Dried bovine placenta improves spermatozoa count in a rat model of male reproductive aging

Surya Agus Prihatno1, Yosua Kristian Adi2, Teguh Budipitojo3, Topas Wicaksono Priyo Jr4 and Yonathan Alvin Maruli Asi Sihotang5

1. Department of Reproduction and Obstetric, Faculty of Veterinary Medicine, Universitas Gadjah Mada, Yogyakarta 55281, Indonesia; 2. Department of Anatomy, Faculty of Veterinary Medicine, Universitas Gadjah Mada, Yogyakarta 55281, Indonesia; 3. Sains Veteriner Magister, Faculty of Veterinary Medicine, Universitas Gadjah Mada, Yogyakarta 55281, Indonesia.

Corresponding author: Surya Agus Prihatno, e-mail: prihatno@ugm.ac.id
Co-authors: YKA: yosua.kristian.a@ugm.ac.id, TB: budipitojo@ugm.ac.id, TWPJ: topas.wicaksono@ugm.ac.id, YAMAS: yonathan.alvin.m@mail.ugm.ac.id

Received: 11-01-2021, Accepted: 06-05-2021, Published online: 21-06-2021

doi: www.doi.org/10.14202/vetworld.2021.1602-1607 How to cite this article: Prihatno SA, Adi YK, Budipitojo T, Priyo Jr TW, Sihotang YAMA (2021) Dried bovine placenta improves spermatozoa count in a rat model of male reproductive aging, Veterinary World, 14(6): 1602-1607.

Abstract

Background and Aim: In the male reproductive system, the aging process can lead to infertility. Recently, placenta and its derivatives have been researched as regenerative agents. This study aimed to describe the basic components of dried bovine placenta powder and its potential effects as a regenerative agent in a rat model of male reproductive aging with D-galactose induction.

Materials and Methods: We divided 15 male Wistar rats, 2 months of age, into three groups: A, the health control group; B, the D-galactose induction group, and C, the D-galactose induction and 10% dried bovine placenta supplementation group. We measured epididymal sperm concentration and testicular weight and volume and analyzed these using one-way analysis of variance.

Results: Dried bovine placenta was rich in nutrients, with 61.98% protein, 21.25±2.07 carbohydrates, 8.58% water, 4.93% ash, and 3.27% fat. The mean epididymal spermatozoa concentration of the rats in Groups A, B, and C was 3026×10^6/mL, 1492.8×10^6/mL, and 2732.5×10^6/mL, respectively. The average total testicle weights were 2.44 g, 2.72 g, and 2.57 g, respectively. The average total testicle volumes were 2.29 cm^3, 2.49 cm^3, and 2.33 cm^3, respectively.

Conclusion: Dried bovine placenta powder is rich in nutrients, especially protein. Supplementation with dried bovine placenta can improve epididymal spermatozoa concentration that is important in fertility.

Keywords: aging, bovine placenta, male reproductive, sperm count, testicular.

Introduction

Aging is a complex biological process involving organic, cellular, and molecular changes. This process is caused chronologically by increasing age but can also be accelerated by environmental factors that cause oxidative stress [1,2]. In the male reproductive system, the aging process can lead to infertility. Semen volume and sperm motility and concentration decline as age increases [3]. A decrease in the quality of bull semen due to aging is a problem at artificial insemination centers in Indonesia [4]. Older bulls often are retained because the cost to replace them is high. Supplementation to improve reproductive performance is an option, but it has not been done because limited research exists on regenerative agents for male livestock reproductive degeneration caused by aging.

Recently, placenta and its derivatives have been studied as regenerative therapy for various degenerative diseases. Placental components can include cord blood cells [5], mesenchymal stem cells [6,7], amniotic membrane [8], placental extract [9], cord blood serum [10], and amniotic fluid [11]. Dried placenta is also rich in protein, fiber, fat, minerals (e.g., sodium, potassium, phosphorus, calcium, iron, magnesium, zinc, copper, and manganese), and hormones (e.g., estradiol, progesterone, testosterone, and growth hormone) [12]. In traditional Chinese medicine, dried human placenta is one ingredient used to treat infertility in humans [13]. Compared to human placenta, bovine placenta is easier to obtain in large quantities. However, dried bovine placenta has never been studied for its nutritional content or its potential effects as a regenerative agent in infertility caused by aging.

