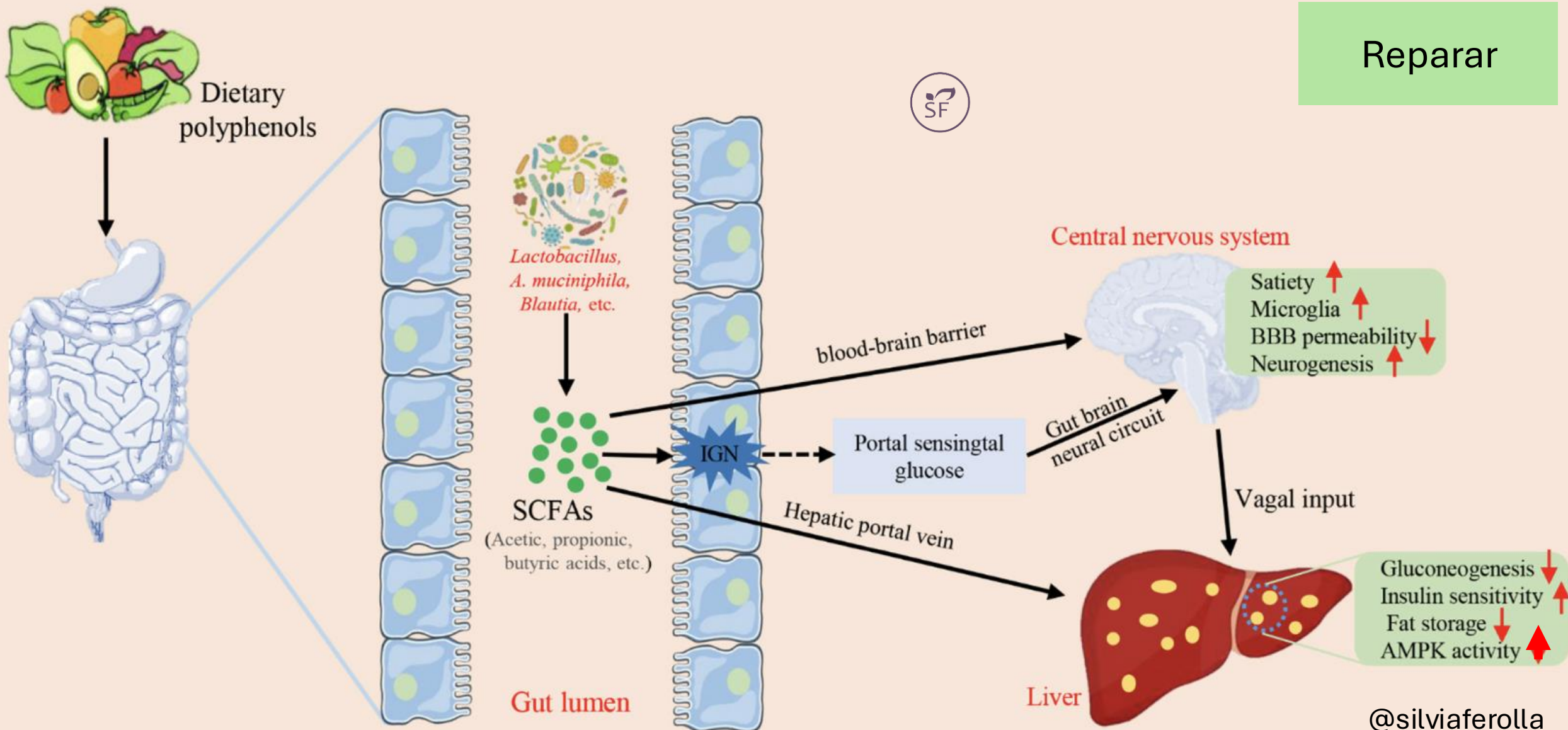


Flavandiols



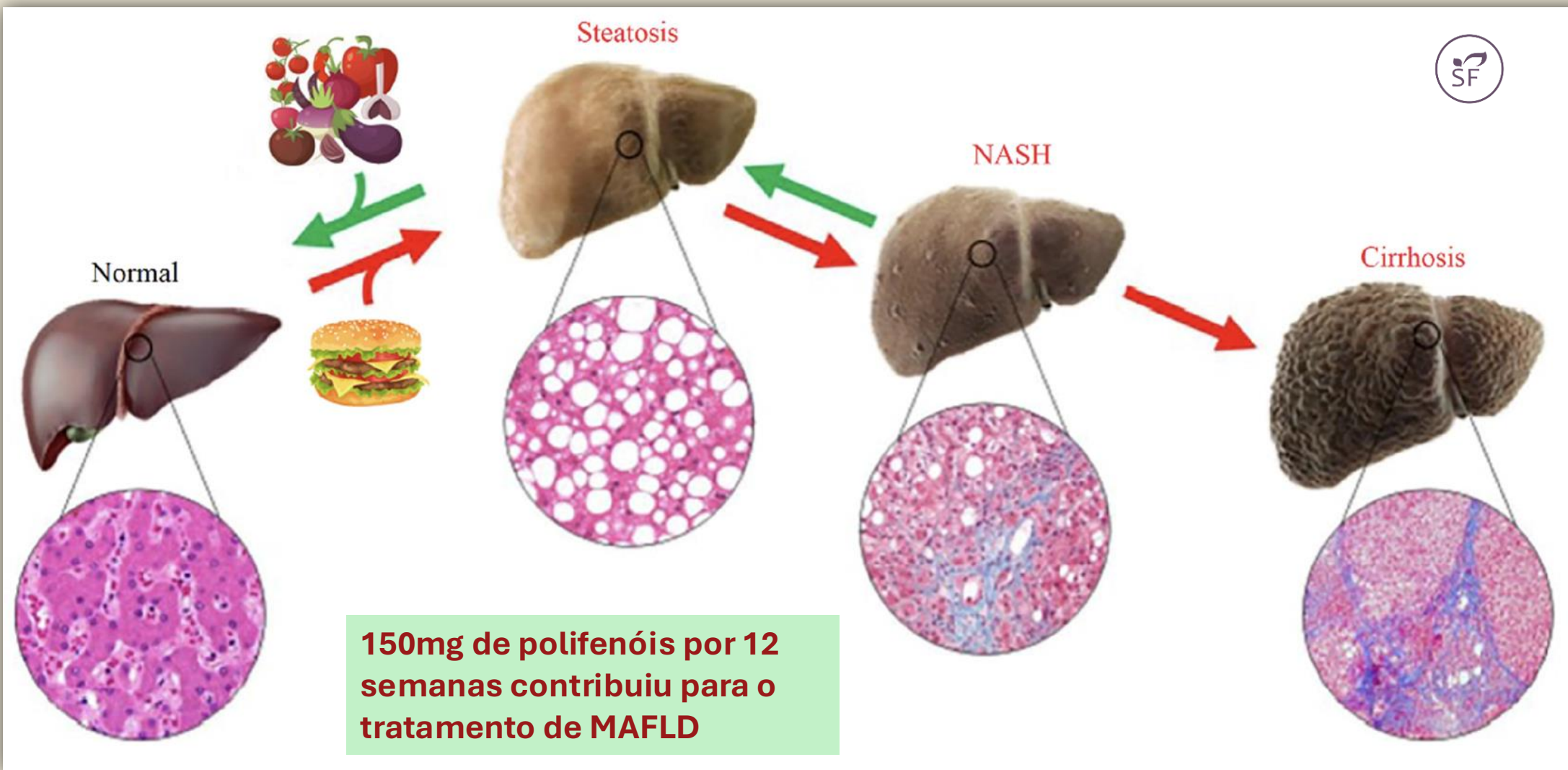
Dietary Polyphenols to Combat Nonalcoholic Fatty Liver Disease via the Gut-Brain-Liver Axis: A Review of Possible Mechanisms

Wang Z, Zeng M, Wang Z, Qin F, Chen J, He Z. Dietary Polyphenols to Combat Nonalcoholic Fatty Liver Disease via the Gut-Brain-Liver Axis: A Review of Possible Mechanisms. *J Agric Food Chem.* 2021 Mar 31;69(12):3585-3600. doi: 10.1021/acs.jafc.1c00751. Epub 2021 Mar 17. PMID: 33729777.



Dietary Polyphenols to Combat Nonalcoholic Fatty Liver Disease via the Gut–Brain–Liver Axis: A Review of Possible Mechanisms

Reparar



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CONTEÚDO DE POLIFENÓIS DOS ALIMENTOS



Maçã
(com casca - 100g):
136 mg de polifenóis



Framboesas (100g):
215 mg de polifenóis



Morangos (100g):
235 mg de polifenóis



240ml de Café:
Entre 200 e 550 mg de
polifenóis



30g de Nozes:
150 mg de polifenóis



**30g de Chocolate
Amargo (70-85%
cacau):**
499.2 mg de polifenóis



**Azeitonas
Pretas (100g):**
569 mg de polifenóis



Mirtilos (100g):
560 mg de polifenóis



30g de Cacau em Pó:
1034.4 mg de polifenóis

AASLD Practice Guidance on the clinical assessment and management of nonalcoholic fatty liver disease



Reparar



Guidance statements:

20. Patients with NAFLD who are overweight or obese should be prescribed a diet that leads to a caloric deficit. When possible, diets with limited carbohydrates and saturated fat and enriched with high fiber and unsaturated fats (e.g., Mediterranean diet) should be encouraged due to their additional cardiovascular benefits.

