Gonadotrophin-responsiveness of granulosa cells from bone morphogenetic protein 15 heterozygous mutant sheep

Kenneth P McNatty, Derek A Heath, Norma L Hudson, Stan Lun1, Jennifer L Juengel2 and Lloyd G Moore2

Victoria University of Wellington, Wellington 6140, New Zealand, 1Mesynthes Ltd, Lower Hutt 5040, New Zealand and 2AgResearch, Invermay Agricultural Centre, Mosgiel 9053, New Zealand

Correspondence should be addressed to K P McNatty; Email: kenneth.mcnatty@vuw.ac.nz

Abstract

The aim of this study was to test the hypothesis that the higher ovulation-rate in ewes heterozygous for a mutation in bone morphogenetic protein 15 (BMP15; FecXI; otherwise known as Inverdale or I C ewes) is due to granulosa cells developing an earlier responsiveness to LH, but not FSH. To address this hypothesis, granulosa cells were recovered from every individual nonatretic antral follicle (>2.5 mm diameter) from I C and wild-type (++) ewes during anoestrus and the luteal and follicular phases and tested for their responsiveness to FSH and human chorionic gonadotrophin (hCG; a surrogate for LH). For the FSH receptor (FSHR) binding study, granulosa cells were harvested in three separate batches from all antral follicles (≥2.5 mm diameter) from I + and ++ ewes. Using a highly-purified ovine FSH preparation, no evidence was found to suggest that I C ewes have a higher ovulation-rate due to enhanced sensitivity of granulosa cells to FSH with respect to cAMP responsiveness or to their FSHR binding characteristics (equilibrium Kd or Bmax). By contrast, a significantly higher proportion of follicles from I + ewes contained granulosa cells responsive to hCG. The higher proportion was due to cells from more small follicles (i.e. >2.5–4.5 mm diameter) developing a response to hCG. It is concluded that the mutation in the BMP15 gene in I + ewes leads to an earlier acquisition of LH responsiveness by granulosa cells in a greater proportion of follicles and this accounts for the small but significantly higher ovulation-rate in these animals.


Introduction

A number of sheep breeds have been identified with point mutations in an oocyte derived growth factor, bone morphogenetic protein 15 (BMP15; Galloway et al. 2000, Hanrahan et al. 2004, Bodin et al. 2007, Martinez-Royo et al. 2008, Monteagudo et al. 2009). In general, animals heterozygous for the mutations in BMP15 have a higher mean ovulation rate than their respective wild-type controls. By contrast, those homozygous for the BMP15 mutation are infertile due to follicular growth being impaired from the primary stage of development. Sheep known as the Hanna line (allele known as FecXH) have a mutation in the BMP15 gene that introduces a stop codon and thus a truncation at the twenty-third amino acid (AA) in the 125 AA sequence of mature BMP15 (Galloway et al. 2000). In Inverdale ewes (allele known as FecXI), the point mutation in the BMP15 gene results in valine being substituted by aspartic acid at AA number 31 of the mature BMP15 protein. Since homozygous Inverdales have an identical ovarian phenotype to that of homozygous Hanna ewes (Galloway et al. 2000), it is reasonable to assume that the overall level of BMP15 or the biological activity of the protein secreted by oocytes is lower in heterozygous Inverdale ewes than in wild-type ewes. Further evidence to support this has been obtained from studies with transfected HEK 293 cells where human BMP15 with the Inverdale mutation was co-expressed with wild-type human GDF9. Secretions of both BMP15 and GDF9 were reduced (see Moore & Shimasaki 2005 for review). However, it seems unlikely that a reduction in both BMP15 and GDF9 is the cause of increased fertility in heterozygous Inverdale ewes since evidence from immunisation studies in wild-type ewes, where antibodies that recognised BMP15 but not GDF9 were generated, suggests that a reduction in secreted bioactive BMP15 alone can lead to increases in ovulation rates and litter size similar to that in heterozygous Inverdale animals (Juengel et al. 2002, McNatty et al. 2006). In vitro studies with rat granulosa cells have shown that BMP15 can inhibit expression of FSH receptor (FSHR) mRNA as well as FSH-induced expression of STAR, P450scc, 3β-hydroxysteroid dehydrogenase, inhibin subunits and LHR (Shimasaki et al. 2004). The evidence from in vitro studies with...
granulosa cells from Inverdale ewes was equivocal with regard to whether BMP15 inhibits the actions of FSH (Shackell et al. 1993). The reason for this was that the FSH preparation used in these studies, namely NIADDK-ovine FSH (oFSH)-17, contained a measurable level of biologically active LH corresponding to around 1.2% of oFSH-S17. It therefore remains to be determined whether the BMP15 mutation in heterozygous Inverdale ewes resulted in a greater proportion of follicles maturing to ovulation by altering the responsiveness of developing follicles to FSH or LH or both. In this study, we wished to test the hypothesis that the higher ovulation rate in heterozygous Inverdale ewes is due to more follicles developing an earlier responsiveness to LH and that the BMP15 mutation did not affect the binding characteristics of the FSHR or the FSH responsiveness of granulosa cells. To test this hypothesis, we examined the effects of FSH at three dose levels and human chorionic gonadotrophin (hCG; as a surrogate for LH) at one dose level to stimulate cAMP synthesis in granulosa cells from all nonatretic ovarian follicles >2.5 mm diameter from both ovaries of each animal. The FSH binding studies were undertaken on pools of granulosa cells from follicles ≥2.5 mm diameter.

