Isolated branched-chain amino acid intake and muscle protein synthesis in humans: a biochemical review

Consumo isolado de aminoácidos de cadeia ramificada e síntese de proteína muscular em humanos: uma revisão bioquímica

Carina de Sousa Santos¹, Fabrício Expedito Lopes Nascimento²

¹ Universidade Federal dos Vales do Jequitinhonha e Mucuri, Diamantina, MG, Brazil.
² Instituto de Pesquisas, Ensino e Gestão em Saúde, Belo Horizonte, MG, Brazil.

DOI: 10.31744/einstein_journal/2019RB4898

How to cite this article:
Santos CS, Nascimento FE. Isolated branched-chain amino acid intake and muscle protein synthesis in humans: a biochemical review. einstein (São Paulo). 2019;17(3):eRB4898. http://dx.doi.org/10.31744/einstein_journal/2019RB4898

Corresponding author:
Carina de Sousa Santos
Universidade Federal dos Vales do Jequitinhonha e Mucuri
Rodovia MGT 367, km 583, 5.000 – Alto da Jacuba
Zip code: 39100-000 – Diamantina, MG, Brazil
Phone: (55 38) 3532-1200
E-mail: carina.sousasantos@gmail.com

Received on:
Dec 17, 2018

Accepted on:
May 30, 2019

Copyright 2019

This content is licensed under a Creative Commons Attribution 4.0 International License.

ABSTRACT

Alongside a proper diet, ergogenic aids with potential direct and/or indirect physical performance enhancing effects are sought after for improved adaptation to physical training. Nutritional ergogenics include diet composition changes and/or dietary supplementation. Branched-chain amino acids valine, leucine and isoleucine are widely popular among products with ergogenic claims. Their major marketing appeal derives from allegations that branched-chain amino acids intake combined with resistance physical exercise stimulates muscle protein synthesis. Evidence supporting the efficacy of branched-chain amino acids alone for muscle hypertrophy in humans is somewhat equivocal. This brief review describes physiological and biochemical mechanisms underpinning the effects of complete protein source and branched-chain amino acid intake on skeletal muscle growth in the postabsorptive and post-exercise state. Evidence in favor of or against potential anabolic effects of isolated branched-chain amino acid intake on muscle protein synthesis in humans is also examined.

Keywords: Leucine; Valine; Isoleucine; Amino acids, branched-chain; Hypertrophy; Muscle, skeletal

RESUMO

No treinamento físico, buscam-se, além de uma dieta adequada, recursos ergogênicos que possam maximizar direta e/ou indiretamente o desempenho físico. Entre as categorias de recursos ergogênicos, o nutricional compreende a modulação da composição dietética e/ou uso de suplementação. A comercialização dos suplementos de aminoácidos de cadeia ramificada valina, leucina e isoleucina possui muita popularidade entre aqueles com alegação ergogênica. O principal marketing está na afirmação de que o consumo isolado de aminoácidos de cadeia ramificada associado ao exercício físico resistido estimula a síntese de proteína muscular. As evidências da eficácia da ingestão isolada de aminoácidos de cadeia ramificada para a hipertrofia muscular em humanos parecem equivocadas. Nesta breve revisão, apresentamos a compreensão fisiológica e bioquímica de como a ingestão de uma fonte completa de proteína e de aminoácidos de cadeia ramificada afeta o crescimento do músculo esquelético no estado pós-absortivo e pós-exercício. Mostramos também as evidências que suportam ou não a afirmação dos potenciais efeitos anabólicos na síntese de proteína muscular dos aminoácidos de cadeia ramificada quando consumidos isoladamente em humanos.

Descritores: Leucina; Valina; Isoleucina; Aminoácidos de cadeia ramificada; Hipertrofia; Músculo esquelético
INTRODUCTION

Alongside proper diets, ergogenic aids with potential direct and/or indirect physical performance enhancing effects are widely sought after for improved adaptation to physical training. These aids are thought to boost energy production and use, support recovery from exercise, improve body composition and increase resistance to peripheral and central fatigue. Nutritional ergogenic aids include diet composition changes and/or dietary supplementation.(1)

According to the International Society of Sports Nutrition (ISSN), ergogenic supplement claims must be backed up by plausible data and supported by solid evidence of efficacy; these supplements must also be legal and safe.(2)

Branched-chain amino acids (BCAA) enjoy great popularity among allegedly ergogenic supplements. The marketing appeal of these products derives from claims that isolated BCAA intake combined with resistance physical exercise stimulates muscle protein synthesis. This long-standing (over 35 years) claim is based on cellular and animal model studies reporting enhanced anabolic intracellular signaling in response to BCAA intake.(3) However, evidence supporting the efficacy of isolated BCAA intake for muscle hypertrophy in humans is equivocal.

