



Introducción

Figure 1. Working

representation

Obese and type 2 diabetic (T2D) humans develop increased insulin resistance (IR), which appears derived from an increased innate immune Toll like receptor 4 (TLR4) activity in skeletal muscle. Besides the role of type 1 innate lymphoid cells (ILC1s) in regulating systemic inflammatory diseases involved in the development of diabetes ¹, monocyte/macrophages play determinant role regulating the systemic availability of lipids for metabolic purposes².

IMMUNONUTRITIONAL PRECISION NUTRITION TO REDUCE THE RISK OF SUFFERING METABOLIC DISEASES

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Obietivos

The purpose of this study was

to examine the role of replacement of critical

nutrients, potential modulators of TLR4 activity, in cookies with the hypothesis that it would result in a biased functional differentiation of **ILC1s** with positive effects in the prevention of early immunonutritional imbalances leading to obesity and T2D (Figure 1).

Métodos

Participants were healthy adult Caucasians of both sexes and included 60 lean subjects with no medical symptoms. Participants were randomly divided into groups, and each group received nutritional instructions to eat cookies (group A, commercial formulation; group B, immunonutritional agonistscontaining formulation) over a 12days study period, while adhere to a normocaloric diet (2057.3 ± 614.7 kcal) according to physical activity.

Resultados

Group B displayed downward trends in fat mass (Group A. +0.58% vs Group B, -0.73%) together with broad variations in muscle mass (Group A, -0.27% vs Group B, -5.5%), suggesting variations in the body extracellular water. However, it was not quantified significant differences in body weight changes. These changes were associated to a **better control of IR** (Δ TyG index; group A, +0.12 vs group B, +0.07 over the study period). These changes could be associated to

significant variations in innate immune effectors (Figure 2).





Conclusiones

Dietary

immunonutritional agonists targeting innate immune biology enable selective functional differentiation of key immune regulators to reduce the risk of suffering dysregulated metabolism leading to obesity and T2D.

Agradecimientos:

Grants Food4ImNut [PID2019-107650RB-C22, Ministry of Science, Innovation and Universities-MICIU] and CYTED [La ValSe-Food, 119RT0S67].

References:

¹Diabetes & Metabolism 2019, 45(4), 341-346; ² Science 2021, 2;373(6550):eabe9383