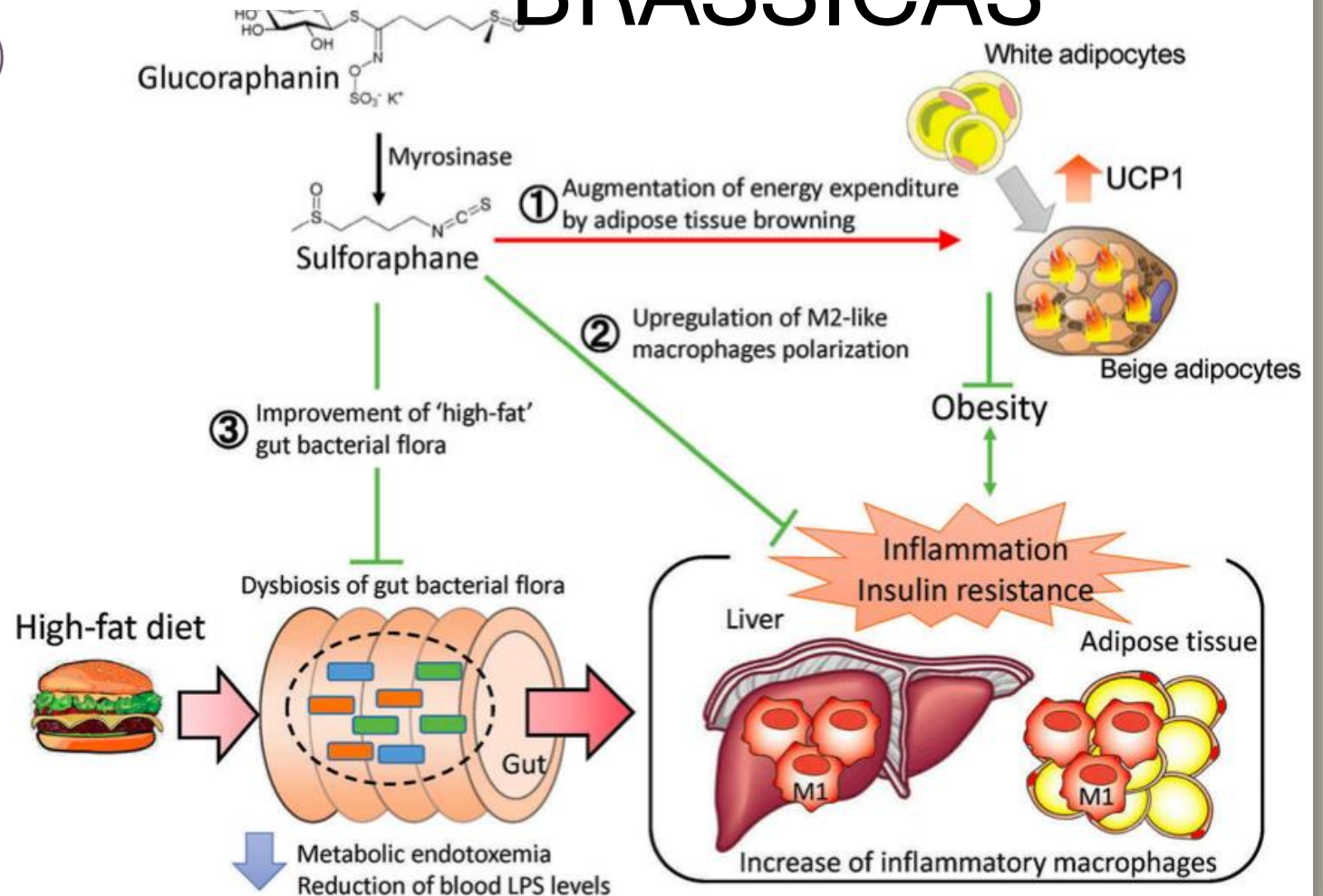
21. Patients with NAFLD should be strongly encouraged to increase their activity level to the extent possible. Individualized prescriptive exercise recommendations may increase sustainability and have benefits independent of weight loss.

