

Kinin B1 and B2 receptor deficiency protects against obesity induced by a high-fat diet and improves glucose tolerance in mice

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Abstract: The kallikrein-kinin system is well known for its role in pain and inflammation, and has been shown recently by our group to have a role also in the regulation of energy expenditure. We have demonstrated that B1 receptor knockout (B1KO) mice are resistant to obesity induced by a high-fat diet (HFD) and that B1 receptor expression in adipocytes regulates glucose tolerance and predisposition to obesity. However, it is also known that in the absence of B1 receptor, the B2 receptor is overexpressed and can take over the function of its B1 counterpart, rendering uncertain the role of each kinin receptor in these metabolic effects. Therefore, we investigated the impact of ablation of each kinin receptor on energy metabolism using double kinin receptor knockout (B1B2KO) mice. Our data show that B1B2KO mice were resistant to HFD-induced obesity, with lower food intake and feed efficiency when compared with wild-type mice. They also had lower blood insulin and leptin levels and higher glucose tolerance after treatment with an HFD. Gene expression for tumor necrosis factor- α and C-reactive protein, which are important genes for insulin resistance, was reduced in white adipose tissue, skeletal muscle, and the liver in B1B2KO mice after the HFD. In summary, our data show that disruption of kinin B1 and B2 receptors has a profound impact on metabolic homeostasis in mice, by improving glucose tolerance and preventing HFD-induced obesity. These novel findings could pave the way for development of new pharmacological strategies to treat metabolic disorders such as insulin resistance and obesity.

Keywords: kallikrein-kinin system, B1/B2 receptors, obesity, glucose tolerance, insulin resistance

Introduction

Obesity and diabetes are serious health problems that are of particular concern given their increasing incidence around the world. The World Health Organization estimates that at least 2.8 million people die each year as a result of being overweight or obese, and currently more than 1.4 billion adults are overweight.¹ The excess energy intake in obese individuals is stored as fat tissue, which is an important organ in the regulation of metabolic homeostasis.² In this scenario, obesity and diabetes are closely related since the accumulation of adipose tissue alters the pattern of secretion of molecules involved in physiological processes, such as glucose uptake, lipid metabolism, inflammation, and vascular homeostasis,^{3,4} which may lead to insulin resistance and development of diabetes.

Study of the kallikrein-kinin system (KKS) has contributed to our understanding of many biological processes, including inflammation, nociception, vascular permeability, blood pressure regulation, and smooth-muscle contraction and relaxation.^{5,6}

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The KKS consists of distinct enzymes called kallikreins, which cleave their precursors, kininogens, producing kinins, a small group of peptides. Kinins activate two specific G-protein coupled receptors, ie, B1 and B2. Although it has been known since 1932 that the KKS participates in energy metabolism,⁷ it has only been recently that these mechanisms have started to be elucidated.^{8,9} Several studies have highlighted the role of the B2 receptor in glucose homeostasis, in particular its effects on insulin-mediated glucose uptake.^{10,11} It was also shown that B1 receptor blockade by a specific antagonist could prevent hyperglycemia and insulinitis, demonstrating that not only the B2 receptor but also the B1 receptor participates in the regulation of glucose uptake.¹²

More recently, the participation of the B1 receptor in insulin secretion has been demonstrated,¹³ as well as in the secretion of leptin by adipose tissue.¹⁴ More evidence concerning the role of KKS in metabolism came from B1 receptor knockout (B1KO) mice which were shown to be hypersensitive to leptin and resistant to obesity induced by a high-fat diet (HFD).^{15,16} Moreover, overexpression of the B1 receptor in adipocytes demonstrated that this receptor regulates glucose tolerance and determines predisposition to obesity.¹⁷ In agreement with these observations, recent findings have highlighted the inflammatory role of the B1 receptor in insulin-resistant glucose-fed rats, in which several pathological symptoms were reversed by treatment with specific B1 receptor antagonists.^{18,19}

We can infer from the above data that KKS participates in development of obesity and related diseases, including insulin resistance and type 2 diabetes. However, the role played by each kinin receptor in these processes remains unclear, given the recurring evidence that in the absence of one receptor, the other is overexpressed and can compensate its functions.^{20,21}

Therefore, we investigated energy metabolism in mice with deletion of both kinin receptors (B1B2KO). We found that B1B2KO mice had lower food and energy intake, lower feed efficiency, and lower serum insulin and leptin levels when compared with wild-type (WT) control mice. Hypothalamic mRNA expression of genes related to regulation of food intake and metabolism was unchanged in B1B2KO mice. However, when these mice were subjected to an HFD, they showed increased glucose tolerance when compared with their WT control littermates. The tumor necrosis factor- α (TNF- α) and C-reactive protein (CRP) genes code for pro-inflammatory proteins that are also known for their role in insulin resistance, and show reduced mRNA expression in white adipose tissue, skeletal muscle, and the liver. These novel data suggest that B1 and B2 receptors have

pivotal roles in energy balance by controlling glucose homeostasis and may be targets for insulin resistance and diabetes therapy.

