Ketone bodies, stress response, and redox homeostasis

Pedro Rojas-Moralesa, José Pedraza-Chaverri a, Edilia Tapia b,∗

a Departamento de Biología, Facultad de Química, Universidad Nacional Autónoma de México, Ciudad de México, 04510, Mexico
b Departamento de Fisiopatología Cardio-Renal, Instituto Nacional de Cardiología Ignacio Chávez, Ciudad de México, 14080, Mexico

ARTICLE INFO

Keywords:
β-Hydroxybutyrate
Redox homeostasis
Stress response molecule
Ischemia and reperfusion

ABSTRACT

The ketone body β-hydroxybutyrate is no longer viewed simply as a metabolic intermediate, as it regulates a broad range of physiological processes at cellular and systemic levels. Particularly, β-hydroxybutyrate functions as a stress response molecule and orchestrates an antioxidant defense program to maintain redox homeostasis in response to environmental and metabolic challenges, such as ischemia. This property of β-hydroxybutyrate might be key for the beneficial effect of calorie restriction on stress response and disease processes.

1. Introduction

Under well-fed, resting conditions, glucose stands as the main physiological fuel supporting energy metabolism, growth, and reproduction. However, in naturally encounter environments prolonged episodes of limited food availability and also moderate physical activity dramatically alter blood glucose levels and provoke baseline physiological functions to be more dependent on fatty acids metabolism [1]. For example, prolonged fasting and endurance exercise cause blood glucose levels to drop while simultaneously lead to increases in ketone bodies at the expense of liver β-oxidation of adipose tissue-derived fatty acids. Also, hepatic ketogenesis—the synthesis of ketone bodies—is enhanced when feeding a high-fat, low-carbohydrate ketogenic diet or in animals undergoing calorie restriction. In any case, the liver takes up and transforms circulating fatty acids into ketone bodies, mainly β-hydroxybutyrate and acetoacetate, which are then exported out to essentially all of the body's tissues, including the brain, where they are primarily metabolized via mitochondria to produce adenosine triphosphate (ATP) (Fig. 1) [2]. In addition to functioning as alternative energy source during energy stress, it is now well established that β-hydroxybutyrate also moonlights as a signaling molecule [2,3] by for example activating the hydroxycarboxylic acid receptor 2 (HCAR2) [4] and inhibiting the nucleotide binding domain leucine-rich repeat-containing receptor, pyrin domain-containing-3 (NLRP3) inflamasome [5] on immune cells, thus blocking the synthesis of inflammatory intermediates. In addition, lysine modification by β-hydroxybutyrylation on histone and non-histone proteins is emerging as an important regulator of cellular physiology and metabolism [6,7]. Interestingly, β-hydroxybutyrate (and its polymerized form poly-β-hydroxybutyrate) is not solely present in eukarya, but also in bacteria and archaea [8], suggesting important roles for this widespread molecule on cellular homeostasis.

One important aspect of β-hydroxybutyrate that is increasingly being appreciated is its impact on reactive oxygen species (ROS) metabolism. In vitro studies have shown that, in the first place, β-hydroxybutyrate functions as a direct antioxidant for hydroxyl radical (•OH) (Fig. 2A) [9], and inhibits mitochondrial ROS production in stressed neurons by facilitating NADH oxidation (Fig. 2B) [10]. NADH oxidation, and hence the increase of the NAD+/NADH ratio, has important implications in the maintenance of cellular redox homeostasis, for example, through the activation of the protein deacetylases sirtuin 1 (SIRT1) and SIRT3. Convincing evidence has demonstrated that SIRT1 regulates redox status when deacetylates and activates the transcription factor forkhead box O3 (FOXO3), which controls the expression of superoxide dismutase 2 (SOD2) and catalase (CAT). SIRT3, on the other hand, reinforces mitochondrial antioxidant defense by deacetylating and increasing the activity of SOD2. Finally, SIRT3 also activates isocitrate dehydrogenase 2 (IDH2) to increase NADPH, which is used by several antioxidant systems to detoxify ROS [11]. β-Hydroxybutyrate also protects against highly oxidative stress conditions by driving the expression of heme oxygenase 1 (HO-1), SOD2, CAT, nicotinamide adenine dinucleotide phosphate (NADPH) quinone oxidoreductase 1 (NQO1), glucose-6-phosphate dehydrogenase (G6PDH), and glutamate-cysteine ligase (GCL), through the regulation of FOXO1, FOXO3, and nuclear factor-erythroid 2-related factor-2 (NRF2) (Fig. 2) [12–14]. β-
Hydroxybutyrate transcriptionally activates FOXO1 and FOXO3 by promoting histone acetylation due to inhibition of class I histone deacetylases (HDACs). NRF2, on the other hand, seems to be activated by increased amounts of the tricarboxylic acid cycle metabolite fumarate resulting after mitochondrial consumption of β-hydroxybutyrate. However, as an enhanced mitochondrial activity is associated with increased ROS production [11], it cannot be excluded the possibility that the β-hydroxybutyrate-mediated NRF2 activation likely involves mild increases in ROS generation from the mitochondrial electron transport chain. Overall, the effect of β-hydroxybutyrate on redox homeostasis fits well with the ability of calorie restriction to reduce also oxidative stress and damage [15].