This study set out to review muscle protein synthesis in the postabsorptive state and after resistance physical exercise from a basic biochemistry perspective, and to examine existing evidence in favor of or against claims of potential anabolic effects of isolated BCAA intake on muscle protein synthesis in humans. PubMed®, Latin American and Caribbean on Health Sciences Literature (LILACS) and Scientific Electronic Library Online (SciELO) electronic databases were searched using the following Medical Subject Headings (MeSH) terms: “leucine”, “valine”, “isoleucine”, “branched-chain amino acids”, “muscle protein” and “exercise”. A secondary search was conducted based on the references listed in selected articles.

Muscle protein synthesis

Proteins consist of 20 amino acids (AA) linked by peptide bonds and arranged in different combinations and amounts. Nine out of these 20 AA are thought to be essential (EAA - essential amino acids), i.e., not synthesized in the body and necessarily obtained through diet. The BCAA leucine, isoleucine and valine account for almost 50% of muscle protein EAA. The remaining 11 EAA are classified as nonessential (NEAA, nonessential amino acids), as they can be synthesized in the body.(4,5)

In healthy individuals with normal mobility, dynamic balance between protein degradation and synthesis orchestrates skeletal muscle protein maintenance. In the postabsorptive (i.e., fasting) state, muscle protein degradation exceeds synthesis, leading to net protein loss. In the postprandial state, synthesis exceeds degradation, since intake of some nutrients, such as proteins and carbohydrates, stimulates muscle protein synthesis and insulin release, suppressing degradation.(6) Therefore, muscle hypertrophy requires a positive net protein balance (i.e., muscle protein synthesis in excess of muscle protein degradation).

Physical exercise and nutrient availability are the major drivers of muscle protein synthesis in adult individuals.(7,8) The anabolic effects of nutrients are boosted primarily by transfer and incorporation of AA obtained through the diet into skeletal muscle proteins.(7) These effects are particularly associated with EAA.(9)

The full range of EAA and the 11 NEAA must be present in proper amounts for muscle protein synthesis. Therefore, muscle protein synthesis is limited by lack or low availability of any of the EAA, whereas lack of NEAA can be offset by increased de novo synthesis.(10)

In the postprandial state, within approximately 30 to 45 minutes of consumption of a protein-rich meal (average time required for digestion, absorption and transport of AA to the systemic circulation), EAA availability increases and muscle protein synthesis rates exceed muscle protein degradation rates, inducing an anabolic state that peaks between 1.5 and 3 hours after meal.(10)

Aminoacidemia-induced muscle protein synthesis is transient. In the postabsorptive state (i.e., 2 to 3 hours after meal), plasma EAA levels drop below postprandial levels if no more dietary protein is consumed. In these circumstances, plasma EAA level maintenance (and hence protein turnover) relies on protein breakdown in skeletal muscles, the major body protein reservoir.(11) The impact of factors such as protein amount and quality, protein intake distribution throughout day and physical exercise on the balance between protein degradation and synthesis must be emphasized.

In humans, muscle protein degradation exceeds synthesis by approximately 30% in the postabsorptive state, since 25% of EAA released into the plasma are captured by other tissues and 5% are oxidized in the muscle. The remaining 70% are reutilized for sustained muscle protein synthesis.(12) Hence, muscle protein degradation always exceeds muscle protein synthesis in the postabsorptive state due to muscle protein catabolism and catabolic conditions determined by lack of dietary EAA intake.
BCAA and muscle protein synthesis in the postabsorptive state

The hypothesis that isolated BCAA intake in the postabsorptive state optimizes muscle protein synthesis is based on the premise that exogenous intake decreases the 30% degradation rate by increasing EAA availability for synthesis – instead of oxidation or release into the plasma. This is thought to be partly due to the AA leucine, as leucine alone is able to induce a muscle protein synthesis response via activation of the mechanistic target of rapamycin complex 1 (mTORC1), a vital cell growth regulator.