Materials and methods

Animals

Healthy 12-week-old C57Bl/6 male B1B2KO mice and their WT littermates were used in this study. They were obtained from the animal house at Universidade Federal de São Paulo, São Paulo, and maintained on a 12-hour light/12-hour dark cycle at 22°C, with free access to standard chow and drinking water. The experimental protocols were approved by the institutional ethics committee (01730/08).

Genotyping of kinin B1 and B2 knockout mice

Genomic DNA was isolated from a mice tail fragment using an NaOH extraction protocol.²² Kinin B1 and B2 receptors were amplified by polymerase chain reaction (PCR) using the following primers: B1 receptor (NM_007539.2) sense 5'-TGAAGCTGTGAGCTCTTTG-3' and antisense 5'-GCTACCCAGATGAGCAGGCA-3'; B2 (NM_009747.2) receptor sense 5'-TGTCCTCAGCGTGTCTTCC-3' and antisense 5'-GGTCCTGAACACCAACATGG-3'.

Kinin B1 and B2 receptor function

The functionality of B1 and B2 receptors was assayed by testing the contractile response of smooth muscle strips (from the fundus of the stomach) to bradykinin and des-Arg(9)-bradykinin (Bachem Inc.). Briefly, smooth muscle strips were extracted from the mice and kept at 37°C in oxygenated (95% O₂, 5% CO₂) modified Krebs buffer (119 NaCl mM, 4.7 KCl mM, 1.2 MgSO₄·7H₂O mM, 2.5 CaCl₂·2H₂O mM, 1.2 KH₂PO₄ mM, 25.0 NaHCO₃ mM, and 5.5 glucose mM). After an equilibration period of 45–60 minutes at a resting tension of 0.5 g, experiments were initiated by checking the viability of the prepared specimen by exposure to carbachol (Sigma-Aldrich) at 10⁻⁵ M. Kinin agonists, ie, bradykinin 10⁻⁶ M and Des-Arg(9)-bradykinin, 10⁻⁶ M, were then applied, and the contractile responses were recorded using an isometric transducer (TRI202P, Panlab) connected to a computerized system for data acquisition and analysis (PowerLab, AD Instruments).

Treatment with a high-fat diet

Mice were fed ad libitum for 12 weeks with a control diet (10% kcal from fat) or an HFD (45% kcal from fat) from Research Diets Inc. Each mouse was put in an individual cage, with weekly weight and food intake recording.

After treatment, the mice were euthanized for blood and tissue collection. Feed efficiency was calculated by the ratio of total weight gain and calories of food ingested in both WT and B1B2KO mice after treatment with the HFD.

Leptin and insulin serum levels

Serum samples were separated and used for leptin (R&D Systems) and insulin (Millipore) measurement as specified by the manufacturers.

Glucose tolerance test

Mice were fasted for 8 hours before the glucose tolerance test. A small fragment of tail tissue was cut for blood collection and glucose was quantified using an Advantage II glucometer (Roche). Glucose measurements were carried out before and at 15, 30, 60, and 120 minutes after intraperitoneal injection of glucose 1 g/kg body weight.

Insulin and leptin sensitivity test

For the insulin sensitivity test, 0.01 U/kg of recombinant human insulin (Lilly) was injected intraperitoneally. Glucose levels were assessed in blood collected from the tail before and 15, 30, and 60 minutes after insulin injection immediately after treatment with the HFD. For leptin sensitivity, recombinant mouse leptin (R&D Systems) was injected as an intraperitoneal bolus of 40 µg/day for 5 days, with food intake measured under basal conditions. Food intake measured 5 days before the leptin injections was considered as basal consumption.

Gene expression

Total RNA (2 µg) was extracted from the hypothalamus, white adipose tissue, skeletal muscle, and liver from WT and B1B2KO mice using TRIzol Reagent (Invitrogen). It was reverse-transcribed using M-MLV (Invitrogen) to cDNA following the manufacturer's instructions. The product was amplified by real-time PCR using the SYBR Green (Fermentas) protocol. Prolylcarboxipeptidase (PRCP), melanocortin 4 receptor (MC4R), neuropeptide Y (NPY), agouti-gene-related protein (AgRP), pro-opiomelanocortin (POMC), cocaine- and amphetamine-regulated transcript (CART), leptin, TNF-α, and CRP mRNAs were quantified as a relative value compared with the internal reference, β-actin. Primers used for real-time PCR were as follows: PRCP (NM_001278581.1) sense 5'-CAGGAGCTCAAGGTCAACCTGTC-3' and antisense 5'-GAACCATAACCATCAAGCTGCCAGAT-3'; MC4R (NM_016977.4) sense 5'-GCTGTGGCAGATATGCTGGTGAG-3' and antisense 5'-CGAGCAAGGAGCTACAGATCACAG-3'; NPY