Therefore, in this study, we describe the basic components of dried bovine placenta of beef cattle from a traditional farm in Yogyakarta, Indonesia. We analyzed the placenta for its potential effects as a regenerative agent in a rat model of male reproductive degeneration caused by aging. We used D-galactose as an inducting agent for male reproductive aging,
adapting a method previously described by Salman et al. [14].

This study aimed to describe the basic components of dried bovine placenta powder and its potential effects as a regenerative agent in a rat model of male reproductive aging with D-galactose induction.

Materials and Methods

Ethical approval

The study was approved by the Research Ethics Committee, Faculty of Veterinary Medicine, Universitas Gadjah Mada with Ethical Clearance Letter Number: 0023/EC-FKH/Int./2020.

Study period and location

This study was conducted from June to November 2020 at Practical Animal Room and Laboratory of Reproduction and Obstetrics, Faculty of Veterinary Medicine, Universitas Gadjah Mada, Yogyakarta Indonesia.

Dried bovine placenta

Fresh bovine placenta that was expelled <6 h post-parturition was obtained from a traditional farm in Yogyakarta, Indonesia. The placenta was washed using sterile water, cut into a smaller piece, and dried under the sun or an oven until the water content decreased. Then, the placenta was ground into a powder.

Ten grams of bovine placenta powder were analyzed for water, ash, fat, protein, and carbohydrate content at the Universitas Gadjah Mada Center for Biotechnology Studies. For use as a supplementation, the placenta powder was mixed with A.D.II pellets in a ratio of 1:9 (10% bovine placenta in A.D.II pellets).

Animal experiment

In this study, 15 male Wistar rats, aged 2 months, were used. The rats were adapted for a new environment from a breeding place to a practical animal room for 1 week and were maintained in a plastic box with fine air circulation in the room with a 12 h dark-light cycle. The room temperature ranged from 24.3°C to 33.3°C, with the lowest humidity at 39% and the highest at 87%. The rats were fed commercial A.D.II pellets and could drink fresh water ad libitum.

After acclimation, the rats were divided into three groups: Group A, healthy controls; Group B, receiving D-galactose induction; and Group C, receiving D-galactose induction with dried bovine placenta supplementation. The rats in Groups B and C were induced with 3 mg/kg body weight D-galactose orally for 6 weeks. After this induction, the rats in Group C were supplemented with 10% dried bovine placenta mixed in the feed for 30 days. After the treatment, all the rats in all the groups were anesthetized to collect the required samples.

Spermatozoa count

The testicular organ, along with the epididymis, was removed from the scrotum. The epididymis was separated from the testicular organ. As much as, 10 μL of semen from the cauda epididymis was taken using a micropipette. Semen was mixed with 990 μL of NaCl solution in a 1.5 mL tube. The semen solution was diluted again using a leukocyte pipette 101. Spermatozoa counting was done using a Neubauer chamber (Hemocytometer: ASSISTANT®, Germany). The calculation was carried out 3 times, and the results were averaged.

Testes weight and volume

The removed testes were weighed using a digital scale (model: EHA401, CAMRY®, Zhongshan, Guangdong, China). After that, the testes were fixed in Bouin’s solution for 24 h. For preservation, the testes were moved in 70% alcohol solution. The length (l), height (h), and width (w) of the testes were measured using a Vernier caliper (Tricle Brand®, Shanghai, China). Testes volume (V) was measured using the following formula: \( V = \frac{4}{3} \pi l h w \).

Statistical analysis

We analyzed the bovine placenta contents descriptively and the spermatozoa count and testes weight and volume statistically using one-way analysis of variance. P values considered significant at \( p<0.05 \).

Results

Dried bovine placenta contents

The dried bovine placenta powder was rich in nutrients. The highest nutritional content was protein, with an average percentage of 61.98%. Carbohydrates were the second highest nutrient content at 21.25% average content. Water, ash, and fat were the lowest content, with 8.58%, 4.93%, and 3.27% average content, respectively (Table-1).

