# Morphological and histological characteristics of ovaries from two genetic groups of guinea pigs (*Cavia porcellus*) from South America

Características morfológicas e histológicas de los ovarios de dos grupos genéticos de cuyes (Cavia porcellus) de América del Sur

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#### Abstract

This study aimed to describe morphological and histological characteristics of ovaries of two genetic groups of guinea pigs from Latin America. Ovaries from 20 improved and 20 native guinea pigs were collected after slaughtering. One ovary from each animal was weighed, measured, counted for visible follicles on the ovarian surface, and used for subsequent oocyte collection by the slicing method. Contralateral ovaries were used to prepare histological sections and quantify follicles. Body and ovary weight and ovary length were significantly greater in improved than in native guinea pigs (p<0.01). Ovarian weight was greater in diestrus than in proestrus (p=0.0632) only in improved animals. The

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©Los autores. Este artículo es publicado por la Rev Inv Vet Perú de la Facultad de Medicina Veterinaria, Universidad Nacional Mayor de San Marcos. Este es un artículo de acceso abierto, distribuido bajo los términos de la licencia Creative Commons Atribución 4.0 Internacional (CC BY 4.0) [https:// creativecommons.org/licenses/by/4.0/deed.es] que permite el uso, distribución y reproducción en cualquier medio, siempre que la obra original sea debidamente citada de su fuente original number of primordial, primary, secondary, and total follicles did not differ between genetic groups. The number of antral follicles was significantly greater in improved than in native guinea pigs. The thickness of zona pellucida and oocyte diameter did not differ between groups. The thickness of the zona pellucida was significantly greater in oocytes of category A than B and C in both groups of guinea pigs. In conclusion, ovaries from improved guinea pigs were heavier and longer than those from native animals. The number of antral follicles was greater in improved than native guinea pigs. Zona pellucida thickness and oocyte diameter were similar in both groups of guinea pigs.

Key words: guinea pigs, ovary morphology, follicles, oocytes

#### RESUMEN

El estudio tuvo como objetivo describir las características morfológicas e histológicas de los ovarios de dos grupos genéticos de cuyes de América Latina. Se recolectaron ovarios de 20 cuyes mejorados y 20 nativos después del sacrificio. Se pesó y midió un ovario de cada animal, se contaron los folículos visibles en la superficie ovárica y se colectaron los ovocitos mediante cortes seriados de la corteza ovárica. Los ovarios contralaterales se utilizaron para preparar cortes histológicos y cuantificar los folículos. El peso corporal y la longitud y peso de los ovarios fueron significativamente mayores en los cuyes mejorados que en los nativos (p<0.01). El peso de los ovarios fue mayor en diestro que en proestro (p=0.0632) en los animales mejorados. El número de folículos primordiales, primarios, secundarios y totales no difirió entre los grupos genéticos. El número de folículos antrales fue significativamente mayor en los cuyes mejorados que en los nativos. El grosor de la zona pelúcida y el diámetro de los ovocitos no difirieron entre los grupos. El grosor de la zona pelúcida fue significativamente mayor en los ovocitos de la categoría A que en los de la B y C en ambos grupos de cuyes. En conclusión, los ovarios de los cuyes mejorados fueron más pesados y largos que los de los animales nativos. El número de folículos antrales fue mayor en los cuyes mejorados que en los nativos. El grosor de la zona pelúcida y el diámetro de los ovocitos fueron similares en ambos grupos de cuyes.

Palabras clave: cuy, morfología del ovario, folículos, ovocitos

#### **INTRODUCTION**

Guinea pigs are domestic mammals, hystricomorph rodents of the Caviidae family, autochthonous to the Andes region (Perú, Ecuador, Bolivia, and Colombia) (Lord *et al.*, 2020). They had a close relationship with the pre-Inca people as a source of food high in protein and low in fat and associated with traditions that persist to this day (Avilés *et al.*, 2014a). These are emblematic animals for their diverse utility. In addition to the role of a pet, guinea pigs are used as an animal model for biomedical research (Dyson *et al.*, 2012; Matsuzawa *et al.*, 2018; Morrison *et al.*, 2018; Sterusky *et al.*, 2020). These mammals also play an important role as a high-quality food source (Lammers *et al.*, 2009; Sánchez-Macías *et al.*, 2018), contributing to the family income of several Latin American countries (Avilés *et al.*, 2014a), and as a source of excellent quality compost (Aliaga *et al.*, 2009).