(NM_023456.2) sense 5'-CGCTCTGCGACACTACATCAA-3' and antisense 5'-GGGCTGGATCTCTTGCCAT-3'; AgRP (NM_001271806.1) sense 5'-GGTGCTAGATC CACA GAACCG-3' and antisense 5'-CCAAGCAGGACTCGTGCAG-3'; POMC (NM_001278581.1) sense 5'-CGAGGCCTTTCCCCTAGAGT-3' and antisense 5'-CCAGGACTTGCTCCAAGCC-3'; CART (NM_013732.7) sense 5'-TGGATGATGCGTCCCATGA-3' and antisense 5'-CGGAATGCGTTTACTCTTGAGC-3'; leptin (NM_008493.3) sense 5'-AGCATTCAGG GCTAACATCC-3' and antisense 5'-TATCCGCCA AGCAGAGGGT-3'; TNF-α (NM_013693.3) sense 5'-CACCACGCTCTTCTGTCTAC-3' and antisense 5'-GATCTGAGTGTGAGGGTCTGG-3'; CRP (NM_007768.4) sense 5'-GGAGAAGCTACTCTGGTGCCTTCT-3' and antisense 5'-CACACAGTAAAGGTGTTCAAGTGGCT-3', and β-actin (NM_007393.3) sense 5'-GAACCTAAGGC CAACCGTGAA-3' and antisense 5'-GACCAGAGGCAT ACAGGGACAAC-3'. Quantitative values for mRNA expression were obtained by the parameter $2^{-\Delta Ct}$, in which ΔCt represents the subtraction of the β-actin Ct values from the ones of the target gene.

Treatment with kinin B1 and B2 receptor antagonists

Healthy 12-week-old male C57Bl/6 mice were injected intraperitoneally daily for 7 days with the B1 receptor antagonist R-715 (1 mg/kg) together with the B2 receptor antagonist HOE-140 (500 µg/kg) or vehicle (saline). Both antagonists were obtained from Bachem Inc. After treatment, the mice in both groups were euthanized and their tissues collected for analysis of leptin, CRP, and TNF-α gene expression.

Statistical analysis

The results are reported as the mean ± standard error of the mean. Statistically significant differences in mean values were evaluated by the independent Student's t-test when two groups were compared, or by the ANOVA test when three or more groups were analyzed. A *P*-value ≤ 0.05 was considered to be statistically significant.

Results

Genetic and pharmacological assays for kinin B1 and B2 receptors

Deletion of both kinin receptors in the mice was confirmed by PCR genotyping, as demonstrated in Figure 1. In the DNA samples from WT mice, B1 (Figure 1A) and B2 receptor (Figure 1B) gene fragments were amplified by PCR, whereas

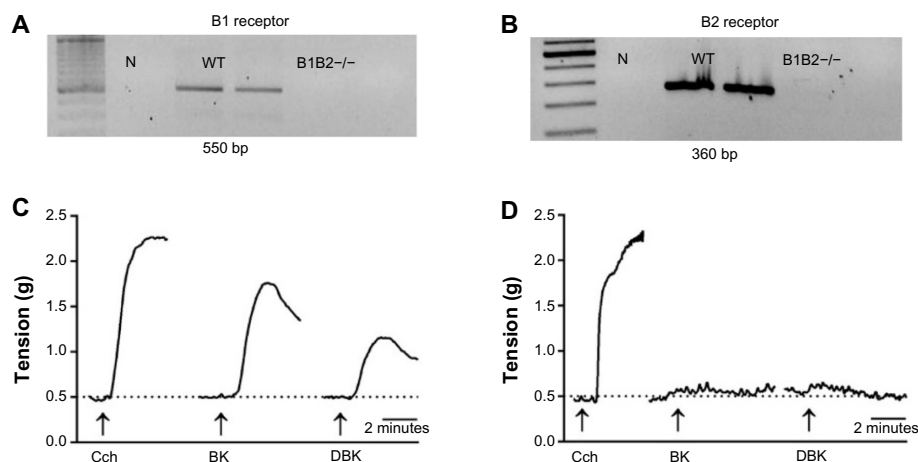


Figure 1 Genotyping the kinin B1 and B2 receptor in B1B2KO mice.

Notes: (A) B1 and (B) B2 receptor gene fragments were amplified by polymerase chain reaction. The contractile response of smooth muscle strips from the gastric fundus to Cch (positive control), BK, and DBK in (C) WT and (D) B1B2KO mice.

Abbreviations: BK, bradykinin; DBK, des-Arg(9)-bradykinin; N, negative control; WT, wild-type; B1B2^{-/-}, B1B2KO; Cch, carbachol.

in B1B2KO mice the fragments could not be amplified. The pharmacological functionality of the B1 and B2 receptors was assayed by testing the contractile response of smooth muscle strips from the gastric fundus to bradykinin and des-Arg(9)-bradykinin. WT mice preparations were responsive to carbachol and to B1 and B2 receptor agonists (Figure 1C). On the other hand, double knockout mice were responsive to carbachol but not to the kinins (Figure 1D), confirming the absence of B1 and B2 receptors in this genetic mouse model.