Results

**FSH binding characteristics in I+ and ++ ewes**

The calculated mean±S.E.M., \( B_{\text{max}} \) for the I+ (\( n=3 \)) and ++ (\( n=3 \)) animals were 18.9±0.6 and 23.3±2.1 fmol/mg protein respectively and the \( K_d \) were 0.14±0.01 and 0.13±0.01 nM respectively. No significant effects of genotype were observed.

**Ovarian characteristics of the I+ and ++ animals used for the cAMP studies**

These are summarised in Table 1. The mean±S.E.M. ovarian weights between the genotypes at each stage of the reproductive cycle were not different except during the follicular phase (+++I+; \( P<0.05 \)). The mean±S.E.M. numbers of corpora lutea (CL) were higher in I+ compared to +++ genotype during both the luteal and follicular phases. However, at both stages of the cycle, the mean±S.E.M. total weight of luteal tissue per ewe in I+ and +++ genotypes was not significantly different. Irrespective of time of the reproductive cycle, the geometric mean (95% confidence limits) numbers of follicles >2.5 mm diameter in I+ (\( N=41 \) animals) was 5.1 (4.4, 5.9) and 7.5 (6.4, 8.8) in the +++ genotype (\( N=38 \) animals); these values were significantly different (I+<+++; \( P<0.001 \)). Likewise, the mean (95% confidence limits) overall numbers of non-atretic follicles in I+ and +++ animals were 3.5 (2.9, 4.2) and 5.2 (4.3, 6.2) respectively (I+<+++; \( P<0.01 \)). The proportion of follicles that were nonatretic for each genotype were similar (i.e. ~69%) and not different from one another. With respect to time of the reproductive cycle, both the total number of follicles and the number of nonatretic follicles >2.5 mm diameter were significantly higher in +++ compared to I+ animals during anoestrus (Table 1). However, during the luteal or follicular phases, no differences were noted.

**Numbers of follicles with respect to follicular diameter, genotype and time of year**

The geometric mean (95% confidence limits) for numbers of nonatretic follicles with respect to stage of the reproductive cycle, genotype and follicular diameter are shown in Table 2. During anoestrus, there were significant effects of genotype (\( P<0.01 \)) and follicular diameter (\( P<0.01 \)) and there was also a significant genotype by follicular diameter interaction (\( P<0.01 \)). However, during the luteal or follicular phases, there were no significant effects of follicular diameter or genotype and no interactions were noted.