In Ecuador and other Latin American countries, there are three production systems of guinea pigs: family-traditional, familycommercial, and commercial, being the first the most prominent (Avilés *et al.*, 2014b). In general, these production systems manage two genetic groups: native or improved. The former are small, rustic animals and undemanding in terms of food quality, and the latter are considerably larger and more productive and are the result of a process of genetic improvement (Chauca, 1995).

Guinea pigs are non-seasonal breeders and spontaneous ovulators, with a gestation length allowing several calvings by year and several offspring per calving (Lammers et al., 2009). The length of the estrous cycle is approximately 16 days (13-22 days) (Shi et al., 1999; Shomer et al., 2015). Ovulation rate is  $2.8 \pm 0.9$  and  $3.6 \pm 0.9$  for Peru and Andean breeds (Araníbar and Echevarría, 2014). The gestation period is an average 68-day length (59-72 days) (Shomer et al., 2015). Litter size varied according to the genetic group:  $2.4 \pm$ 0.22 in two local native strains (Azuay y Cañar) and  $3.4 \pm 0.32$  in Peru breed females (Cedillo and Quizhpi, 2017), however litter size varies according to parity (Fernández, 2019). The weight of the ovary varied throughout the estrous cycle between 90 and 125 mg, and number of follicles in both ovaries averaged  $101 \pm 9$  (60-105 follicles) (Labhsetwar and Diamond, 1970). Follicular growth occurs in two waves (Bland, 1980; Hamilton and Tam, 1990); the first ending in atresia between days 9-12 and the second wave culminating in ovulation at the end of the estrous cycle (Hamilton and Tam, 1990).

Characterizing the reproductive aspects of both genetic groups is an important step for further plans on *in vitro* production and cryopreservation of guinea pig embryos. Information on morphological and histological features of native and improved guinea pig ovaries is scarce or non-existent. Accordingly, this study aimed to characterize morphologically and histologically the ovary of two genetic groups, native and improved, of guinea pigs from Latin America Andes.

#### MATERIALS AND METHODS

#### **Study Location and Animal Management**

This research was conducted at the Faculty of Agricultural Sciences, University of Cuenca, Ecuador. Forty adult female guinea pigs were used, 20 from native («Azuay and Cañar» native genetic lines) and 20 from improved genetic groups («Auquicuy and Macabeo» Ecuadorian lines), clinically healthy, 1 to 3 calving, with a minimum adult weight of at least 80%, reproductively active and not pregnant.

Animals were handled according to procedures approved by the Veterinary Science Committee, Faculty of Agricultural Sciences, University of Cuenca, and the research was performed following chapter 7.8 of the Terrestrial Animal Health Code -2019© OIE (07/08/2019), regarding the protection of animals used in scientific experiments. All animals were housed under uniform handling and feeding conditions. They were fasted for a period of 12-14 h overnight and then were slaughtered by cervical neck dislocation, one of the euthanasia methods approved by the American Veterinary Medical Association (AVMA, 2020).

# Vaginal Cytology

Immediately after slaughter, vaginal smears were obtained to determine the stage of the estrous cycle following the criteria described by Stockard and Papanicolau (1917) for Cavia porcellus. Smears were taken from the proximal third of the vaginal epithelium by inserting a wet sterile swab roughly 2 cm into the vagina and twice rotating it against the vaginal wall. Each swab was rotated onto a microscope slide, and each smear was air-dried and fixed with ethyl alcohol to preserve cellular morphology. Vaginal smears were stained with Wright® staining (JQWRG-1 K0-00, Quimical, Ecuador), and observed under a microscope (Olympus CX31, Germany) at x10, 20 and 40 magnifications.

### **Ovary Collection and Evaluation**

To account for the effect of anatomical location, the right or left ovaries of the first guinea pig incorporated into the study was randomly assigned to study histological and morphological characteristics respectively and then alternated successively in subsequent animals so that each aspect studied in each genetic group consisted of 10 left and 10 right ovaries. Accordingly, the ovaries selected for morphology and oocyte micrometry were immersed in physiological solution at 37 °C and transported to the laboratory for weighing, measuring, and quantifying the follicles visible on the ovarian surface. Ovaries to describe the follicular population were cut in half and immersed in refrigerated Bouin fixative solution for six hours (Osman, 1985).