B1B2KO mice protected against HFD-induced obesity

Body weight was measured for 12 weeks in B1B2KO mice on a control diet or the HFD. The mice had a reduced body weight and smaller visceral fat depots after the HFD but not after the control diet (Figure 2A and B). To investigate why B1B2KO mice were protected against HFD-induced weight gain, food and energy intake was measured every week during the HFD and chow diet regimen. As shown in Figure 2C and D, WT mice showed higher food intake and calories ingested when compared with B1B2KO mice regardless of the type of diet. Also, the B1B2KO mice on the HFD showed lower feeding efficiency (Figure 2E), indicating more energy expenditure after the HFD.

Gene expression in the hypothalamus was unchanged in B1B2KO mice

Given the reduced food and energy intake found in the B1B2KO mice, we also investigated the mRNA expression of important hypothalamic genes associated with satiety and energy balance. However, expression of orexigenic (NPY, AgRP), anorexigenic

(POMC, CART), PRCP, and MC4R genes was not significantly different between B1B2KO and WT mice treated with a control diet or the HFD (data not shown).

Insulin and leptin levels were reduced in B1B2KO mice

In order to better understand the metabolic profile in B1B2KO mice, we measured two important hormones involved in energy homeostasis, ie, insulin and leptin. Insulin and leptin levels (protein in the serum and mRNA from white adipose tissue) were decreased in B1B2KO mice when compared with WT mice using both regimens (Figure 3A–C). No difference in leptin sensitivity was observed in B1B2KO mice after the HFD (Figure 3D). Also, B1B2KO mice on the control diet or HFD did not present changes in insulin sensitivity when compared with WT mice (Figure 3E).

B1B2KO mice showed improved glucose tolerance

Differences in insulin levels between B1B2KO and WT mice suggested a potential impact of the B1 and B2 receptors on blood glucose levels and glucose tolerance. B1B2KO mice showed lower glucose levels than WT mice after treatment with the HFD (Figure 4A). Accordingly, in this condition, B1B2KO mice also showed higher glucose tolerance after glucose stimulation (Figure 4B).

Genes related to insulin resistance were downregulated after HFD

Given the importance of inflammation in the pathogenesis of insulin resistance, we measured the expression of two inflammatory mediators in different metabolic tissues of