Spermatozoa count

The concentration of spermatozoa from the caudal epididymis was determined using a Neubauer chamber. The average spermatozoa count in the rats in Groups A, B, and C was 3026×10^3, 2.49 cm^3, and 2.33 cm^3, respectively (Table-3). The average total testicle volume of the rats in Groups A, B, and C was 2.29 cm^3, 2.49 cm^3, and 2.33 cm^3, respectively (Table-4).

Discussion

In mammals, a placenta is an organ for the maternal-fetal exchange of gas and nutrients, hormone secretion, and immune responses [15]. The gross
Table-1: Contents nutrient analysis of dried bovine placenta powder.

| Samples                            | Water (%) | Ash (%) | Fat (%) | Protein (%) | Carbohydrates (by different %) |
|------------------------------------|-----------|---------|---------|-------------|--------------------------------|
| Bovine placenta powder 1           | 8.21      | 5.57    | 3.32    | 62.94       | 19.96                          |
| Bovine placenta powder 2           | 9.05      | 4.14    | 3.43    | 61.49       | 21.89                          |
| Average                            | 8.58±0.41 | 4.93±0.91 | 3.27±0.33 | 61.98±1.83 | 21.25±2.07                    |

Table-2: The number of spermatozoa from the caudal epididymis.

| Rat number | Spermatozoa count (10⁶/mL) |
|------------|--------------------------|
|            | Group A | Group B | Group C |
| Rat 1      | 3466    | 1266    | 2266    |
| Rat 2      | 3200    | 1200    | 3266    |
| Rat 3      | 2600    | 1532    | 2666    |
| Rat 4      | 2932    | 2000    |         |
| Rat 5      | 2932    | 1466    | 2732    |
| Average    | 3026±25 | *1492.8±315 | 2732.5±411 |

*p<0.05

Table-3: Testicle weight of rats in Group A, Group B, and Group C.

| Rat number | Testicle weight (g) |
|------------|---------------------|
|            | Group A | Group B | Group C |
|            | Left testis | Right testis | Left testis | Right testis | Left testis | Right testis |
| Rat 1      | 1.20±0.18 | 1.28±0.20 | 1.40±0.24 | 1.40±0.25 | 1.46±0.26 | 1.40±0.25 |
| Rat 2      | 1.36±0.20 | 1.43±0.21 | 1.50±0.25 | 1.46±0.21 | 1.34±0.21 | 1.34±0.21 |
| Rat 3      | 0.98±0.18 | 1.06±0.20 | 1.24±0.24 | 1.24±0.24 | 1.22±0.22 | 1.22±0.22 |
| Rat 4      | 1.10±0.20 | 1.04±0.20 | 1.40±0.24 | 1.38±0.24 | -         | -         |
| Rat 5      | 1.40±0.36 | 1.36±0.31 | 1.30±0.28 | 1.28±0.28 | 1.18±0.10 | 1.10±0.10 |
| Average    | 1.21±0.23 | 1.23±0.24 | 1.37±0.35 | 1.35±0.35 | 1.30±0.27 | 1.27±0.27 |
|            | 0.18±0.16 | 0.17±0.16 | 0.10±0.09 | 0.09±0.09 | 0.13±0.13 | 0.13±0.13 |
| Total      | 2.44±0.31 | 2.72±0.17 | 2.57±0.22 |         |           |           |

Table-4: Testicle volume of rats in Group A, Group B, and Group C.

| Rat number | Testicle volume (cm³) |
|------------|----------------------|
|            | Group A | Group B | Group C |
|            | Left testis | Right testis | Left testis | Right testis | Left testis | Right testis |
| Rat 1      | 1.25±0.18 | 1.08±0.20 | 1.26±0.31 | 1.31±0.31 | 1.28±0.35 | 1.35±0.35 |
| Rat 2      | 1.30±0.20 | 1.24±0.21 | 1.33±0.31 | 1.31±0.31 | 1.18±0.11 | 1.11±0.11 |
| Rat 3      | 1.00±0.10 | 1.01±0.10 | 1.17±0.13 | 1.13±0.13 | 1.12±0.14 | 1.14±0.14 |
| Rat 4      | 1.00±0.10 | 1.03±0.10 | 1.24±0.29 | -         | -         | -         |
| Rat 5      | 1.26±0.26 | 1.26±0.26 | 1.23±0.19 | 1.19±0.19 | 1.09±0.04 | 1.04±0.04 |
| Average    | 1.16±0.12 | 1.12±0.12 | 1.25±0.25 | 1.25±0.25 | 1.17±0.16 | 1.16±0.16 |
|            | 0.15±0.12 | 0.12±0.12 | 0.06±0.08 | 0.08±0.08 | 0.08±0.13 | 0.13±0.13 |
| Total      | 2.29±0.23 | 2.49±0.12 | 2.33±0.18 |         |           |           |