The major and minor diameter of each ovary was measured with a digital caliper and weighed on an analytical balance (Boeco, BAS 31plus, Germany). All follicles located on the ovarian surface were counted. Immediately, cumulus-oocyte complexes (COCs) were collected by the slicing method and rinsed three times in synthetic oviductal fluid (SOF). Cumulus-oocytes complexes (improved: n=249; native: n=259) were identified and classified into three categories: A) five or more compact layers of cumulus cells surrounding the oocyte and cytoplasm with homogeneous granulation (improved: n=71; native: n=73); B) less than five compact layers of cumulus cells and totally or partially homogeneous cytoplasm (improved: n=87; native: n=101); and C) oocytes partially or totally free of cumulus cells and/or with expanded cumulus cells and/or mostly heterogeneous cytoplasm (improved: n=91; native: n=85). Cumulus-oocytes complexes were exposed to hyaluronidase (Sigma-Aldrich) (1 mg/ml) for 2-5 min and the cumulus cells removed by gentle pipetting. Oocytes were photographed with a digital camera (Exelis AU-60 0-HD, USA) coupled to a phase contrast microscope at x100 magnification (Olympus CX31, Japan). The thickness of the zona pellucida and oocyte diameter (excluding zona pellucida) were measured using the AmScope (v. 3.7) software.

#### **Histological Processing**

After fixation, the ovaries were embedded in paraffin, cut into 5  $\mu$ m sections, and five sections mounted on each microscope slide. Serial sections mounted on each microscope slice were separated from each other by 50  $\mu$ m of ovarian tissue. After deparaffinization with xylene and rehydration through descending grades of alcohol, tissue specimens were stained with hematoxylin and eosin.

To avoid enumerating the same follicle twice or more times, only antral follicles showing an oocyte inside, and secondary and primary follicles with a visible nucleolus were considered. Primordial follicles were quantified even in the absence of a nucleolus but with an easily identifiable nuclear membrane (Bosch et al., 2004). Primordial follicles were identified as structures having one layer of flattened granulosa cells surrounding the oocyte (the presence of at least one flattened cell within the layer was sufficient to include the follicle in this category), primary follicles were those with one layer of cuboidal granulosa cells, secondary follicles had two or more layers of cuboidal granulosa cells, and antral follicles had a fluid-filled antral cavity inside (Hernandez-Fonseca et al., 2005). The gonadosomatic index was determined (GSI =  $[ovary weight / body weight] \times 100$ ). The follicle count reported in this study corresponds to the average number per ovary.

#### **Statistical Analysis**

Variables showing a skewed distribution (as determined by the Shapiro-Wilks test) were Log<sup>10</sup> transformed. The effect of the genetic group on morphological characteristics of ovaries (size and weight) and oocytes (zona pellucida thickness and oocyte diameter), and follicle categories (histological

Table 1.	Body and ovary weight, gonadosomatic index (GSI), ovary size and number of
	follicles in the ovarian surface in native and improved guinea pigs

Genetic group	Weight		Ovary size (mm)		GSI	Follicles in ovarian	
	Animal (g)	Ovary (mg)	Length	Width	(×10 <sup>-2</sup> )	surface	
Improved	$1986.4\pm55.4^{\mathrm{a}}$	$118.3\pm7.7^{\rm a}$	$7.8\pm0.3^{\text{b}}$	$4.7\pm0.2^{\rm a}$	$1.2\pm0.09^{\rm a}$	$35.3\pm2.8^{\rm a}$	
Native	$942.4\pm58.5^{\text{b}}$	$62.3\pm8.1^{\text{b}}$	$6.4\pm0.3^{\text{c}}$	$4.3\pm0.2^{\rm a}$	$1.3\pm0.10^{\rm a}$	$28.7\pm2.9^{\rm a}$	

Different letters in the same column differ, <sup>a-b</sup> p<0.0001; <sup>b-c</sup> p=0.0052 GSI = (ovary weight / body weight)  $\times$  100

sections) were analyzed by analysis of variance using the GLM (General Linear Model) procedure of SAS (Statistical Analysis System, 2012; version 9.4). Ovarian location (right or left), estrous cycle phase (diestrus and proestrus), and the genetic group  $\times$  estrous cycle phase interaction were considered as independent variables. In oocyte morphometry, in addition to the genetic group, the oocyte category was considered as an independent variable. Statistical significance was considered as p<0.05, and p values between 0.051 and 0.1 were considered as tendency. Values are expressed in mean  $\pm$  standard error.