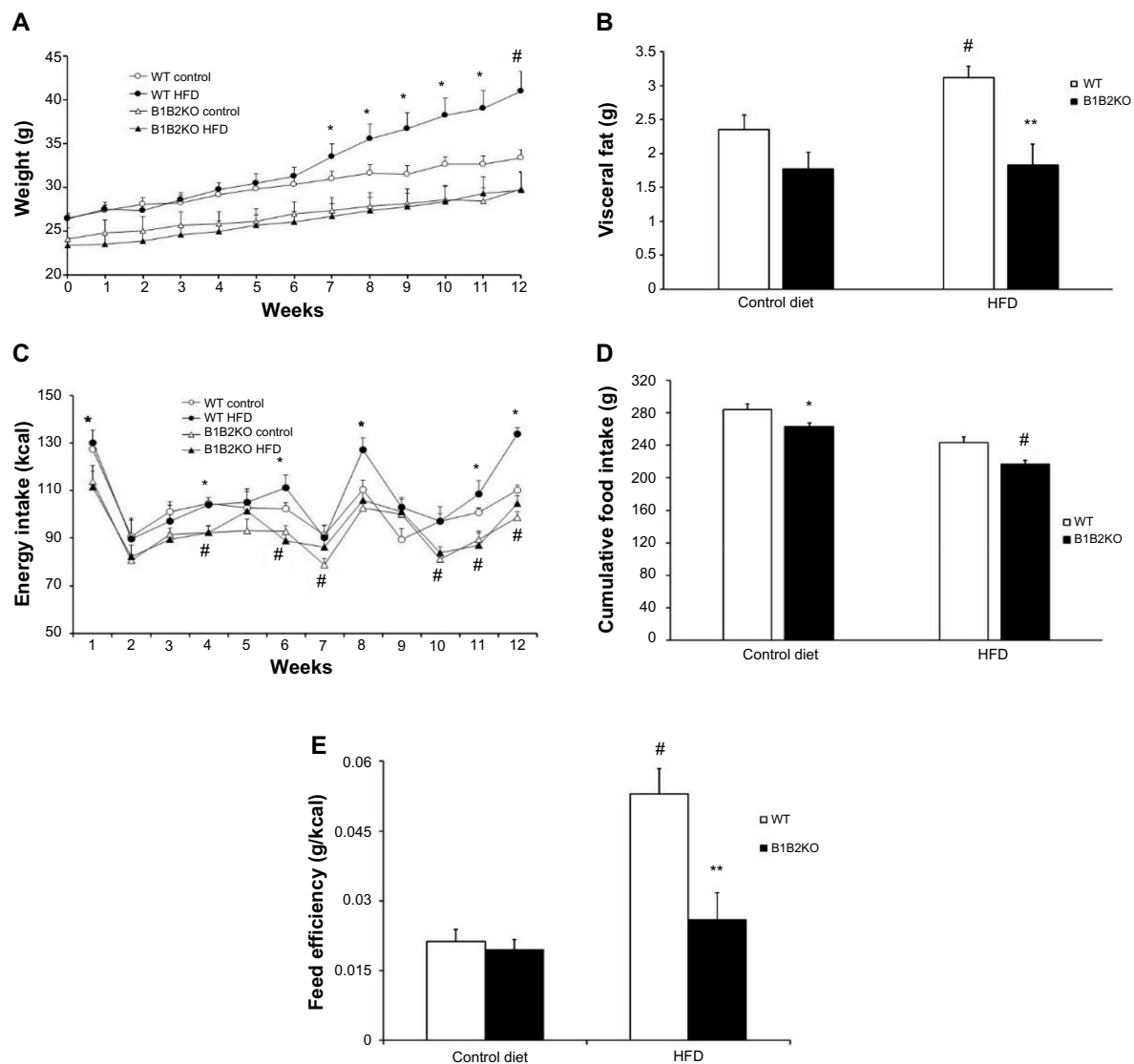


Figure 2 B1B2KO mice are protected against obesity induced by HFD treatment and present lower food intake and feed efficiency.

Notes: (A) Body weight after 12 weeks of HFD treatment. * $P < 0.05$ WT HFD versus B1B2KO groups; # $P < 0.05$ versus all groups. Data are expressed as the mean $\pm$ standard error of the mean and represent mouse weight per gram. White circles indicate WT control, black circles indicate WT HFD, white triangles indicate B1B2KO control, and black triangles indicate B1B2KO HFD. (B) Visceral fat weight after 12 weeks of HFD treatment. White bars indicate WT control and HFD, and black bars indicate B1B2KO control and HFD. # $P < 0.05$ WT HFD versus WT control; ** $P < 0.01$ WT HFD versus B1B2KO HFD. Data are expressed as the mean $\pm$ standard error of the mean and represent visceral fat depot weight per gram. (C) Energy intake after 12 weeks of HFD treatment. Energy intake was calculated based on information from the chow manufacturer. White circles indicate WT control, black circles indicate WT HFD, white triangles indicate B1B2KO control, and black triangles indicate B1B2KO HFD. * $P < 0.05$ WT HFD versus B1B2KO HFD; # $P < 0.05$ WT control versus B1B2KO control. Data are expressed as the mean $\pm$ standard error of the mean and represent energy intake (kCal). (D) Cumulative food intake after 12 weeks of HFD treatment. White bars indicate WT control and HFD, black bars indicate B1B2KO control and HFD. * $P < 0.05$ WT control versus B1B2KO control; # $P < 0.05$ WT HFD versus B1B2KO HFD. Data are expressed as the mean $\pm$ standard error of the mean and represent chow per gram. (E) Feed efficiency after 12 weeks of HFD treatment. Feed efficiency was calculated by the ratio of total weight gain and calories of food ingested after the HFD treatment in both WT and B1B2KO mice. White bars indicate WT control and HFD, black bars indicate B1B2KO control and HFD. $P < 0.01$ WT HFD versus control diet group. ** $P < 0.01$ WT HFD versus B1B2KO HFD. Data are expressed as the mean $\pm$ standard error of the mean and represent energy intake (kCal).

Abbreviations: HFD, high-fat diet; WT, wild-type.

double knockout mice. TNF- α mRNA expression was reduced in both white adipose tissue and skeletal muscle in B1B2KO mice after the HFD (Figure 5A and B). Similarly, CRP mRNA expression was decreased in the livers of the B1B2KO mice after the HFD (Figure 5C). In addition, WT mice were treated for 7 days with specific B1 (R-715) and B2 (HOE-140) receptor antagonists. Gene expression of leptin and TNF- α in white adipose tissue and skeletal muscle was

measured and CRP expression was evaluated in the liver, but no differences were observed (data not shown).

Discussion

Using both genetic and pharmacological approaches, the role of kinin receptors in energy balance has started to be elucidated in recent years. However, it is known that when one of the kinin receptors is blocked or deleted, the other receptor