In vivo experiments have further illustrated that β-hydroxybutyrate is an anti-ischemic molecule. Ischemia follow by reperfusion supposes a tremendous metabolic challenge to cells and tissues leading to severe organ damage and death, owing in large part to oxidative burst [16]. Administration of β-hydroxybutyrate before or after ischemia and reperfusion results in strong protection of the heart, brain, liver, and the kidney of rodents [13,17–19]. Mechanisms like reduction of oxidative stress, but also mitochondrial protection, suppression of endoplasmic reticulum stress, and enhanced autophagy, as well as the inhibition of cell death processes like necrosis, apoptosis, and pyrroptosis, account for such protective effect of β-hydroxybutyrate (Fig. 3). In this case β-hydroxybutyrate also replicates the multi-systemic protective effect of calorie restriction against ischemia and reperfusion-associated injury [20].

The exact mechanism linking calorie restriction to enhanced stress resistance, and particularly of reducing oxidative stress, is still missing. It has been hypothesized that β-hydroxybutyrate mediates the beneficial effects of calorie restriction [3,21]. This idea seems plausible because in addition to its anti-inflammatory and anti-oxidative properties, β-hydroxybutyrate also leads to lifespan extension [22]. In fact, besides to FOXO1/3, and NRF2, β-hydroxybutyrate actually seems to interact with signaling pathways activated by reduced nutrient intake, like the adenosine monophosphate (AMP)-activated protein kinase (AMPK) [22,23], and SIRT1/3 [22,24], and interferes with both insulin [25] and mammalian target of rapamycin (mTOR) signaling [26]. Thus, β-hydroxybutyrate helps organisms to overcome stressful/pathological
Currently, calorie restriction remains the most powerful tool to increase lifespan in various species [27]. More importantly, calorie restriction reduces risk of chronic diseases in both experimental animals and humans [28]. However, implementing a lifelong calorie restriction regimen in the general population is a bit of a challenge. In light of the contribution of oxidative damage in diverse pathologies [29], increasing physiological amounts of β-hydroxybutyrate might prove useful to promote cellular adaptation to stress and possibly overall human health.

Declaration of competing interest

“The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.”
Acknowledgements

PRM is a doctoral student from Programa de Doctorado en Ciencias Bioquímicas, Universidad Nacional Autónoma de México (UNAM) and receives a scholarship from Consejo Nacional de Ciencia y Tecnología (CONACyT). This study was funded by grants from CONACyT-Mexico (A1-S-7495), Programa de Apoyo a Proyectos de Investigación e Innovación Tecnológica (PAPIIT-Mexico, IN202219), Programa de Apoyo a la Investigación y Posgrado (PAIP-Mexico, 5000-9105), and Fondos del Gasto Directo autorizado a la Subdirección de Investigación Básica, Instituto Nacional de Cardiología Ignacio Chávez.

Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.redox.2019.101395.