Key points:

- *Weight loss improves hepatic steatosis, NASH, and hepatic fibrosis in a dose-dependent manner.*
- *The necessary support to manage comorbid disease and foster the adoption of liver protective health behaviors is best achieved using a multidisciplinary approach.*
- *Coffee consumption (caffeinated or not) of at least 3 cups daily is associated with less advanced liver disease.*

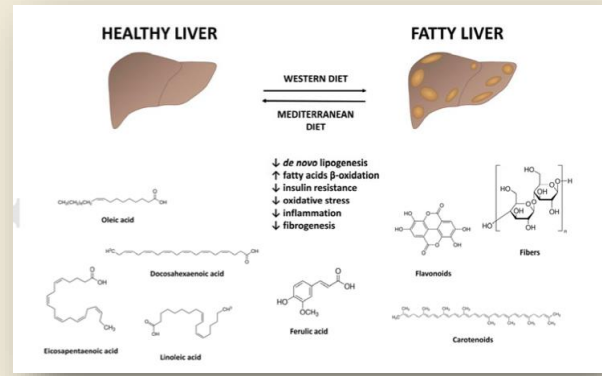
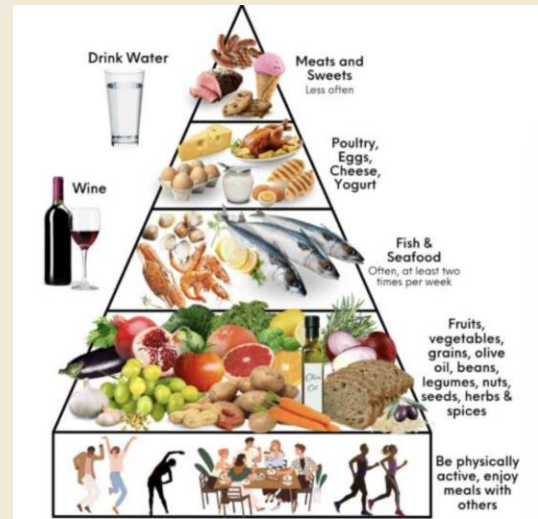
Glucoraphanin: a broccoli sprout extract that ameliorates obesity-induced inflammation and insulin resistance

- A glucorafanina, um gliocosinolato presente nas brássicas, é transformada em **sulforafano** pela mirosinase da microbiota intestinal.
- Aumento do "escurecimento" do tecido adiposo.
- Promove equilíbrio saudável de macrófagos no fígado e tecido adiposo branco com redução da inflamação.
- Reduz os níveis de LPS no sangue e endotoxemia metabólica.



Reparar

Dieta mediterrânea



nutrients *Nutrients* 2023, 15, 2250. <https://doi.org/10.3390/nu15102250>

Review

Does the Mediterranean Diet Have Any Effect on Lipid Profile, Central Obesity and Liver Enzymes in Non-Alcoholic Fatty Liver Disease (NAFLD) Subjects? A Systematic Review and Meta-Analysis of Randomized Control Trials

British Journal of Nutrition, page 1 of 9
© The Author(s), 2021. Published by Cambridge University Press on behalf of The Nutrition Society

doi:10.1017/S0007114521002270

Effect of Mediterranean diet on liver enzymes: a systematic review and meta-analysis of randomised controlled trials

Downloaded from <https://www.cambridge.org/core>

Clinical Nutrition 41 (2022) 1913–1931

Contents lists available at [ScienceDirect](https://www.sciencedirect.com)

Clinical Nutrition

journal homepage: <http://www.elsevier.com/locate/clnu>

Meta-analyses

The effectiveness and acceptability of Mediterranean diet and calorie restriction in non-alcoholic fatty liver disease (NAFLD): A systematic review and meta-analysis

Contents lists available at [ScienceDirect](https://www.sciencedirect.com)

Clinical Nutrition ESPEN

journal homepage: <http://www.clinicalnutritionespen.com>

Original article

Effects of the Mediterranean diet on cardiovascular risk factors in non-alcoholic fatty liver disease patients: A systematic review and meta-analysis

Clin Nutr ESPEN. 2020 Jun;37:148-156. doi: 10.1016/j.clnesp.2020.03.003.