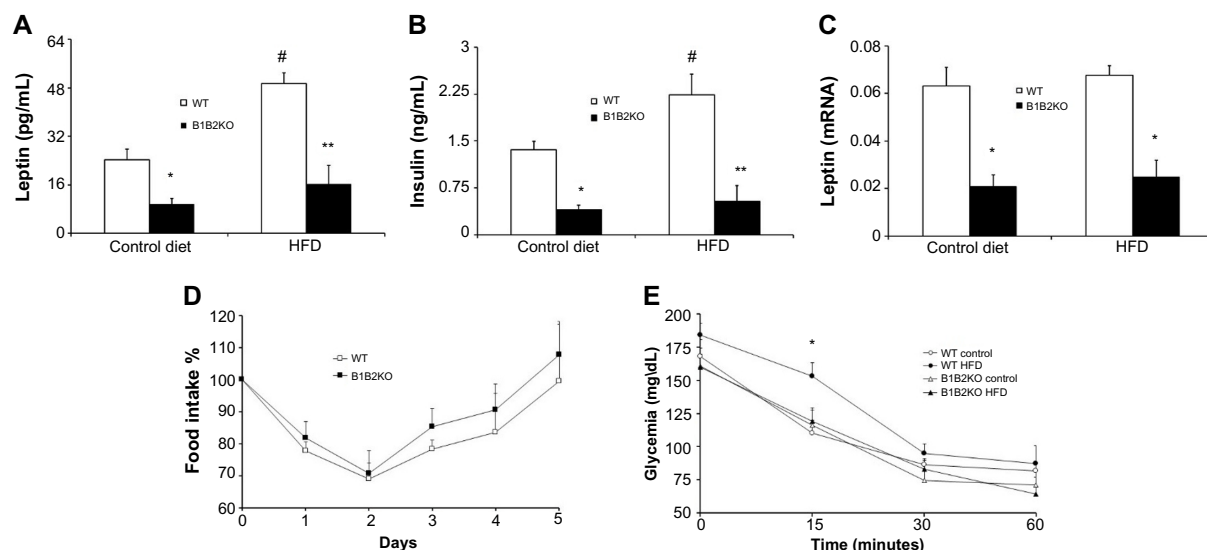


Figure 3 B1B2KO mice have low insulin and leptin levels.

Notes: (A) Serum leptin levels after 12 weeks of HFD treatment. (B) Serum insulin levels after 12 weeks of HFD treatment. (C) Leptin mRNA expression in white adipose tissue after 12 weeks of HFD treatment. White bars indicate WT control and HFD, and black bars indicate B1B2KO control and HFD. * $P < 0.05$ B1B2KO control versus WT control; ** $P < 0.01$ B1B2KO HFD versus WT HFD; # $P < 0.05$ WT HFD versus WT control. (D) Leptin sensitivity test in WT and B1B2KO mice. Consumption was measured after intraperitoneal injection of 40 μ g/day of leptin and compared with basal consumption. Data are expressed as the mean $\pm$ standard error of the mean and represent percentage of food intake. White squares indicate WT, black squares indicate B1B2KO. (E) Insulin sensitivity test after 12 weeks of HFD treatment. Glycemia was measured before and after intraperitoneal injection of insulin 0.01 U/kg at following times: 0, 15, 30, and 60 minutes. Data are expressed as the mean $\pm$ standard error of the mean. White circles indicate WT control, black circles indicate WT HFD, white triangles indicate B1B2KO control, and black triangles indicate B1B2KO HFD. * $P < 0.05$ WT HFD versus all groups.

Abbreviations: HFD, high-fat diet; WT, wild-type.

may compensate by upregulating its expression, which may introduce a confounding factor when interpreting data. To resolve part of this issue, B1B2KO mice were generated and characterized. These double knockout mice are healthy, fertile, normotensive, and protected against endotoxin-induced hypotension.²³

In our present work, we have shown that B1B2KO mice are resistant to HFD-induced obesity, showing less food intake and a smaller weight gain per calorie intake when compared with WT mice. Based on these observations, we analyzed the

mRNA expression of genes in the hypothalamus associated with metabolism. NPY and AgRP, both orexigenic genes, were found to be present at the same levels in B1B2KO and WT mice whether on the control diet or the HFD. Similarly, expression of anorexigenic genes, POMC and CART, was not different between knockout and control mice. Recently, our group found that CART mRNA expression is elevated in the lateral hypothalamic area of B1KO mice.²⁴ Even though B1KO and B1B2KO mice show similarities in some aspects of metabolism, expression of CART seems to be different in

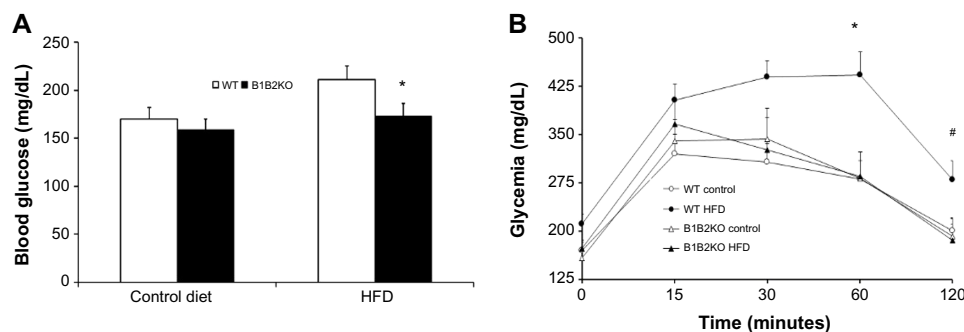


Figure 4 B1B2KO mice show high glucose tolerance.

Notes: (A) Glucose levels after 12 weeks of HFD treatment. White bars indicate WT control and HFD, black bars indicate B1B2KO control and HFD. * $P < 0.05$ WT HFD versus B1B2KO HFD. (B) Glucose tolerance test after 12 weeks of HFD treatment. Glycemia was measured before and after intraperitoneal injection of glucose 1 g/kg at the following times: 0, 15, 30, 60, and 120 minutes. Data are expressed as the mean $\pm$ standard error of the mean. White circles indicate WT control, black circles indicate WT HFD, white triangles indicate B1B2KO control, and black triangles indicate B1B2KO HFD. * $P < 0.05$ WT HFD versus all groups and # $P < 0.05$ WT HFD versus B1B2KO HFD.