## RESULTS

As expected, the guinea pigs of the improved genetic group had greater body weight and heavier ovaries than the native guinea pigs (p<0.0001). The ovary length in improved animals was statistically greater but the ovary width was quite similar between genetic groups. The GSI did not differ between groups. The follicle count in the ovarian surface was greater but not significant in improved than in native guinea pigs (Table 1).

In native guinea pigs, the ovary length and width, the GSI, and the follicular count were similar in diestrus and proestrus (Table 2). In improved guinea pigs, ovarian weight was greater in diestrus than in proestrus (p=0.0632), while the other variables were similar between phases of the estrous cycle. Ovarian weight was greater in both phases of the estrous cycle in improved than in native guinea pigs (p<0.0001). The ovary length and width were significantly greater only in diestrus of improved guinea pigs compared with the same phase of the estrous cycle in the native group (Table 2).

The number of primordial, primary, secondary, and total follicles did not differ between genetic groups. However, there was a significantly greater number of antral follicles in improved than in native guinea pigs (Table 3). In improved guinea pigs, the number of all categories of follicles did not differ between both phases of the estrous cycle (Table 4). In native guinea pigs, there was a greater number of primordial (p=0.0770, primary (p=0.0091), and total (p=0.0222) follicles in diestrus than in proestrus. Greater number of primary follicles was observed in proestrus in improved than in native guinea pigs (p=0.0802). Also, a significantly greater number of antral follicles was observed in proestrus (p=0.0389) in

	Ovary weight	Ovar (m	y size m)	GSI	Follicles in ovarian surface	
	(ing)	Length	Width	(~10)		
Improved						
Diestrus	$129.9\pm9.1^{\mathtt{a},1}$	$8.2\pm0.4^{\text{a},1}$	$4.8\pm0.2^{\text{a},1}$	$1.3\pm0.11^{\text{a},1}$	$32.4\pm3.5^{\text{a},\ 1}$	
Proestrus	$100.1\pm12.4^{\text{b},1}$	$7.2\pm0.5^{\text{a},1}$	$4.4\pm0.3^{\text{a},1}$	$1.0\pm0.16^{\text{a, 1}}$	$38.8\pm4.7^{\text{a},1}$	
Native						
Diestrus	$56.8\pm11.4^{\mathrm{a},2}$	$6.0\pm0.5^{\text{a},3}$	$3.7\pm0.3^{\text{a},4}$	$1.2\pm0.11^{\text{a, 1}}$	$27.2\pm4.3^{a,\ 1}$	
Proestrus	$68.4 \pm 11.4^{a,2}$	$6.7\pm0.5^{\text{a},1}$	$4.8\pm0.5^{\text{a},1}$	$1.4 \pm 0.16^{a, 5}$	$30.2\pm4.3^{\mathrm{a},1}$	

Table 2.Ovary weight, gonadosomatic index (GSI), ovary size and number of follicles in<br/>the ovarian surface in native and improved guinea pigs, according to the phase of<br/>estrous cycle

Different letters in the same column for each genetic group differ (<sup>a-b</sup> p= 0.0632) Different numbers in the column between the same estrous cycle phase differ (<sup>1-2</sup> p<0.0001; <sup>1-3</sup> p=0.0009; <sup>1-4</sup> p=0.0115; <sup>1-5</sup> p=0.0516)

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Table 3.	Number	of follicles	of native	and imr	proved	guinea	pig	ovaries
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Constia group	Number of follicles					
Genetic group	Primordial	Primary	Secondary	Antral	Total	
Improved	$12.7\pm2.4^{\rm a}$	$15.1\pm2.3^{\text{a}}$	$12.0\pm2.9^{\rm a}$	$6.1\pm0.9^{\text{a}}$	$45.8\pm6.6^{\rm a}$	
Native	$11.1\pm2.3^{a}$	$13.2\pm2.2^{a}$	$14.8\pm2.7^{\rm a}$	$2.7\pm0.9^{\text{b}}$	$41.9\pm 6.3^{a}$	

Different letters in the same column differ (a-b p=0.0158)

 Table 4.
 Number of follicles in native and improved guinea pig ovaries according to the phase of estrous cycle