**Effects of FSH on granulosa cell cAMP production with respect to follicular diameter, genotype or stage of reproductive cycle**

When the cAMP responses by granulosa cells over the follicular size ranges >2.5–3.5, >3.5–4.5 and >4.5 mm diameter were examined overall or with

<table>
<thead>
<tr>
<th>Ovarian characteristic</th>
<th>Anoestrus</th>
<th>Luteal phase</th>
<th>Follicular phase</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>I+</td>
<td>+++</td>
<td>I+</td>
</tr>
<tr>
<td>Ovarian weight (g)</td>
<td>1.10±0.16</td>
<td>1.26±0.10</td>
<td>3.66±0.18</td>
</tr>
<tr>
<td>CL no.</td>
<td>3.00±0.06</td>
<td>1.90±0.18</td>
<td>1.39±0.06</td>
</tr>
<tr>
<td>Total follicle no. &gt;2.5 mm diameter</td>
<td>4.1 (3.3, 5.1)</td>
<td>7.4 (5.6, 9.7)</td>
<td>6.3 (5.1, 7.7)</td>
</tr>
<tr>
<td>Non-atretic follicle no. &gt;2.5 mm diameter</td>
<td>2.7 (2.1, 3.5)</td>
<td>6.1 (4.3, 8.7)</td>
<td>4.2 (3.4, 5.1)</td>
</tr>
<tr>
<td>No of ewes</td>
<td>21</td>
<td>18</td>
<td>10</td>
</tr>
</tbody>
</table>

Values for ovarian weight, CL number (no), and total CL weight are means±S.E.M., whereas the values for follicle number (no) are geometric means with 95% confidence limits in brackets. For each row and stage of reproductive cycle: \(^*P<0.01\); \( ^{*P}<0.05 \).
respect to each stage of the reproductive cycle, significant effects of FSH dose \((P<0.01)\) but no effects of follicular diameter, genotype or interactions were noted. The mean overall effects of FSH dose on cAMP production by granulosa cells with respect to genotype are shown in Fig. 1.

The proportions of nonatretic follicles \((>2.5 \text{ mm diameter})\) with granulosa cells capable of producing \(\geq 5 \text{ pmol cAMP/million cells}\) with respect to genotype and stage of the reproductive cycle were also investigated. During anoestrus, luteal and follicular phases, the respective proportions of follicles from \(+\) animals producing \(\geq 5 \text{ pmol cAMP/million granulosa cells}\) in response to FSH were \(23/61 (38\%)\), \(25/44 (57\%)\) and \(31/43 (72\%)\) and for the \(+ +\) ewes they were \(31/102 (30\%)\), \(24/43 (56\%)\) and \(36/51 (71\%)\). The highest proportion of follicles producing \(\geq 5 \text{ pmol cAMP/million granulosa cells}\) was found during the follicular phase and the lowest during anoestrus for both genotypes. However, when each of these data sets was tested by \(\chi^2\) analysis no genotype effects were noted either overall or at any stage of the reproductive cycle.

**Effects of hCG on granulosa cell cAMP production with respect to follicular diameter, genotype or stage of reproductive cycle**

Less than 4\% of \(+\) (4/174) and \(+ +\) (5/152) follicles \(\leq 2.5 \text{ mm in diameter}\) were found to have granulosa cells capable of producing \(\geq 5 \text{ pmol cAMP/million cells}\) (data not shown). Therefore, the hCG responsiveness was examined in more detail from follicles \(>2.5 \text{ mm diameter}\). When the overall effects of hCG on granulosa cell cAMP production were examined with respect to follicular diameter and genotype (stages of the reproductive cycle pooled) significant effects of genotype \((P<0.05)\) and follicular diameter \((P<0.001)\) but no interactions were noted (Fig. 2). During both anoestrus and the luteal, but not follicular phases, significant effects of genotype were noted (anoestrus, \(P<0.05\); luteal phase, \(P<0.02\)). At all reproductive stages, significant effects of follicular diameter on hCG-induced cAMP responsiveness were noted \((P<0.01)\).

The proportions of nonatretic follicles with granulosa cells producing \(\geq 5 \text{ pmol cAMP/million cells}\) with respect to genotype and stage of the reproductive cycle were also investigated. During anoestrus, luteal and follicular phases, the respective proportions of follicles from \(+\) animals producing \(\geq 5 \text{ pmol cAMP/million granulosa cells}\) were \(20/60 (33\%)\), \(23/44 (52\%)\) and \(24/43 (56\%)\) and for the \(+ +\) ewes they were \(18/102 (18\%)\), \(10/43 (23\%)\) and \(24/51 (47\%)\). When each of these data sets were tested by \(\chi^2\) analysis, significant genotype effects were noted overall \((P<0.01)\), during anoestrus \((P<0.01)\) and the luteal phase \((P<0.05)\) but not during the follicular phase.