Abbreviations: HFD, high-fat diet; WT, wild-type.

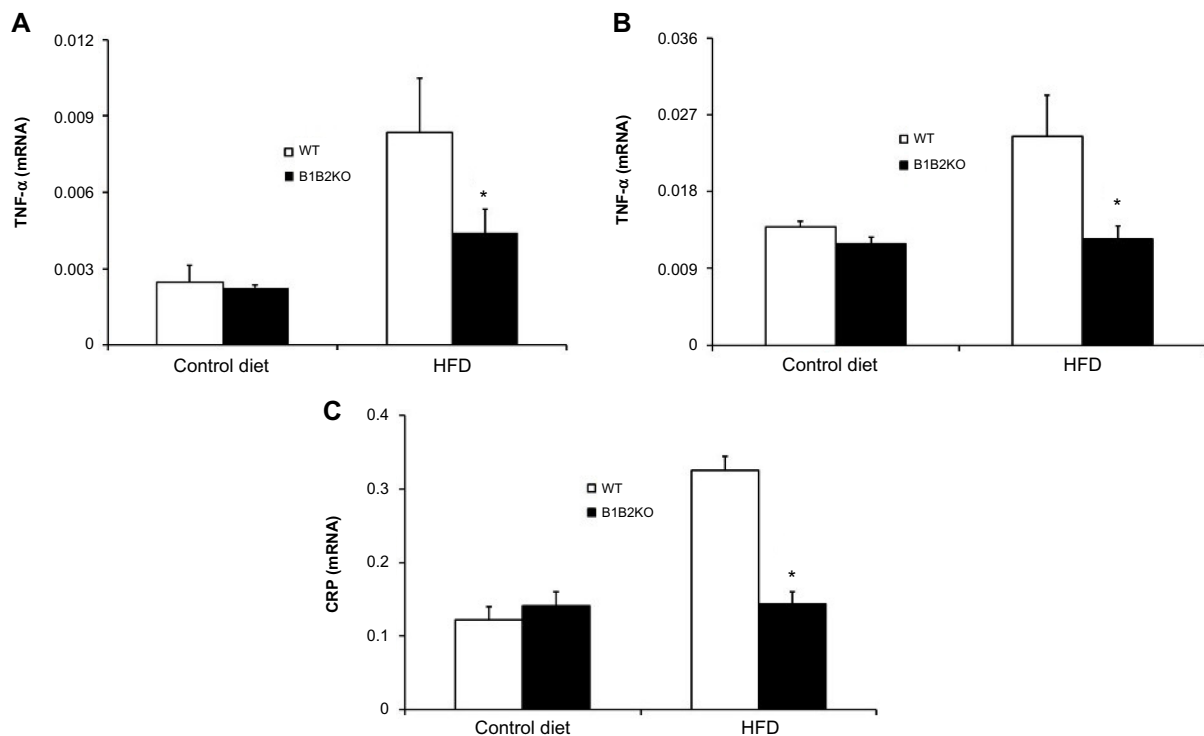


Figure 5 Inflammatory cytokines are downregulated in tissues of B1B2KO mice.

Notes: TNF- α mRNA expression in white adipose tissue (A) and skeletal muscle tissue (B) after 12 weeks of HFD treatment. (C) CRP mRNA expression in the liver after 12 weeks of HFD treatment. Data are expressed as the mean $\pm$ standard error of the mean of the $2^{-\Delta\Delta C_t}$ parameter and represent the relative expression between TNF- α /CRP and β -actin. White bars indicate WT control and HFD, black bars indicate B1B2KO control and HFD. * $P < 0.05$ B1B2KO HFD versus WT HFD.

Abbreviations: HFD, high-fat diet; WT, wild-type; CRP, C-reactive protein; TNF- α , tumor necrosis factor- α .

the two strains. However, it should be taken into account that we analyzed expression of CART in the whole hypothalamus of B1B2KO mice, and not only in the lateral hypothalamic area, which may have contributed to the observed lack of differences. Further, Heijboer et al demonstrated that gene expression for NPY, AgRP, POMC, and CART is not altered in the mouse hypothalamus after treatment with an HFD,²⁵ suggesting that regulation of these genes may be not affected by an HFD in WT or B1B2KO mice. PRCP and MC4R gene expression was also tested, given the role of these genes in metabolism. PRCP is an enzyme associated with inactivation of alpha-melanocortin-stimulating hormone in rodents,²⁶ which in turn binds to MC4R to regulate food intake behavior and energy balance.²⁷ Together with other orexigenic and anorexigenic genes, expression of PRCP and MC4R was unchanged in the B1B2KO mouse hypothalamus after treatment with the control diet or the HFD, indicating that the reduced food consumption observed in these mice was probably related to an alternative metabolic pathway not examined in this study.