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Reparar

AASLD

Déficit calórico 500-1000 kcal
Perda de peso de pelo menos 3%-5% melhorar a esteatose
Maior perda de peso (7%-10%) é necessária para melhorar a maioria das características histopatológicas da NASH, incluindo fibrose

EASL

Déficit calórico 500-1000 kcal
Perda de peso de 500 a 1000g/ semana
7 a 10% perda de peso total
Baixa a moderada ingestão de gordura e em carboidrato
Evitar bebidas e alimentos processados contendo frutose

APASL

Déficit calórico 500-1000 kcal
Contra indica dieta muito baixa em caloria

Current guidelines for the management of non-alcoholic fatty liver disease: A systematic review with comparative analysis

Leoni S, Tovoli F, Napoli L, Serio I, Ferri S, Bolondi L. Current guidelines for the management of non-alcoholic fatty liver disease: A systematic review with comparative analysis. World J Gastroenterol. 2018 Aug 14;24(30):3361-3373. doi: 10.3748/wjg.v24.i30.3361. PMID: 30122876; PMCID: PMC6092580.

AASLD

Atividade aeróbica e treinamento de resistência
mais de 150 min por semana

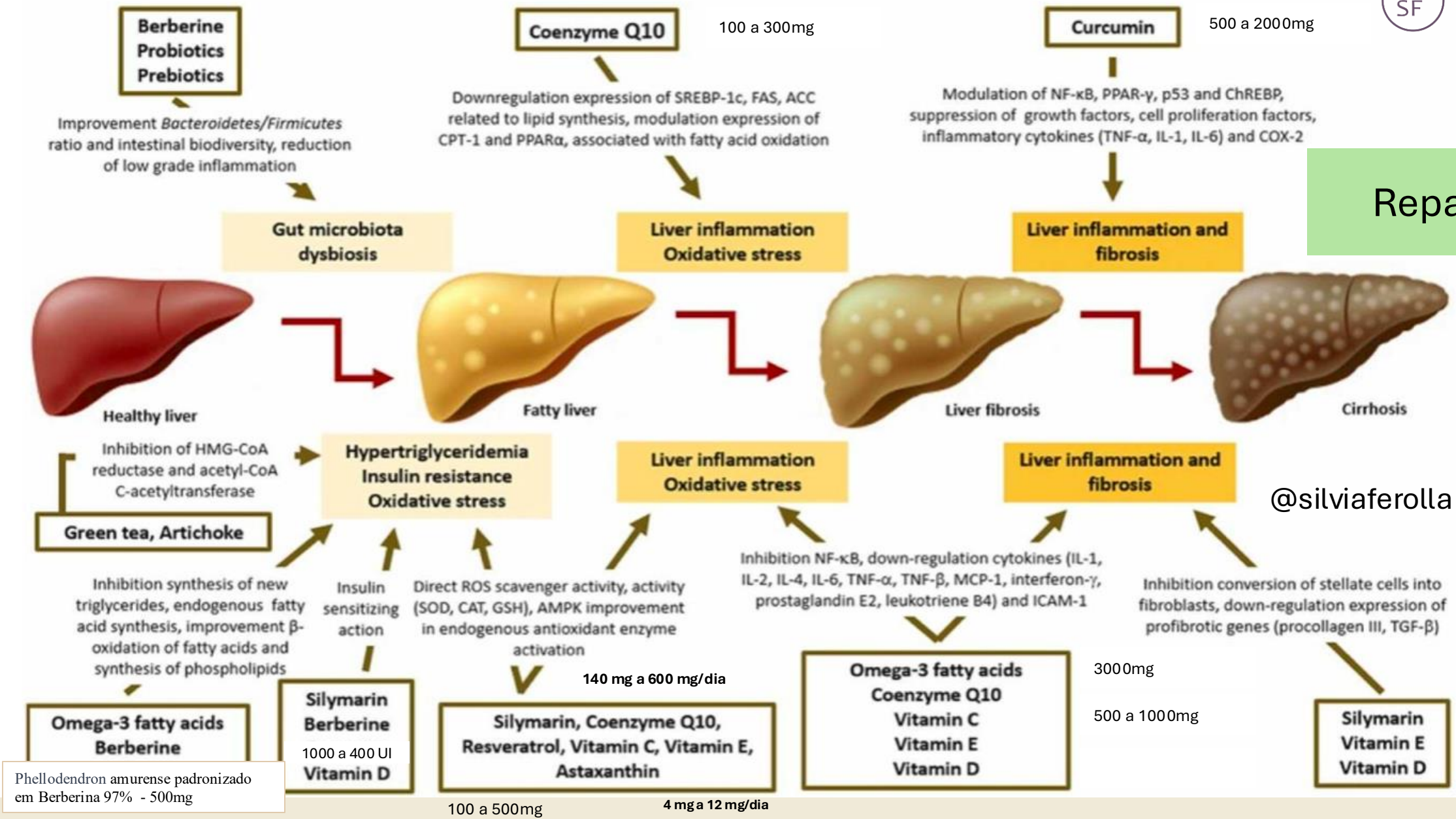
EASL

Atividade aeróbica e treinamento de resistência
150 a 200 min por semana em 3 a 5 sessões