Genetic	Number of follicles					
group	Primordial	Primary	Secondary	Antral	Total	
Improved						
Diestrus	$14.5\pm2.8^{\mathrm{a},1}$	$14.4\pm2.7^{\text{a},1}$	$12.6\pm3.4^{\text{a},1}$	$6.0\pm1.0^{\text{a},1}$	$47.6\pm7.8^{\mathrm{a},1}$	
Proestrus	$10.9\pm3.8^{\rm a,1}$	$15.8\pm3.7^{\mathrm{a},1}$	$11.3\pm4.7^{\mathrm{a},1}$	$6.3\pm1.4^{\text{a},1}$	$44.0\pm10.7^{\text{a},1}$	
Native						
Diestrus	$15.4\pm3.2^{\mathrm{a},1}$	$19.6\pm3.2^{\text{b},1}$	$18.5\pm4.0^{\text{a},1}$	$3.6\pm1.4^{\text{a},1}$	$57.7\pm9.1^{\text{c},1}$	
Proestrus	$6.8\pm3.2^{\text{b},1}$	$6.8\pm3.2^{\text{c},2}$	$11.0\pm4.0^{\text{a},1}$	$1.8\pm1.5^{\text{a},3}$	$26.1\pm9.1^{\text{d},1}$	

Different letters in the same column for each genetic group differ (a-b p=0.0770; b-c p=0.0091; c-d p=0.0222)

Different numbers in the column between the same estrous cycle phase differ (1-2 p=0.0802; 1-3 p=0.0389)

improved than in native guinea pigs. There was no effect of ovarian location (right or left) on follicle count in either follicle category in both groups of guinea pigs (data not shown).

The proportion of A, B, and C oocytes was similar in improved (28.5, 34.9, and 36.5% respectively) and native (28.1, 38.9, and 32.8% respectively) guinea pigs. The thickness of zona pellucida ( $14.8 \pm 0.18$  and  $15.1 \pm 0.18 \,\mu m$  for improved and native guinea pigs, respectively) and oocyte diameter  $(78.6 \pm 0.32 \text{ and } 78.6 \pm 0.31 \text{ } \mu\text{m} \text{ for improved}$ and native guinea pigs, respectively) were quite similar between genetic groups. The oocyte diameter was statistically similar between oocyte categories in both genetic groups (data not shown). The thickness of the zona pellucida was significantly greater in oocytes of category A than in oocytes of categories B and C in both improved (15.9  $\pm$  $0.34^{\rm a},\ 14.3\ \pm\ 0.30^{\rm b}$  and  $14.1\ \pm\ 0.31^{\rm b}$ respectively; <sup>a-b</sup> p<0.001) and native (15.6  $\pm$  $0.33^{\circ,e}$ ,  $14.7 \pm 0.30^{d,e}$ , and  $15.0 \pm 0.31^{e}$ , respectively; <sup>c-d</sup> p<0.05) guinea pigs.

# DISCUSSION

Statistical analysis showed marked differences in terms of body and ovarian weight and ovarian size between improved and native genetic groups. In general, the morphometric and histological traits studied were quite similar. In some cases, however, variation prevented the numerical differences, which were sometimes notable, from being statistical as well. The zootechnical characteristics of improved and native guinea pigs have been previously described (Chauca, 1995; Aliaga et al., 2009). The improved guinea pigs in this study resulted from many years of genetic improvement in traits related to meat production. The native variety, consisting of smaller and more rustic animals, was not subjected to systematic genetic selection, so their body weight was slightly less than half that of the former. The

weights of 90-day-old nulliparous guinea pigs from the same native population and improved guinea pigs of a different genetic line than the used in this study were reported a 542.0  $\pm$  17.1 and 874.2 $\pm$  27.9 g, respectively (Cedillo and Quizhpi, 2017).

Body and ovarian weights were about twice as great in females from improved than from native guinea pigs. As these two traits had a similar relationship in both genetic groups, the gonadosomatic index was also very similar. This index represents the percentage ratio between the ovarian and the body weight and varied between 0.012 and 0.013% in improved and native guinea pigs, respectively. In other rodent species, the GSI was ~3.5 folds lower in rats (Ajala *et al.*, 2013) and ~4 folds greater in mice (Wu *et al.*, 2015) than in the guinea pigs in this study.