Louard et al.,(14,15) tested this hypothesis in humans submitted to overnight fasting. In their study, the effects of intravenous BCAA infusion for 3 and 16 hours on muscle protein synthesis and degradation were investigated. Both infusion protocols increased plasma BCAA levels, whereas plasma levels of other EAA decreased. Muscle protein degradation and synthesis also decreased by the same degree, but the synthesis/degradation balance remained negative. The fact that muscle protein degradation was mitigated by isolated intake of three out of 11 EAA, but remained higher than muscle protein synthesis, suggested the catabolic state prevailed in order to release other EAA required for synthesis. It seems therefore plausible to assume BCAA intake alone cannot create an anabolic state leading to muscle protein synthesis in excess of degradation, at least in theory.

Another important question is whether anabolic pathway activation and increased muscle protein synthesis are separate events. Rising insulin levels are a potent anabolic signaling pathway activator, but are not associated with enhanced muscle protein synthesis in the absence of EAA.(16) In contrast, intake of small amounts (3g) of EAA stimulates synthesis regardless of anabolic signaling pathway activation.(17) Should muscle protein synthesis be limited by activation of factors triggering pathway initiating, increases in plasma EAA levels, however small, would not have this effect. These findings demonstrated muscle protein synthesis in humans is limited by availability of the full range of EAA rather than anabolic signaling pathway activation.(3)

BCAA and post-exercise muscle protein synthesis

Combination of physical exercise (resistance physical exercise in particular) with protein intake maximizes and prolongs muscle protein synthesis stimulation for approximately 24 hours post-workout due to increased tissue sensitivity to anabolic properties of AA.

The effects of leucine on human muscle protein synthesis were investigated by Churchward-Venne et al.(20) Participants of that study (young, fit men) consumed 25g of whey protein enriched with 3g of leucine (enough to induce maximal muscle protein synthesis stimulation after resistance physical exercise)(21) or one quarter of that dose after one resistance physical exercise session. Suboptimal and optimal doses of leucine-enriched whey protein similarly increased muscle protein synthesis for 1 to 3 hours post-workout; however, only the optimal (25g) whey protein dose was able to sustain increased synthesis up to 5 hours.

The muscle protein synthesis enhancing potential of higher leucine doses was investigated by the same research group.(22) Participants of that study (young, fit men) consumed leucine-enriched whey protein after one session of resistance physical exercise, as follows: 25g of whey protein containing 3g of leucine; one quarter of that whey protein dose containing 0.75g, 3g or 5g of leucine, or 5g leucine plus similar amounts of isoleucine and valine (3g of each). All treatments equally increased muscle protein synthesis for 1.5 hours post-workout. However, only the suboptimal dose of whey protein containing 5g of leucine was as effective as 25g of whey protein in sustaining increased muscle protein synthesis for 4.5 hours post resistance physical exercise.

Potential reasons why the suboptimal whey protein dose containing 5g of leucine and high amounts of valine and isoleucine failed to sustain muscle protein synthesis were also investigated. It was argued that BCAA may compete for the same carrier in intestinal and muscle cells.(23,24) Therefore, addition of the other two BCAA may have limited sustained muscle protein synthesis due to lower leucine uptake by cells, as suggested by lower blood and intracellular leucine levels in the first hour post-intake in participants receiving this treatment.

Leucine apparently stimulates muscle protein synthesis after resistance physical exercise; however, as with other BCAA, the full range of EAA must be available in levels that will not promote competition for cell carriers or limit their action.

Muscle protein synthesis in response to isolated intake of 5.6g of BCAA or placebo following resistance physical exercise was recently investigated in humans (young, fit men). Within 4 hours, muscle protein synthesis was 22% higher in the treated group as compared to the placebo group.(25) In spite of statistical significance, this increase was six times lower compared to muscle protein synthesis of similar duration in response to consumption of whey protein containing similar amounts of BCAA and EAA.
International organization recommendations
The International Olympic Committee makes no specific recommendations regarding BCAA intake. In contrast, the ISSN does not recommend the use of BCAA to maximize muscle protein synthesis given the limited evidence and inconsistent results in favor of efficacy.\(^{(2,27,28)}\)