The anatomic structure of the bovine placenta is classified as the cotyledonary type [16]. The bovine placenta has 75-125 placentomes. Histological classification of the bovine placenta in early pregnancy has been known to be the epitheliocorial type, which becomes progressively syncytiotrophoblastic at the beginning of the second trimester [17]. In this study, we collected placenta that had been expelled <6 h earlier. The bovine fetal membrane is normally expelled within 12 h after delivery of the calf.

Analysis of the dried bovine placenta powder showed that it was still rich in nutrients. This finding is in line with a report by Phuapradit et al. [12] on the nutrients and hormones in a heat dried human placenta, which was rich in protein and minerals. However, the protein hormone concentration was reduced during the steaming and dehydration processes [18]. At least 16 hormones can be detected in human placenta samples processed for encapsulation, and some in concentration can induce physiological effects [19]. In addition, antioxidant peptides can be extracted from human and goat placenta [20, 21]. In this study, 10% dried bovine placenta in commercial feed could increase the protein content from 15.39% to 22.95%.

Our statistical analysis of the spermatozoa count from the epididymal semen showed a significant difference among the groups. Spermatozoa concentration in Groups A and C was significantly different from that in Group B (p<0.05). However, we found no significant difference in the spermatozoa concentration of the rats in Groups A and C (Figure-1). This result confirmed an earlier finding that the administration of D-galactose at a dosage of 3 mg/kg Body Weight orally for 6 weeks can reduce epididymal sperm count [14]. The administration of D-galactose systemically can induce aging in vitro and in vivo and has been widely utilized in anti-aging therapeutic studies [22].

Many researchers have studied the role of oxidative stress in D-galactose-induced animal models [23-29]. In our study, we proved that supplementation with dried bovine placenta can improve spermatozoa count in rats induced into aging by D-galactose. However, the mechanisms involved in the improvement of spermatogenesis are still unclear. During pregnancy, the placenta produces cytokines such as tumor necrosis factor (TNF)-alpha, interleukin (IL)-1, IL-6, and IL-10 [30-32]. Cytokines, which are small proteins secreted and released by cells, have a specific effect on the interaction and communication between cells [33]. Cytokines also have an important role in spermatogenesis.

TNF produces Sertoli cells (Scs) and germ cells within the testis to stimulate and maintain spermatogenesis [34]. IL-1 secreted by Scs and germ cells is known to inhibit Leydig cell steroidogenesis, stimulate Sc transferrin and IL-6 production, and promote Sc proliferation [35]. IL-1 along with IL-6 can regulate...
Sc and spermatogenic cell development [36]. In addition, IL-10 has been proposed to maintain an immune tolerant testicular environment due to its immunosuppressive properties [37].

The placenta produces large amounts of estrogen during human and non-human primate pregnancy, especially in late gestation [38]. In males, estradiol also is essential for modulating libido, erectile function, and spermatogenesis, although inappropriate estradiol levels can lead to a decrease in testicular size and sperm production [39]. However, cytokines and hormone concentrations in dried bovine placenta need to be investigated further.

Statistical analysis of testicle weight and volume of rats in Groups A, B, and C showed no significant differences, although the average sperm concentration from the cauda epididymal among the groups was significantly different. The average testicle weight and volume of the rats in Group B were slightly higher than those (Figures-2 and 3) of the rats in Groups A and C. Similar results have been previously reported by Amini et al. [40], indicating that no significant differences were found in the testicular weight and volume of rats fed a sesame seed diet although the mean cell number in the left epididymis was significantly different. Trautwein et al. [41] reported that testicle volume and weight did not influence the quality of the epididymal sperm of cats. Other reports have stated that some factors such as age, season, environment, and body size can influence testicular weight and volume [42-44]. In our study, we controlled for age, season, and environment.

