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MICRORABITS: A FACTORIAL DESIGN TO EVALUATE GENETIC AND MATERNAL EFFECTS ON GROWTH AND FEED EFFICIENCY IN A LINE SELECTED FOR RESIDUAL FEED INTAKE

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MICRORABITS: A FACTORIAL DESIGN TO EVALUATE GENETIC AND MATERNAL EFFECTS ON GROWTH AND FEED EFFICIENCY IN A LINE SELECTED FOR RESIDUAL FEED INTAKE

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ABSTRACT

The aim of this experiment was to evaluate the significance of neonatal environment (ultimately including the microbiota composition) on feed efficiency. For that purpose, rabbits of the G10 line, selected for 10 generations on residual feed intake (RFI), were fostered by does of a non-selected control line G0, and vice versa. In parallel, collaterals were adopted by mothers from their original line. 900 animals were produced in 3 successive batches and raised in individual or collective cages. Traits analyzed in this preliminary study were body weight at weaning (32 days) and at the end of the test (63 days), average daily gain (ADG), feed intake between weaning and 63 days (FI), feed conversion ratio (FCR) and RFI. Line of the rabbit, type of housing and batch were significant effects for all traits. G10 does had a negative effect on FCR (+0.06, \( P = 0.04 \)), irrespective the line of young rabbits. G10 animals are weaker than G0 at 32 days (-82.9 g) and at 63 days (-161 g). They have also a lower ADG (-2.36 g/day), FCR (-0.36), RFI (-548 g/day) and a lower FI (-839 g), illustrating a better feed efficiency.

Key words: feed efficiency, residual feed intake, growth, genetics

INTRODUCTION

Performances of growing rabbits are determined by their genotype and their environment. The effect of maternal environment is particularly important in this species. Improvement of feed efficiency is essential to increase the competitiveness of the rabbit industry but also to reduce the animal excretion, and consequently decrease the environmental impact of the production. It can be achieved in rabbit by selection on residual feed intake (RFI) or on growth under restricted feeding (Drouilhet et al., 2013, 2015). However these selection strategies do not take into account the contribution of gut microbiota improved feed efficiency, although some previous results have demonstrated its relation with digestive efficiency in chicken (Mignon-Grasteau et al., 2015). To further investigate the effects of the animal genotype and maternal environment on feed efficiency, an experiment based on cross fostering between a line selected on RFI and a non-selected control line was performed. Ultimately, it should allow disentangling the effect of animal genetic and dam microbiota transmission on the traits. The objective of this preliminary study was to estimate both host genotype effect and maternal environment effect on growth and feed efficiency in rabbit.

MATERIALS AND METHODS

Animal management

The experimental rabbit populations were issued from the INRA 1001 line (Larzul and De Rochambeau, 2005) and bred in the experimental INRA farm Pôle Expérimentation Cunicole Toulousain (Castanet-Tolosan, France), in accordance with the national regulations for human care and use of animals in agriculture. Two lines were used in this study: the G10 line, selected for 10 generations on RFI and the G0...
line produced from frozen embryos of the ancestor population of the selected line. The 490 G10 and 410 G0 rabbits were produced in 3 batches with a 42 days interval. Within 48 hours following birth, every kit was fostered. In each batch, half of kits was fostered by G0 does and the second half of kits was fostered by G10 does. Does adopted alternatively kits from one line and from the other line in successive batches. Litters of 5 to 7 kits were made up, mixing sires families of kits within fostered litters.

At weaning (32 days), in each batch, 152 kits were placed in individual cages, 48 in digestibility cages and the rest in collective cages of 4 to 5 animals. All animals were fed ad libitum the same commercial pelleted diet until the end of the fattening period (63 days).

**Traits**

Animals were weighed at weaning (BW32) and at 63 days of age (BW63). Individual feed intake (FI) was recorded in individual and digestibility cages and estimated in collective cages by dividing total feed consumption by the number of animals in the cage, taking into account death of animals when occurring prior to the end of the test. Average daily gain (ADG) was obtained by dividing the body weight gain during the test by the number of days of the growing period. Feed conversion ratio (FCR) was calculated as total feed intake divided by the body weight gain.

**Statistical analyses**

The RFI was computed as the residual of the multiple linear regression of total feed intake on average metabolic body weight (average body weight between weaning and end of the test to the power 0.75) to account for maintenance requirements and ADG to account for production requirements (REG procedure; SAS software).

Fixed effects to be accounted for in the statistical analyses were tested using a linear model (GLM procedure, SAS, 2008):

\[
Y_{ijklm} = \mu + kit\ line_i + doe\ line_j + batch_k + housing_l + batch_k \times housing_l + e_{ijklm}, \quad (1)
\]

with \(Y_{ijklm}\) the trait value for animal \(k\), kitline \(i\) the line of the animal (2 levels), doe line \(j\) the line of the foster doe (2 levels), batch \(k\) the batch of the animal (3 levels), housing \(l\) the type of cage in which the animal was raised (3 levels). The only significant interaction between all fixed effects was \(batch_k \times housing_l\) (\(P < 0.05\)), therefore no other interaction was retained in the models.

**RESULTS AND DISCUSSION**

Levels of significance of fixed effects are presented in Table 1. The batch effect and the batch \(\times\) housing interaction, being significant for all traits, are not mentioned in this table.