Circulating levels of leptin and insulin were also measured in the B1B2KO and WT mice. Decreased serum levels of both hormones were found in the double knockout mice on

the control diet or the HFD. These data suggest hypersensitivity to these hormones, although no changes in insulin and leptin sensitivity were observed in B1B2KO mice. On the other hand, B1B2KO mice showed less glycemia and increased glucose tolerance after treatment with the HFD. We believe the mechanisms underlying these findings may be related to the control of energy intake and its expenditure. Indeed, energy expenditure is increased in B1B2KO mice, as suggested by the feed efficiency data. Studies analyzing the impact of B1 receptor on metabolism have found that B1KO mice benefit from deletion of the B1 receptor, with better leptin and insulin sensitivity.^{16,17} In agreement with this, blockade of the B1 receptor by the specific inhibitor SSR240612 reverted insulin and glucose concentrations to baseline levels in insulin-resistant glucose-fed rats.¹⁹

By analyzing the B2 receptor knockout (B2KO) mouse phenotype, our group has found decreased serum insulin levels and increased glucose uptake after stimulation with insulin (unpublished data). Schweitzer et al²⁸ reported that B2KO mice also have lower serum insulin levels, but show decreased glucose tolerance. Conversely, it has been shown that B2KO mice have higher serum insulin levels and are resistant to the effects of insulin on mediating glucose uptake.²⁹ Taking

these data together, the role of the B2 receptor in glucose uptake remains controversial. We can conclude that disruption of this receptor leads to significant alterations in glucose homeostasis, probably by compensatory mechanisms promoted by other related genes. Indeed, recent gene expression data from microarray analysis of B2KO mice have shown dysregulation of several genes related to pathological pathways involved in endothelial injury, oxidative stress, insulin and leptin metabolism, and the inflammatory process.³⁰ Therefore, we propose that deletion of the B2 receptor in mice regulates expression of not only the B1 receptor,²⁰ but also genes linked to insulin and glucose homeostasis. In agreement with this idea, we believe that disruption of the B1 receptor in mice leads not only to altered expression of B2 receptor,²¹ but also other genes related to control of the energy balance. In this scenario, our double knockout model shows that metabolic parameters measured in the B1B2KO mice were not exclusively dependent on B1 or B2 receptor regulation, indicating that ablation of the B1 and/or B2 receptor might also have a profound impact on several other gene pathways related to disorders of metabolism.

To determine the link between abrogation of KKS and genes involved in metabolic homeostasis, we analyzed gene expression of proteins related to insulin resistance and the inflammatory process. We focused on inflammatory genes because the B1 receptor plays an important role in the immune response in many disorders.^{31–33} Thus, we measured gene expression of TNF- α and CRP, given that these proteins are present in elevated concentrations in insulin-resistant and obese individuals. In these cases, white adipose tissue and skeletal muscle are the main sources of TNF- α ,^{34,35} which is directly related to impaired glucose uptake and insulin resistance mechanisms via activation of nuclear factor kappa B and IKK β signaling.³⁶ Here we found that B1B2KO HFD-treated mice had decreased TNF- α mRNA expression in white adipose tissue and skeletal muscle. Similarly, increased CRP levels, with liver as the main source, are related to impaired insulin resistance.¹⁶ We observed that CRP mRNA expression was decreased in the liver of the B1B2KO mouse after the HFD. Recently, Diaz et al demonstrated that blockade of the B1 receptor is able to reverse inflammatory biomarkers related to insulin resistance in insulin-resistant glucose-fed rats.¹⁹ These findings suggest that reduced mRNA expression of TNF- α and CRP could be related to deletion of the B1 receptor in B1B2KO mice. Absence of the pro-inflammatory B1 receptor could be mandatory to modify the expression of inflammatory genes involved in the pathology of insulin resistance.

Accordingly, in order to observe the effect of pharmacological blockade of the B1 and B2 receptors, we treated WT animals with specific antagonists, ie, R-715 and HOE-140. No change in mRNA expression was observed after pharmacological treatment for leptin, TNF- α , and CRP gene expression in white adipose tissue, skeletal muscle, or the liver. Conversely, leptin mRNA expression was not decreased in white adipose tissue after treatment with specific antagonists, indicating that the decreased baseline leptin levels found in B1B2KO mice probably resulted from more profound and prolonged inactivation of kinin receptors.

In conclusion, we found that a deficiency of B1 and B2 receptors protects against obesity induced by an HFD and promotes a decrease in circulating levels of insulin and leptin. In addition, deletion of both B1 and B2 receptors has specific consequences with regard to mechanisms involving glucose homeostasis, ie, improving glucose tolerance. We propose that compensatory effects, as a consequence of disruption of the KKS, could dysregulate expression of inflammatory genes related to metabolic disorders such as insulin resistance. Therefore, our findings reinforce the role of the KKS in metabolism by showing that B1 and B2 receptor regulation is crucial for glucose tolerance control in mice, and suggest that new drugs focused on the KKS might be effective as therapeutic tools for conditions related to obesity, such as insulin resistance and diabetes.

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Disclosure

The authors report no conflicts of interest in this work.

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