Conclusion

Bovine placenta is rich in nutrients, especially protein. Supplementation of 10% dried bovine placenta powder in a rat model of male reproductive aging induced by D-galactose can improve epididymal spermatozoa concentration that is important in fertility.

Authors’ Contributions

SAP and YKA: Designed the experimental procedures. TWPJ and YAMAS: Conducted the research work. YKA and TB: Performed sample and statistical analysis. SAP, YKA, and TB: Drafted the manuscript. All authors read and approved the final manuscript.

Acknowledgments

This research was financially supported by Faculty of Veterinary Medicine, Universitas Gadjah Mada, Indonesia, through Faculty Competitive Research (Grant no. 1700/UN1/FKH/HK4/2020).

Competing Interests

The authors declare that they have no competing interests.

Publisher’s Note

Veterinary World remains neutral with regard to jurisdictional claims in published institutional affiliation.

References

1. Sulistyoningrum, E. (2017) D-galactose-induced animal model of male reproductive aging. JKKI, 8(1): 19-27.
2. Liguori, I., Russo, G., Curcio, F., Bulli, G., Aran, L., Della-Morte, D., Gargiulo, G., Testa, G., Cacciarelli, F., Bonaduce, D. and Abete, P. (2018) Oxidative stress, aging, and diseases. Clin. Interv. Aging, 13: 757-772.
3. Kovac, J.R., Addai, J., Smith, R.P., Coward, R.M.,
1. Argiris, A., Ondho, Y.S., Santoso, S.I. and Kurnianto, E. (2018) Effect of age and bulls on fresh semen quality and frozen semen production of Holstein bulls in Indonesia. IOP Conf. Ser. Earth Environ. Sci., 119(1): 012033.

3. Togashi, S.I., Takahashi, N., Iwama, M., Watanabe, S., and consumption.

17. Santos, R.B., Silva, J.M. and Beletti, M.E. (2017) Ultrastructure of bovine placenta during all gestational period. Arch. Bras. Med. Vet. Zootec., 69(6): 1376-1384.

18. Johnson, S.K., Groten, T., Pastuschek, J., Rödel, J., and consumption.

19. Young, S.M., Gryder, L.K., Zava, D., Kimball, D.W. and Pinkert, C.A. (2010) D-galactose effectiveness in modeling aging and therapeutic antioxidant treatment in mice. Rejuvenation Res., 13(6): 729-735.

20. Togashi, S.I., Takahashi, N., Iwama, M., Watanabe, S., Tamagawa, K. and Fukushima, T. (2002) Antioxidative collagen-derived peptides in human-placenta extract. Placenta, 23(6): 497-502.

21. Hou, Y., Zhou, J., Liu, W., Cheng, Y., Wu, L. and Yang, G. (2014) Preparation and characterization of antioxidant peptides from fermented goat placenta. Korean J. Food Sci. Anim. Resour., 34(6): 769-776.

22. Azman, K.F. and Zakaria, R. (2019) D-galactose-induced accelerated aging model: An overview. Biogerontology, 20(6): 763-782.

23. Sun, K., Yang, P., Zhao, R., Bai, Y. and Guo, Z. (2018) Matrine attenuates D-galactose-induced aging-related behavior in mice via inhibition of cellular senescence and oxidative stress. Oxid. Med. Cell. Longev., 2018: 7108604.

24. Liu, H., Zhang, X., Xiao, J., Song, M., Cao, Y., Xiao, H. and Liu, X. (2020) Astaxanthin attenuates d-galactose-induced brain aging in rats by ameliorating oxidative stress, mitochondrial dysfunction, and regulating metabolic markers. Food Funct., 11(5): 4103-4113.

25. Bo-Huy, C., Palec, S., Apaijai, N., Chattipakorn, S.C. and Chattipakorn, N. (2018) Effects of D-galactose-induced aging on the heart and its potential interventions. J. Cell. Mol. Med., 22(3): 1392-1410.

26. Zhao, L., Yang, H., Xu, M., Wang, X., Wang, C., Lian, Y., Mehmood, A. and Dai, H. (2019) Stevia residue extract ameliorates oxidative stress in d-galactose-induced aging mice via Akt/Nrf2/HO-1 pathway. J. Funct. Foods, 52(2): 587-595.