Overall, ovarian morphology, GSI, and the number of follicles on the ovarian surface did not vary between diestrus and proestrus in both genetic groups. However, the ovarian weight in improved guinea pigs was greater in diestrus than in proestrus. The weight and number of follicles in both ovaries did not change significantly between days 1, 6, 11, and 15 of the estrous cycle in nulliparous guinea pigs (Labhsetwar and Diamond, 1970). In that study, the weight of both ovaries varied between 90 and 120 mg, and the follicular count per animal between 70 and 90 follicles across the estrous cycle. In 2-4 month aged cyclic guinea pigs, the average length, width, and volume of ovaries was  $4.9 \pm 0.1$  mm, 3.4 $\pm$  0.1 mm, and 30.5  $\pm$  1.7 mm<sup>3</sup>, respectively (Wang et al., 2019).

Histological examinations showed no difference in the number of each category of follicles between groups of guinea pigs, except for antral follicles whose number was significantly greater in proestrus of the improved guinea pigs. Even though the follicular count in all categories was similar between diestrus and proestrus in improved guinea pigs, the count of primordial and primary follicles and the total number of follicles in native guinea pigs was greater in diestrus than in proestrus. As follicular growth in guinea pigs occurs in two waves (Bland, 1980; Hamilton and Tam, 1990), a large variation in the number of follicles can be observed throughout the estrous cycle (Logothetopoulos et al., 1995). For instance, the number of nonantral follicles (two-layer of granulosa cell) on days 1, 3, 6, 9, 13, and 16 of the estrous cycle was 125.4, 97.3, 110, 87.2, 67, 112.8 follicles respectively (Logothetopoulos et al., 1995). Also, Hamilton and Tam (1990) observed the greatest number of growing follicles on days 0 and 9, and the lowest number on days 6 and 14 of the estrous cycle. Interestingly, the number of follicles around the time of proestrus in the Logothetopoulos et al. (1995) report was considerably lower than the equivalent period to diestrus, in accordance with the present study

The number of follicles (total and per category) in this study is comparable with that reported by Labhsetwar and Diamond (1970). However, Peddie (1980), observed that the total number of antral follicles did not change significantly during the cycle and averaged  $271.9 \pm 37.8$  on day 5,  $232.2 \pm 29.9$  on day 10 and  $221.1 \pm 24.9$  on day 15; however, the maximum size of the follicles varied substantially. Although not specifically stated by Hamilton and Tam (1990), the total number of follicles ranged from 112 to 200 follicles during the estrous cycle of guinea pigs. These discrepancies in the total number of follicles in different studies are mainly due to methods of follicle counting and the number of slices analyzed. In the present study the objective was not to count as many follicles as possible per ovary but to count a sample that would allow the comparison of two genetic groups under the same criteria.

An interesting finding, not previously reported in guinea pigs was to observe the similarity of the zona pellucida thickness and the oocyte diameter between both groups of guinea pigs. It was also interesting to note that although the oocyte diameter did not vary between the oocyte categories in native and improved guinea pigs, category A oocytes (of greater quality) in both groups had a slightly thicker zona pellucida than the other oocyte categories. In cows, a lesser thickness of the zona pellucida was associated with a corpus luteum in the ovary from which the oocyte was collected, and with greater oocyte competence (Argudo *et al.*, 2020). In guinea pigs, however, there is no evidence linking zona pellucida thickness with oocyte competence.

In the current study, the average diameter of oocytes collected mainly from antral follicles was  $78.6 \pm 0.31 \ \mu\text{m}$ . In a coincidence, a study in adult guinea pigs (Dunk in-Hartley strain with body weights between 562 and 750 g) determined an oocyte diameter of  $78.9 \pm 9.3 \ \mu\text{m}$  in antral follicles (Sadeu *et al.*, 2007). A comparative study in several mammalian species reported oocyte diameters from antral follicles of 72.3, 79, 105.1, and 120 \ \mu\)m for mice, hamsters, pigs, and humans respectively (Griffin *et al.*, 2006). In cows, the oocyte diameter in antral follicles was 124.8 \mu\)m (Argudo *et al.*, 2020).

# CONCLUSIONS

Ovaries from improved guinea pigs were heavier and longer than from natives. The gonadosomatic index and the number of follicles visible on the ovarian surface were similar between groups. With few exceptions, the number of follicles of different categories and the total follicles did not vary between genetic groups. There was a great similarity in zona pellucida thickness and oocyte diameter in native and improved guinea pigs. The proportion of class A, B, and C oocytes was similar between genetic groups. The likeness in the ovarian morphological characteristics of these two genetic groups suggests the use of similar reproductive management, with the further purpose of applying assisted reproduction techniques for improving productive and reproductive performance.