Asia-
pacific

Atividade aeróbica e treinamento de resistência

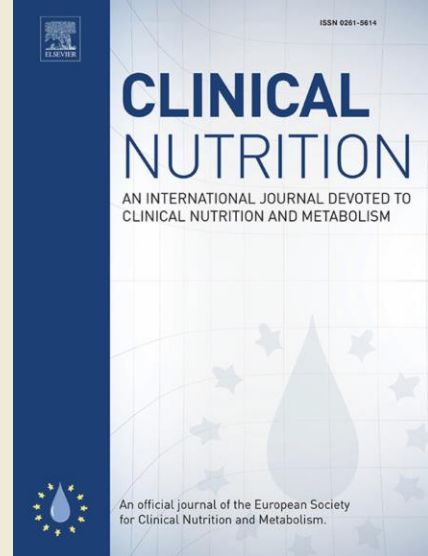
Nutraceutical actions on the four stages of NAFLD



Reparar

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Optimal probiotic combinations for treating nonalcoholic fatty liver disease: A systematic review and network meta-analysis



Background

Probiotic administration is a promising therapy for improving conditions in NAFLD patients. This network meta-analysis aimed to compare and estimate the relative effects of probiotic interventions and identify the optimal probiotic species for the treatment of NAFLD (Nonalcoholic fatty liver disease) patients.

Methods

The PubMed, Web of Science, Embase, and Cochrane databases were searched from inception to 29 January 2024 to identify RCTs that were published in English. The GRADE framework was used to assess the quality of evidence contributing to each network estimate.

Results

A total of 35 RCTs involving 2212 NAFLD patients were included in the analysis. For primary outcomes, *Lactobacillus + Bifidobacterium + Streptococcus* exhibited the highest probability of being the finest probiotic combination in terms of enhancing acceptability as well as reducing AST (SMD: -1.95 95% CI: -2.90, -0.99), ALT (SMD = -1.67, 95% CI: -2.48, -0.85), and GGT levels (SMD = -2.17, 95% CI: -3.27, -1.06). In terms of the secondary outcomes, *Lactobacillus + Bifidobacterium + Streptococcus* was also the best probiotic combination for reducing BMI (SMD = -0.45, 95% CI: -0.86, -0.04), LDL levels (SMD = -0.45, 95% CI: -0.87, -0.02), TC levels (SMD = -1.09, 95% CI: -1.89, -0.29), and TNF- α levels (SMD = -1.73, 95% CI: -2.72, -0.74).

Conclusion

This network meta-analysis revealed that *Lactobacillus + Bifidobacterium + Streptococcus* may be the most effective probiotic combination for the treatment of liver enzymes, lipid profiles, and inflammation factors. These findings can be used to guide the development of a probiotics-based treatment guideline for NAFLD since there are few direct comparisons between different therapies.

Reinocular

- 35 ECRs com 2212 pacientes com DHGNA.
 - Lactobacillus + Bifidobacterium + Streptococcus mostrou-se a melhor combinação probiótica.
 - AST: Redução significativa com DME de -1,95 (IC 95%: -2,90 a -0,99).
 - ALT: Redução significativa com DME de -1,67 (IC 95%: -2,48 a -0,85).
 - GGT: Redução significativa com DME de -2,17 (IC 95%: -3,27 a -1,06).
 - IMC: Leve redução com DME de -0,45 (IC 95%: -0,86 a -0,04).
 - LDL: Leve redução com DME de -0,45 (IC 95%: -0,87 a -0,02).
 - CT (Colesterol Total): Redução com DME de -1,09 (IC 95%: -1,89 a -0,29).
 - TNF- α : Redução significativa com DME de -1,73 (IC 95%: -2,72 a -0,74).
- Conclusão :** Lactobacillus + Bifidobacterium + Streptococcus pode ser a combinação mais eficaz para tratar enzimas hepáticas, perfis lipídicos e fatores inflamatórios em pacientes com DHGNA.