27. Guo, B., Guo, Q., Wang, Z., Shao, J.B., Liu, K., Du, Z.D. and Gong, SS. (2020) D-galactose-induced oxidative stress and mitochondrial dysfunction in the cochlear basilar membrane: An in vitro aging model. Biogerontology, 21(3): 311-323.

28. Parameswaran, K., Irwin, M.H., Steliou, K. and Pinkert, C.A. (2010) D-galactose effectiveness in modeling aging and therapeutic antioxidant treatment in mice.

29. Umbayev, B., Askarova, S., Almabayeva, A., Saliev, T., Masoud, A.R. and Bulanin, D. (2020) Galactose-induced skin aging: The role of oxidative stress. Oxid. Med. Cell. Longev., 2020(8): 7145656.

30. Benyo, D.F., Smarason, A., Redman, C.W.G., Sims, C. and Conrad, K.P. (2001) Expression of inflammatory cytokines in placenta from women with preeclampsia. J. Clin. Endocrinol. Metab., 86(6): 2505-2512.

31. Street, M., Seghini, P., Fieni, S., Ziveri, M., Volta, C., Martorana, D., Viani, I., Gramellini, D. and Bernasconi, S. (2006) Changes in interleukin-6 and IGF system and their relationships in placenta and cord blood in newborns with fetal growth restriction compared with controls. Eur. J. Endocrinol., 155(4): 567-574.

32. Baergen, R., Benirschke, K. and Ulich, T.R. (1994) Cytokine expression in the placenta. The role of interleukin 1 and interleukin 1 receptor antagonist expression in chorioamnionitis and parturition. Arch. Pathol. Lab. Med., 118(1): 52-55.

33. Zhang, J.M. and An, J. (2007) Cytokines, inflammation, and pain. Int. Anesthesiol. Clin., 45(2): 27-37.

34. Loveland, K.L., Klein, M.J., Pueschel, D., Indumathy, S., Bergmann, M., Loveland, B.E., Hedger, M.P. and Schuppe, H.C. (2017) Cytokines in male fertility and reproductive pathologies: Immunoregulation and beyond. Front. Endocrinol., 8: 307.

35. Huleihel, M. and Lunenfeld, E. (2002) Involvement of estradiol in male reproductive function. Mol. Cell. Endocrinol., 187(1-2): 125-132.

36. Hedger, M.P. and Meinhardt, A. (2003) Cytokines and the intratesticular IL-1 system in the regulation of Sertoli cell function. Cell. Mol. Endocrinol., 86(6): 2505-2512.

37. Mittal, S.K. and Roche, P.A. (2015) Suppression of antigen presentation by IL-10.

38. Albrecht, E.D. and Pepe, G.J. (2010) Estrogen regulation of placental angiogenesis and fetal ovarian development during pregnancy. Curr. Opin. Immunol., 22(3): 1392-1410.

39. Schulster, M., Bernie, A.M. and Ramasamy, R. (2016) The role of estradiol in male reproductive function. Asian J. Androl., 18(3): 435-440.

40. Amini, M.J., Hassani, B.H., Nikzad, H., Taherian, A. and Salehi, M. (2013) Effect of diet contains sesame seed on adult wistar rat testis. Int. J. Morphol., 31(1): 197-202.

41. Trautwein, L.G.C., Souza, A.K., Bordo, S., Aguilar, J.B., Paranzini, C.S. and Martins, M.I.M. (2017) Volume and
testicular weight correlation with sperm quality of domestic cats. *Anim. Reprod.*, 14(1): 322.

42. Johnson, L., Petty, C.S. and Neaves, W.B. (1984) Influence of age on sperm production and testicular weights in men. *J. Reprod. Fertil.*, 70(1): 211-218.

43. Mock, A.J. and Frankel, A.I. (1978) A seasonal influence on testes weight and serum gonadotropin levels of the mature male laboratory rat. *Biol. Reprod.*, 18(5): 772-778.

44. Aissanou, S. and Ayad, A. (2020) Influence of age, body weight and season on testicular and epididymis biometrics in donkeys (*Equus asinus*). *Int. J. Morphol.*, 38(5): 1434-1443.

**********