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#### REFERENCES

- Ajala OO, Oyeyemi MO, Fayomi AP, Olaniyan D. 2013. Gonadosomatic index of female Wister rats treated with graded concentration of Aloe Vera gel. J Med Plants Res 7: 339-342. doi: 10.5897/JMPR12.333
- 2. Aliaga L, Moncayo R, Rico E, Caycedo A. 2009. Producción de cuyes. Lima, Perú: Fondo Editorial UCSS. 808 p.
- Araníbar E, Echeverría C. 2014. Número de ovulaciones por ciclo estrual en cuyes (*Cavia porcellus*) Andina y Perú. Rev Inv Vet Perú 25: 29-36. doi: 10.15381/rivep.v25i1.8465
- 4. Argudo DE, Tenemaza MA, Merchán SL, Balvoa JA, Méndez MS, Soria ME, Galarza LR, et al. 2020. Intraovarian influence of bovine corpus luteum on oocyte morphometry and developmental competence, embryo production and cryotolerance. Theriogenology 155: 232-239. doi: 10.1016/ j.theriogenology.2020.05.044
- Avilés DF, Martínez AM, Landi V, Delgado JV. 2014a. The guinea pig (Cavia porcellus): an Andean resource of interest as an agricultural food source. Anim Gen Res 55: 87-91. doi: 10.1017/ S2078633614000368.
- Avilés DF, Landi V, Delgado JV, Martínez AM. 2014b. El pueblo ecuatoriano y su relación con el cuy. Actas Iberoame Conserv Anim 4: 38-40.

- [AVMA] American Veterinary Medical Association. 2020. Guidelines for the euthanasia of animals. 2020 ed. [Internet]. Available in: https://www.avma.org/sites/default/files/2020-01/ 2020\_Euthanasia\_Final\_1-15-20.pdf
- Bland KP. 1980. Biphasic follicular growth in the guinea-pig oestrous cycle. J Reprod Fertil 60: 73-76. doi: 10.1530/ jrf.0.0600073
- Bosch P, Hernandez-Fonseca HJ, Miller DM, Wininger JD, Massey JB, Lamb SV, Brackett BG. 2004. Development of antral follicles in cryopreserved cat ovarian tissue transplanted to immunodeficient mice. Theriogenology 61: 581-594. doi: 10.1016/s0093-691x(03)00244-9
- 10. Cedillo J, Quizhpi J. 2017. Caracterización zoométrica, parametría productiva y reproductiva de dos ecotipos de cuy criollo provenientes de la provincia de Azuay y Cañar a través de la conformación de núcleos exsitu y su comparación con una línea mejorada. Tesis de Médico Veterinario Zootecnista. Ecuador: Univ. de Cuenca. 80 p.
- Chauca L. 1995. Producción de cuyes (Cavia porcellus) en los países andinos. World Anim Rev 83: 9-19.
- 12. Dyson RM, Palliser HK, Kelleher MA, Hirst JJ, Wright IM. 2012. The guinea pig as an animal model for studying perinatal changes in microvascular function. Pediatr Res 71: 20-24. doi: 10.1038/pr.2011.9
- 13. Fernández AE. 2019. Factores que influyen en el desempeño reproductivo de cuyes (*Cavia porcellus*) mejorados en una granja comercial del cantón Sigsig de la provincia del Azuay. Tesis de Médico Veterinario Zootecnista. Ecuador: Univ. de Cuenca. 58 p.
- 14. Griffin J, Emery BR, Huang I, Peterson CM, Carrell DT. 2006. Comparative analysis of follicle morphology and oocyte diameter in four mammalian species (mouse, hamster, pig, and human). J Exp Clin Assist Reprod 3: 2. doi: 10.1186/1743-1050-3-2