**Table 1: Level of significance of fixed effects**

<table>
<thead>
<tr>
<th>Trait</th>
<th>Kit line</th>
<th>Foster doe line</th>
<th>Type of housing</th>
</tr>
</thead>
<tbody>
<tr>
<td>Body weight at 32 days</td>
<td>***</td>
<td>ns</td>
<td>/</td>
</tr>
<tr>
<td>Body weight at 63 days</td>
<td>***</td>
<td>ns</td>
<td>***</td>
</tr>
<tr>
<td>Average Daily Gain</td>
<td>***</td>
<td>ns</td>
<td>***</td>
</tr>
<tr>
<td>Feed Conversion Ratio</td>
<td>***</td>
<td>*</td>
<td>***</td>
</tr>
<tr>
<td>Residual Feed Intake</td>
<td>***</td>
<td>ns</td>
<td>***</td>
</tr>
<tr>
<td>Feed Intake</td>
<td>***</td>
<td>ns</td>
<td>***</td>
</tr>
</tbody>
</table>

ns = non significant ;*: \(P<0.05\);**: \(P<0.01\); ***: \(P < 0.001\).

The kit line effect and the type of housing effect were significant for all traits (\(P < 0.001\)). The foster doe effect was significant only for FCR, G10 foster does showing an unfavorable effect (-0.06±0.02). Least square means of the kit line and of the type of housing effects are presented in Table 2. The G10 animals were lighter than G0 rabbits at 32 days (-82.9 g) and at 63 days (-161 g). They also had a lower ADG (-2.36 g/day), FCR (-0.36), RFI (-548 g) and a lower FI (-839 g), illustrating a better feed efficiency. These
results demonstrate that selection on RFI was efficient, as already reported (Drouilhet et al., 2013, 2015). Nguyen et al. (2005) have also reported a successful selection experiment on RFI in pig.

Table 2: Least square means for kit line and type of housing

<table>
<thead>
<tr>
<th>Trait</th>
<th>G0</th>
<th>G10</th>
<th>collective digestibility</th>
<th>individual</th>
</tr>
</thead>
<tbody>
<tr>
<td>BW32 (g)</td>
<td>916 ± 6</td>
<td>833 ± 6</td>
<td>2,436 ± 14</td>
<td>2,596 ± 20</td>
</tr>
<tr>
<td>BW63 (g)</td>
<td>2,624 ± 13</td>
<td>2,463 ± 12</td>
<td>47.77 ± 0.32</td>
<td>51.88 ± 0.25</td>
</tr>
<tr>
<td>ADG (g/day)</td>
<td>51.76 ± 0.28</td>
<td>49.40 ± 0.26</td>
<td>3.14 ± 0.02</td>
<td>2.69 ± 0.03</td>
</tr>
<tr>
<td>FCR</td>
<td>3.02 ± 0.02</td>
<td>2.66 ± 0.02</td>
<td>333 ± 20</td>
<td>-117 ± 29</td>
</tr>
<tr>
<td>RFI (g)</td>
<td>298 ± 18</td>
<td>-250 ± 17</td>
<td>4,850 ± 26</td>
<td>4,628 ± 21</td>
</tr>
<tr>
<td>FI (g)</td>
<td>5,127 ± 23</td>
<td>4,288 ± 21</td>
<td>4,645 ± 38</td>
<td>4,628 ± 21</td>
</tr>
</tbody>
</table>

*a, b* means with different letters are significantly different (*P* < 0.05).

Concerning the type of housing, performances of rabbits raised in individual cages were similar to those raised in digestibility cages. However, rabbits raised in collective cages were lighter at 63 days (-162 g approx.) and presented a lower ADG (-4.21 g/day approx.) than rabbits raised individually. They had also higher FCR, RFI and FI (around +0.45, +464 g et +205 g, respectively). Coulmin et al. (1982) obtained similar results by decreasing the number of rabbit per cage: heavier animals with a higher ADG associated to smaller number of animals per cage, probably due to decreased loss of energy in relation with activity, but they reported no modification of FCR.

Kit and foster doe lines effects are shown in Figure 1. Compared to G0, G10 foster does had an unfavorable effect on FCR, irrespective of the kit line (-0.06). The maternal effect included the permanent environmental effect offered by the doe to the kits, its own genetic effect and its microbiota transmitted to kits. Our results reflect a negative maternal effect of the selected line G10 on feed efficiency. At this stage of the study, it is not possible to identify which component of maternal effect (milk, maternal behavior, microbiota...) was degraded by the selection (Combes et al., 2013). This can be related to negative correlations previously estimated in some studies between direct and maternal effects on production traits in rabbits (David et al., 2015). In conclusion, FCR was strongly influenced by the genotype of the kit (*Δ* = 0.36, *P* < 0.001) and to a lesser extent by the maternal environment (*Δ* = 0.06, *P* < 0.05).

![Figure 1: Kit line and foster doe line effects on feed conversion ratio.*: *P*<0.05 ; ***: *P* < 0.001.](image.png)
CONCLUSIONS

Our results demonstrate that selection on feed efficiency was successful. However maternal effects were degraded by the selection. Further investigations are undergoing to better understand the effect of selection on direct and maternal effects. These investigations include host genotyping and microbiota sequencing.

ACKNOWLEDGEMENTS

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REFERENCES