References

[1] G.F. Cahill, Fuel metabolism in starvation, Annu. Rev. Nutr. 26 (2006) 1–22.
[2] P. Puchalska, P.A. Crawford, Multi-dimensional roles of ketone bodies in fuel metabolism, signaling, and therapeutics, Cell Metabol. 25 (2017) 262–284.
[3] J.C. Newman, E. Verdin, Ketone bodies as signaling metabolites, Trends Endocrinol. Metab. 25 (2014) 42–52.
[4] M. Rahman, S. Muhammad, M.A. Khan, H. Chen, D.A. Ridder, H. Muller-Fielitz, B. Pokorna, T. Vollbrandt, I. Stolting, R. Nadrowitz, J.G. Okun, S. Offermanns, M. Schwaninger, The beta-hydroxybutyrate receptor HCA2 activates a neuroprotective subset of macrophages, Nat. Commun. 5 (2014) 3944.
[5] Y.H. Youm, K.Y. Nguyen, R.W. Grant, E.L. Goldberg, M. Bodogai, D. Kim, D. D’Agostino, N. Planasvky, C. Lupfer, T.D. Kamneganti, S. Kang, T.L. Horvath, T.M. Fahmy, P.A. Crawford, A. Biragyn, E. Alnemri, V.D. Dixit, The ketone metabolite β-hydroxybutyrate blocks NLPR3 inflammasome-mediated inflammatory disease, Nat. Med. 21 (2015) 263–269.
[6] Z. Xie, D. Zhang, D. Chung, Z. Tang, H. Huang, L. Dai, S. Qi, J. Li, G. Colak, Y. Chen, C. Xia, C. Peng, H. Ruan, M. Kirkey, D. Wang, L.M. Jensen, O.K. Kwon, S. Lee, S.D. Fletcher, M. Tan, D.B. Lombard, K.P. White, H. Zhao, J. Li, R.G. Roeder, X. Yang, Y. Zhao, Metabolic regulation of gene expression by histone lysine β-Hydroxybutyrylation, Mol. Cell 62 (2016) 194–206.
[7] K. Liu, F. Li, Q. Sun, N. Lin, H. Han, K. You, F. Tian, Z. Mao, T. Li, T. Tong, M. Geng, Y. Zhao, W. Gu, W. Zhao, p53 β-hydroxybutyrylation attenuates p53 activity, Cell Death Dis. 10 (2019) 243.
[8] E.N. Dedkova, L.A. Blatter, Role of β-hydroxybutyrate, its polymer poly-β-hydroxybutyrate and inorganic polyphosphate in mammalian health and disease, Front. Physiol. 5 (2014) 260.
[9] M.L. Haces, K. Hernández-Foncea, O.N. Medina-Campos, T. Montiel, J. Pedraza-Chaverri, L. Massieu, Antioxidant capacity contributes to protection of ketone bodies against oxidative damage induced during hypoglycemic conditions, Exp. Neurol. 211 (2008) 85–96.
[10] M. Moolten, P.G. Sullivan, L. Davis, D.Y. Kim, J.M. Rho, Ketones inhibit mitochondrial production of reactive oxygen species production following glutamate excitotoxicity by increasing NADH oxidation, Neuroscience 145 (2007) 256–264.
[11] H. Luo, H.H. Chiang, M. Louw, A. Susanto, D. Chen, Nutrient sensing and the oxidative stress response, Trends Endocrinol. Metab. 28 (2017) 449–460.
[12] T. Shimizu, M.D. Hirschey, J. Newman, W.H. K. Shirakawa, N. Moan, C.A. Grueter, H. Lim, L.R. Saunder, R.D. Stevens, C.B. Newgard, R.V. Farese Jr., R. de Cabo, S. Ulrich, K. Akassoglou, E. Verdin, Suppression of oxidative stress by β-hydroxybutyrate, an endogenous histone deacetylase inhibitor, Science 339 (2013) 211–214.

Fig. 3. The ketone body β-hydroxybutyrate is an anti-ischemic molecule. β-Hydroxybutyrate maintains organ integrity in response to ischemia and reperfusion by suppressing oxidative stress, mitochondrial dysfunction, and inflammation. Also, β-hydroxybutyrate inhibits apoptosis, necrosis, and pyroptosis induced by ischemia and reperfusion. ROS: reactive oxygen species, ER: endoplasmic reticulum.