Resistance physical exercise increases muscle tissue sensitivity to AA for up to 24 hours post-exercise, regardless of whether protein is consumed prior to, during or 1 to 3 hours post-workout. The major factor is evenly spread protein intake throughout the day, according to individual recommendations. Incremental induction of an anabolic state by repeated resistance physical exercise and diet is what boosts skeletal muscle remodeling and hypertrophy.\(^{(29,30)}\)

For this reason, both organizations support recommendations of combined resistance physical exercise and daily consumption of 1.6g to 2.2g/kg of body mass of complete, high quality EAA-rich protein containing 700 to 3,000mg of leucine in evenly distributed amounts throughout the day (every 3 to 4 hours) for maximal muscle protein synthesis, including one dose after exercise and one close to bed time.\(^{(2,27,28)}\)

\section*{CONCLUSION}
Existing evidence suggests BCAA stimulate muscle protein synthesis following resistance physical exercise. However, in the absence of other essential amino acids, BCAA are not able to sustain maximal synthesis responses. Lack of sustained muscle protein synthesis stimulation equals poor physiologic value. Therefore, a growing body of literature emphasizes that BCAA supplementation alone does not enhance muscle protein synthesis any more than consumption of complete, high quality protein containing the full range of essential amino acids.

\section*{AUTHORS´ INFORMATION}
Santos CS: http://orcid.org/0000-0002-3755-6358
Nascimento FE: http://orcid.org/0000-0001-9612-5163

\section*{REFERENCES}
1. Beck KI, Thomson JS, Swift RJ, von Hurst PR. Role of nutrition in performance enhancement and postexercise recovery. Open Access J Sports Med. 2015;6:259-67. Review.
2. Kerkvliet SI, Wilborn CD, Roberts MD, Smith-Ryan A, Klein CR, Jäger R, et al. ISSN exercise & sports nutrition review update: research & recommendations. J Int Soc Sports Nutr. 2018;15(1):38. Review.
3. Wolfe RR. Branched-chain amino acids and muscle protein synthesis in humans: myth or reality? J Int Soc Sports Nutr. 2017;14(1):30.
4. Moran LA, Horton HR, Scrimgeour KG, Perry MD. Amino acids and the primary structure of proteins. In: Moran LA, Horton HR, Scrimgeour KG, Perry MD, editors. Biochemistry São Paulo: Pearson Education of Brazil; 2013. p. 56-85.
5. Nelson DL, Cox MM. Amino acids, peptides and proteins. In: Nelson DL, Cox MM, editors. Principles of Biochemistry 6 ed. Porto Alegre: Artmed Publisher; 2014. p. 75-114.
6. Rennie MJ, Wackerhage H, Spangenberg EE, Booth FW. Control of the size of the human muscle mass. Annu Rev Physiol. 2004;66(1):799-828. Review.
7. Atherton PJ, Smith K. Muscle protein synthesis in response to nutrition and exercise. J Physiol. 2012;590(5):1049-57. Review.
8. McGlory C, Van Vliet S, Stokes T, Mittendorfer B, Phillips SM. The impact of exercise and nutrition on the regulation of skeletal muscle mass. J Physiol. 2019;597(5):1251-8.
9. Smith K, Barua JM, Watt PW, Scrimgeour CM, Rennie MJ. Flooding with L-[1-\(^{13}\)C]leucine stimulates human muscle protein incorporation of continuously infused L-[1-\(^{13}\)C]valine. Am J Physiol. 1992;262(3 Pt 1):E372-6.
10. Atherton PJ, Etheridge T, Watt PW, Wilkinson D, Selby A, Rankin D, et al. Muscle full effect after oral protein: time-dependent concordance and discordance between human muscle protein synthesis and mTORC1 signaling. Am J Clin Nutr. 2010;92(5):1080-8.
11. Cahill GF Jr, Aoki TT. Starvation and body nitrogen. Trans Am Clin Climatol Assoc. 1971;82:43-51. Review.
12. Biolo G, Gastaldelli A, Zhang XJ, Wolfe RR. Protein synthesis and breakdown in skin and muscle: a leg model of amino acid kinetics. Am J Physiol. 1994;267(3 Pt 1):E467-74.
13. Crozier SJ, Kimball SR, Emmert SW, Anthony JC, Jefferson LS. Oral leucine administration stimulates protein synthesis in rat skeletal muscle. J Nutr. 2005;135(3):376-82.
14. Louard RJ, Barrett EJ, Gelfand RA. Effect of infused branched-chain amino acids on muscle and whole-body amino acid metabolism in man. Clin Sci (Lond). 1990;79(5):457-66.
15. Louard RJ, Barrett EJ, Gelfand RA. Overnight branched-chain amino acid infusion causes sustained suppression of muscle proteolysis. Metabolism. 1995;44(4):424-9.
16. Greenhaff PL, Karagounis LG, Peice N, Simpson EJ, Hazel M, Layfield R, et al. Disassociation between the effects of amino acids and insulin on signaling, ubiquitin ligases, and protein turnover in human muscle. Am J Physiol Endocrinol Metab. 2008;295(3):E585-604.
17. Bukhari SS, Phillips BE, Wilkinson DJ, Limit MC, Rankin D, Mitchell WK, et al. Intake of low-dose leucine-rich essential amino acids stimulates muscle anabolism equivalently to bolus whey protein in older women at rest and after exercise. Am J Physiol Endocrinol Metab. 2015;308(12):E1565-65.
18. Burd NA, West DW, Moore DR, Atherton PJ, Stokes T, Prior T, et al. Enhanced amino acid sensitivity of myofibrillar protein synthesis persists for up to 24 h after exercise resistance in young men. J Nutr. 2011;141(4):568-73.
19. Churchward-Venne TA, Burd NA, Phillips SM. Nutritional regulation of muscle protein synthesis with resistance exercise: strategies to enhance anabolism. Nutr Metab (Lond). 2012;9(1):40.
20. Churchward-Venne TA, Burd NA, Mitchell CJ, West DW, Phillip A, Marcotte GR, et al. Supplementation of a suboptimal protein dose with leucine or essential amino acids: effects on myofibrillar protein synthesis at rest and following resistance exercise in men. J Physiol. 2012;590(11):2761-85.
21. Moore DR, Robinson MJ, Fry JL, Tang JE, Glover EI, Wilkinson SB, et al. Ingested protein dose response of muscle and albumin protein synthesis after resistance exercise in young men. Am J Clin Nutr. 2009;89(1):161-8.
22. Churchward-Venne TA, Breen L, Di Donato DM, Hector AJ, Mitchell CJ, Moore DR, et al. Leucine supplementation of a low-protein mixed macronutrient beverage enhances myofibrillar protein synthesis in young men: a double-blind, randomized trial. Am J Clin Nutr. 2014;99(2):276-86.
23. Szmelcman S, Guggenheim K. Interference between leucine, isoleucine and valine during intestinal absorption. Biochem J. 1966;100(1):7-11.
24. Hyde R, Taylor PM, Hundal HS. Amino acid transporters: roles in amino acid sensing and signalling in animal cells. Biochem J. 2003;373(Pt 1):1-18. Review.

