PHLPP1 PHOSPHATASE INHIBITION IN HYPOTHALAMUS RESTORES INSULIN SIGNALING AND ACTION AND REDUCES ADIPOSITY IN OBESE RATS

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Abstract

PH Domain and Leucine Rich Repeat Protein Phosphatase 1 (PHLPP1) regulates the Akt activity in hypothalamus of adult rats. Obesity is responsible by the increase of PHLPP1 in the hypothalamus, inhibiting Akt activity, impairing the insulin action in the hypothalamus, resulting in insulin resistance. Thus, the silencing of this phosphatase could act in the regulation of Akt activity.

Key words: *PHLPP1, Obesity, Hypothalamus.*

Introduction

Obesity results from imbalance between food intake and energy expenditure. The energy homeostasis is regulated by hypothalamic neurons that receive different neural, hormonal and metabolic signals. Insulin is one of the main hormones that regulate energy homeostasis and acts through a cascade of intracellular signaling that depends on the activation of several proteins, such as Akt. Our hypothesis is that PHLPP1 inhibits Akt activity by dephosphorylating serine 473 residues, impairing the insulin action in the hypothalamus^{1,2}.

The objectives are investigate PHLPP1 protein expression in the hypothalamus of diet induced obesity (DIO) rats and to assess whether the PHLPP1 silencing improves insulin action and decreases body adiposity

Results and Discussion

Image 1: Diet induced obesity

A. Intolerance Insulin Test (ITT), standard chow (Chow) and high fat diet (HFD). *p= 0,0002. B. PHLPP1 expression in hypothalamus. *p= 0,007.

Image 2 Treatment to silence PHLPP1 expression.

A: Percent of weight loss after the treatment with siRNA. *p=0,0098. B: Food intake. C: Mass of retroperitoneal adipose tissue. *p=0,0497. D: Fasting blood glucose after treatment with siRNA. *p=0,0359.

Conclusions

Hypothalamic PHLPP1 silencing in DIO animals restored, at least in part, the insulin signaling, promoting reduction in body weight and adiposity.

Acknowledgement

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