- **15.** Hamilton GS, Tam WH. 1990. The bimodular development of ovarian follicles in cyclic guinea pigs and differences between antral follicles developed at different phases of the cycle. Can J Zool 68: 1031-1036. doi: 10.1139/z90-150
- 16. Hernandez-Fonseca HJ, Bosch P, Miller DM, Wininger JD, Massey JB, Brackett BG.2005. Time course of follicular development after bovine ovarian tissue transplantation in male nonobese diabetic severe combined immunodeficient mice. Fertil Steril 83(Suppl 1): 1180-1187. doi: 10.1016/ j.fertnstert.2004.07.985
- 17. Labhsetwar AP, Diamond M. 1970. Ovarian changes in the guinea pig during various reproductive stages and steroid treatments. Biol Reprod 2: 53-57. doi: 10.1095/biolreprod2.1.53
- 18. Lammers PJ, Carlson SL, Zdorkowski GA, Honeyman MS. 2009. Reducing food insecurity in developing countries through meat production: the potential of the guinea pig (Cavia porcellus). Renew Agr Food Syst 24: 155-162.
- 19. Logothetopoulos J, Dorrington J, Bailey D, Stratis M. 1995. Dynamics of follicular growth and atresia of large follicles during the ovarian cycle of the guinea pig: fate of the degenerating follicles, a quantitative study. Anat Rec 243: 37-48. doi: 10.1002/ar.1092430106
- 20. Lord E, Collins C, deFrance S, LeFebvre MJ, Pigière F, Eeckhout P, Erauw C, et al. 2020. Ancient DNA of guinea pigs (*Cavia* spp) indicates a probable new center of domestication and pathways of global distribution. Sci Rep 10: 8901. doi: 10.1038/s41598-020-65784-6
- 21. Matsuzawa S, Isobe M, Kurosawa N. 2018. Guinea pig immunoglobulin VH and VL naïve repertoire analysis. PLoS One 13: e0208977. doi: 10.1371/journal.pone.0208977

- 22. Morrison JL, Botting KJ, Darby JRT, David AL, Dyson RM, Gatford KL, Gray C, et al. 2018. Guinea pig models for translation of the developmental origins of health and disease hypothesis into the clinic. J Physiol 596: 5535-5569. doi: 10.1113/JP274948
- 23. Osman P. 1985. Rate and course of atresia during follicular development in the adult cyclic rat. J Reprod Fertil 73: 261-270. doi: 10.1530/jrf.0.0730261
- 24. Peddie MJ. 1980. Follicular development in the adult guinea-pig and responses to human chorionic gonadotrophin. J Endocrinol 85: 9-16. doi: 10.1677/joe.0.0850009
- 25. Sadeu JC, Adriaens I, Cortvrindt R, Smitz J. 2007. Study of folliculogenesis in vivo in guinea pig. Theriogenology 68: 1228-1239. doi: 10.1016/j.theriogenology.2007.08.028
- 26. Sánchez-Macías D, Barba-Maggi L, Morales-delaNuez A, Palmay-Paredes J. 2018. Guinea pig for meat production: a systematic review of factors affecting the production, carcass and meat quality. Meat Sci 143: 165-176. doi: 10.1016/j.meatsci.2018.05.004
- 27. Shi F, Ozawa M, Komura H, Yang P, Trewin AL, Hutz RJ, Watanabe G, Taya K. 1999. Secretion of ovarian inhibin and its physiologic roles in the regulation of follicle-stimulating hormone secretion during the estrous cycle of the female guinea pig. Biol Reprod 60: 78-84. doi: 10.1095/biolreprod60.1.78
- 28. Shomer NH, Holcombe H, Harkness JE. 2015. Biology and diseases of guinea pigs. Lab Anim Med 2015: 247-283.
- 29. Sterusky M, Plevkova J, Grendar M, Buday T. 2020. Female guinea pig model for cough studies and its response to most common tussive substances. Physiol Res 69(Suppl 1): S171-S179. doi: 10.33549/physiolres.934409

- *30. Stockard C, Papanicolaou G. 1917.* The existence of a typical oestrous cycle in the guinea pig - with a study of its histological and physiological changes. Am J Anat 22: 225-283. doi: 10.1002/ aja.1000220204
- *31. Wang J, Liu Z, Sun Q, Xia S, Cui J, Yang L, An L, Zhang J, et al. 2019.* Combined treatment with cysteamine and

leukemia inhibitory factor promotes guinea pig oocyte meiosis *in vitro*. Am J Transl Res 11: 7479-7491.

32. Wu J, Yuan M, Song Y, Sun F, Han X, MC-LR 2015. MC-LR exposure leads to subfertility of female mice and induces oxidative stress in granulosa cells. Toxins 7: 5212-5223. doi: 10.3390/toxins7124872