25. Jackman SR, Witard OC, Philp A, Wallis GA, Baar K, Tipton KD. Branched-chain amino acid ingestion stimulates muscle myofibrillar protein synthesis following resistance exercise in humans. Front Physiol. 2017;8:390.

26. Witard OC, Jackman SR, Breen L, Smith K, Selby A, Tipton KD. Myofibrillar muscle protein synthesis rates subsequent to a meal in response to increasing doses of whey protein at rest and after resistance exercise. Am J Clin Nutr. 2014;99(1):86-95.

27. Maughan RJ, Burke LM, Dvorak J, Larson-Meyer DE, Peeling P, Phillips SM, et al. IOC consensus statement: dietary supplements and the high-performance athlete. Int J Sport Nutr Exerc Metab. 2018;28(2):104-25.

28. Jäger R, Kerksick CM, Campbell BJ, Cribb PJ, Wells SD, Skwiat TM, et al. International society of sports nutrition position stand: protein and exercise. J Int Soc Sports Nutr. 2017;14(1):20. Review.

29. Aragon AA, Schoenfeld BJ. Nutrient timing revisited: is there a post-exercise anabolic window? J Int Soc Sports Nutr. 2013;10(1):5.

30. Schoenfeld BJ, Aragon AA, Krieger JW. The effect of protein timing on muscle strength and hypertrophy: a meta-analysis. J Int Soc Sports Nutr. 2013;10(1):53.