Fig. 4. The ketone body β-hydroxybutyrate links calorie restriction and stress resistance. β-Hydroxybutyrate, which is produced during calorie restriction, activates adenosine monophosphate-activated protein kinase (AMPK), sirtuins (SIRTs), forkhead box O (FOXO), and nuclear factor-erythroid 2-related factor-2 (NRF2) stress response signaling pathways while inhibiting insulin and mammalian target of rapamycin (mTOR) signaling. IGF-1: insulin growth factor like-1.
T. Miyauchi, Y. Uchida, K. Kadono, H. Hirao, J. Kawase, T. Watanabe, S. Ueda, H. Okajima, H. Terajima, S. Uemoto, Up-regulation of FOXO1 and reduced inflammation by β-hydroxybutyric acid are essential diet restriction benefits against liver injury, Proc. Natl. Acad. Sci. U.S.A. 116 (2019) 13533–13542.

Y. Inuta, T. Imada, R. Hisamura, E. Osinski, S. Nakamura, E. Inagaki, M. Ito, T. Soga, K. Yuiyota, Ketone body 3-hydroxybutyrate mimics calorie restriction via the Nrf2 activator, fumarate, in the retina, Aging Cell 17 (2018) e12699.

R.S. Sobal, R. Weindruch, Oxidative stress, caloric restriction, and aging, Science 273 (1996) 59–63.

D.N. Granger, P.R. Kvietys, Reperfusion injury and reactive oxygen species: the evolution of a concept, Redox Biol. 6 (2015) 524–551.

Y. Yu, Y. Yu, Z. Zhang, Z. Wang, W. An, X. Zhao, Treatment with D-β-hydroxybutyrate protects heart from ischemia/reperfusion injury in mice, Eur. J. Pharmacol. 829 (2018) 121–128.

M. Suzuki, M. Suzuki, K. Sato, S. Debi, T. Sato, A. Matsuura, A. Hiraide, Effect of beta-hydroxybutyrate, a cerebral function improving agent, on cerebral hypoxia, anoxia and ischemia in mice and rats, Jpn. J. Pharmacol. 87 (2001) 143–150.

T. Tajima, A. Yoshifujii, A. Matsui, T. Itoh, K. Uchiyama, T. Kanda, H. Tokuyama, S. Wakino, H. Itoh, β-hydroxybutyrate attenuates renal ischemia-reperfusion injury through its anti-pyroptotic effects, Kidney Int. 95 (2019) 1120–1137.

B.L. Veech, P.C. Bradshaw, K. Clarke, W. Curtis, R. Pavlosky, M.T. King, Ketones bodies mimic the life span extending properties of caloric restriction, JUBMB Life 69 (2017) 305–314.

C. Edwards, J. Canfield, N. Copes, M. Rehan, D. Lips, P.C. Bradshaw, D-beta-hydroxybutyrate extends lifespan in C. elegans, Aging (Albany NY) 6 (2014) 621–644.

H.B. Bae, D.H. Kim, M.H. Park, B. Lee, M.J. Kim, E.K. Lee, K.W. Chung, S.M. Kim, D.S. Im, H.Y. Chung, β-Hydroxybutyrate suppresses inflammasome formation by ameliorating endoplasmic reticulum stress via AMPK activation, Oncotarget 7 (2016) 66444–66454.

J. Yin, P. Han, Z. Tang, Q. Liu, J. Shi, Sirtuin 3 mediates neuroprotection of ketones against ischemic stroke, J. Cereb. Blood Flow Metab. 35 (2015) 1783–1789.

D.H. Kim, M.H. Park, S. Ha, E.J. Bang, Y. Lee, A.K. Lee, J. Lee, B.P. Yu, H.Y. Chung, Anti-inflammatory action of β-hydroxybutyrate via modulation of PGC-1α and FoxO1, mimicking calorie restriction, Aging (Albany NY) 11 (2019) 1283–1304.

Q. Wang, Y. Zhou, P. Rychahou, T.W. Fan, A.N. Lane, H.L. Weiss, B.M. Evers, Ketogenesis contributes to intestinal cell differentiation, Cell Death Differ. 24 (2017) 458–468.

L. Fontana, L. Partridge, V.D. Longo, Extending healthy life span—from yeast to humans, Science 328 (2010) 321–326.

J. Most, V. Tosti, L.M. Redman, L. Fontana, Calorie restriction in humans: an update, Ageing Res. Rev. 39 (2017) 36–45.

I. Liguori, G. Russo, F. Curcio, G. Bulli, L. Aran, D. Della-Morte, G. Gargiulo, G. Testa, F. Cacciarelli, D. Bonaduce, P. Abete, Oxidative stress, aging, and diseases, Clin. Interv. Aging 13 (2018) 757–772.