@silviaferolla

Reinocular



L. acidophilus

L. bulgaricus

B. breve

B. lactis

S. thermophilus

L. plantarum

L. paracasei

L. rhamnosus

L. casei

B. bifidum

**PROBIÓTICO E
NASH/MASH**

**mínimo de
SEMANAS!**

8

Obstructive sleep apnea and severity of nonalcoholic fatty liver disease

Results: Fifty-one patients were evaluated, 80.4% had systemic arterial hypertension (SAH), 68.6% type 2 diabetes mellitus, 62.7% dyslipidemia and 96.1% MS. Regarding the histological evaluation (n = 48), all had steatosis, 95.8% steatohepatitis and 83.3% fibrosis. In polysomnography, 80.4% were group 1 and 19.6% group 2. In univariate analysis, no correlation was found between steatosis severity, NASH and presence or severity of fibrosis with OSA. A multivariate analysis adjusted for obesity level, found that patients with moderate to severe OSA had an increased risk of hepatic fibrosis (odds ratio 1.22, 95% confidence interval: 1.02-1.45, P = 0.027).

Conclusion: The present study demonstrated an association between fibrosis and moderate to severe OSA, regardless of obesity.

n=51

Na análise multivariada ajustada por obesidade, foi verificado que pacientes com apneia do sono moderada a grave apresentavam risco aumentado de fibrose hepática (OR: 1.22)

Reequilibrar

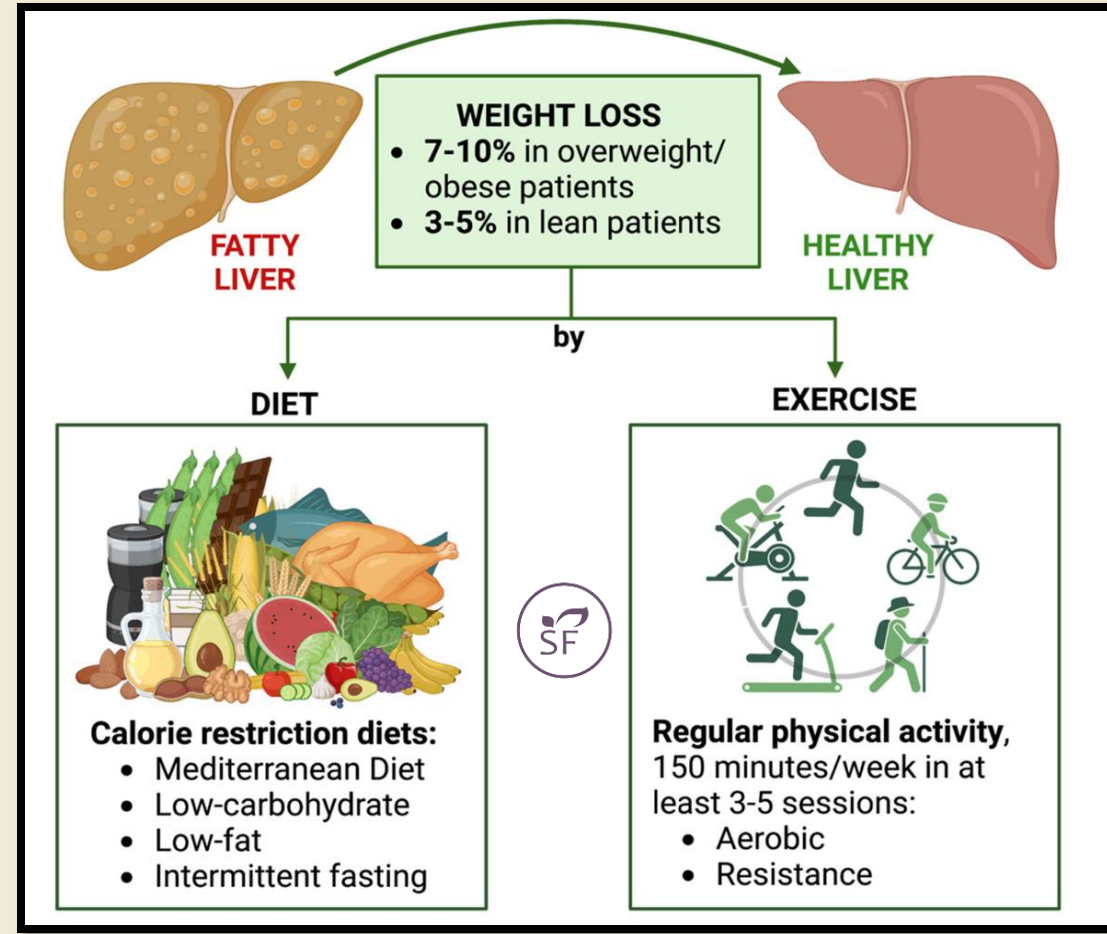
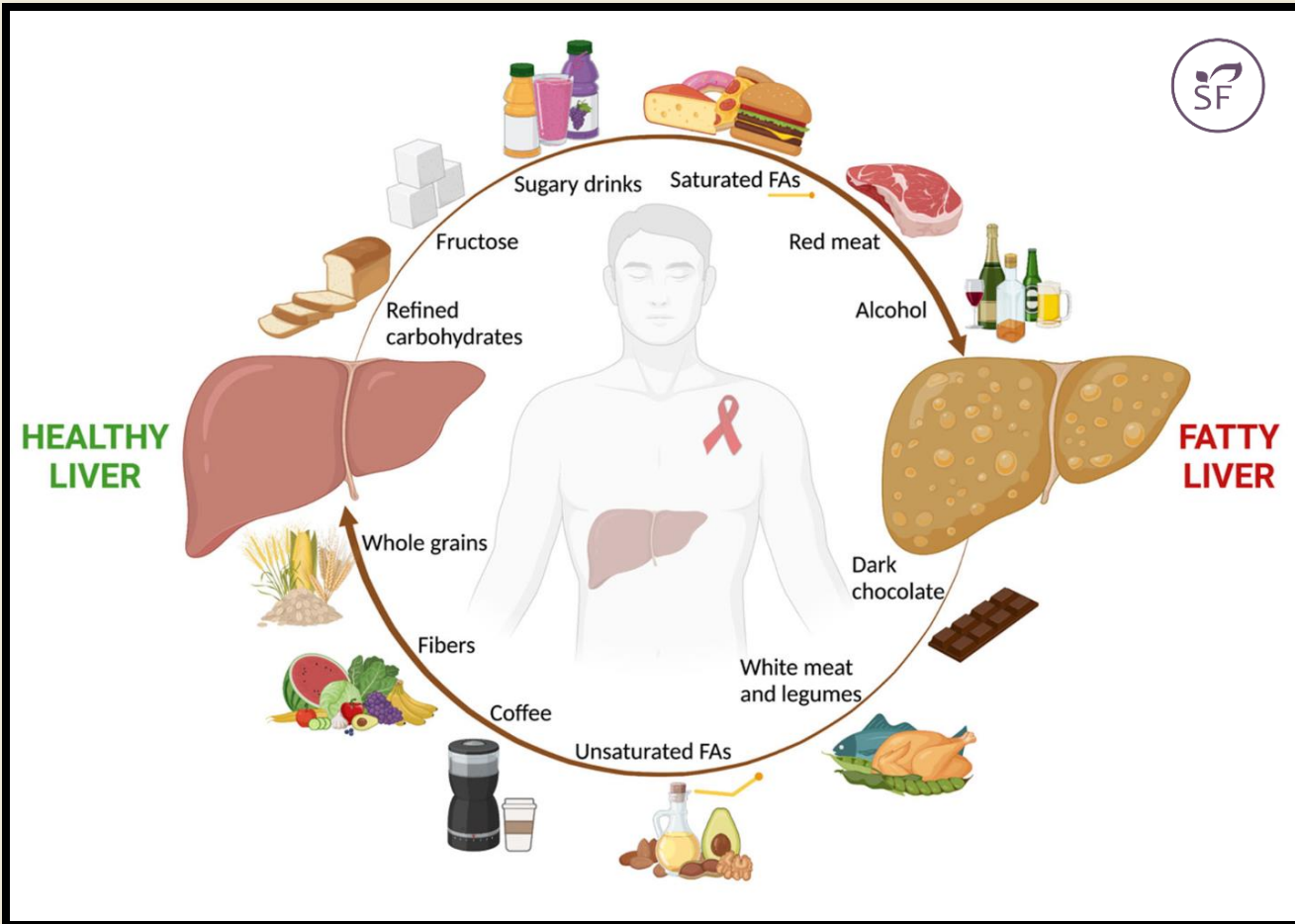


SONO



HIPÓXIA INTERMITENTE - ALTERA NÍVEIS DE EIXO HPA-CORTISOL E AUMENTA RI!

Reavaliar SEMPRE!





NAFLD

A ESTEATO-HEPATITE METABÓLICA



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