**Discussion**

The ovulation rates and luteal characteristics of the Inverdale ewes in the present study were similar to those previously reported by Shackell *et al.* (1993). While \(+\) ewes had an average ovulation rate approximately one higher than the wild-types, the total mass of luteal tissues was not different between the genotypes. This arose despite the fact that there were no differences in the number or range of follicular diameters in nonatretic...
follicles present during either the luteal or follicular phases of the oestrous cycle. The only exception noted in follicular activity was observed for the total number and number of nonatretic >2.5–3.5 mm diameter follicles during anoestrus (+ + >1+). Overall, these findings confirm that there is no significant difference in size distribution of nonatretic follicles >2.5 mm diameter favouring 1+ over + + ewes. Previous studies show that treatment of ewes with exogenous FSH can significantly increase the number of nonatretic follicles >4.5 mm diameter without altering the total number of antral follicles (McNatty et al. 1992).

While differences in plasma FSH concentrations have not been reported for 1+ ewes, earlier studies by Shackell et al. (1993) suggest that 1+ animals contained antral follicles with granulosa cells that potentially are more sensitive to FSH. Such an interpretation is consistent with the in vitro results with rat granulosa cells showing that BMP15 can inhibit FSHR mRNA activity (Shimasaki et al. 2004) and with 1+ ewes having reduced levels of BMP15 (Galloway et al. 2000) so that I+ granulosa cells will be more responsive to FSH. However, from a detailed evaluation of the responsiveness of granulosa cells from all individual nonatretic follicles, no genotype effect was observed for FSH responsiveness in granulosa cells. In the present study, a highly purified FSH preparation, devoid of LH contamination, was used and there was no evidence to suggest that the proportion of follicles responding to any FSH dose differed between the genotypes. Indeed, the only effect noted for FSH was that of dose; the higher the dose, the greater the cAMP response. Therefore, it is likely that the genotype effect observed at a high dose of FSH (1000 ng/ml) by Shackell et al. (1993) was due to the LH contamination in the FSH preparation. For example, when 1000 ng/ml of FSH was added to the culture, the LH contamination would have been 12 ng/ml. In addition to the absence of a genotype effect with respect to FSH-induced cAMP synthesis, no genotype differences were noted with respect to the FSH binding characteristics in granulosa cells from follicles ≥2.5 mm diameter. However, it is important to note, when collecting cells for this study, that no attempt was made to distinguish between nonatretic or atretic follicles. Moreover, neither the binding studies nor those examining cAMP responses in individual follicles rule out the possibility that there are effects of the BMP15 mutation on FSH responsiveness in smaller diameter (i.e. ≤2.5 mm) follicles. However, if this was the case, it was not reflected in any differences in cAMP responses of granulosa cells of these small follicles (data not shown) or any differences in the size distributions of antral follicles which might be anticipated if there were differences in sensitivities to FSH. In the Shackell et al. (1993) study, more small antral follicles between 1.0 and 2.5 mm diameter were observed in I+ than in + + ewes but this was not the case in the present study (data not shown).

The key finding from the present study was that 1+ ewes have a greater proportion of nonatretic follicles >2.5 mm diameter with granulosa cells responsive to hCG/LH than in wild-type ewes. Although significant, the effect was small and only observed after screening granulosa cells from all individual non-arectic follicles >2.5 mm diameter, as not all were responsive to hCG. It is likely that this genotype effect would have been masked if granulosa cell from all non-arectic follicles were pooled. It is perhaps not surprising that the effect is small given that the ovulation rate difference between the genotypes is only ~1. It is evident that the overall genotype effect for the hCG/LH-induced cAMP responsiveness was due to a greater proportion of non-arectic follicles between >2.5 and 4.5 mm in 1+ ewes having granulosa cells responding to hCG relative to that in + + animals. With respect to stage of the reproductive cycle, this genotype effect was noted during anoestrus and the luteal phase but not follicular phase. Our interpretation of these data is that during the luteal phase and anoestrus, the 1+ effect influences the number of follicles >2.5 mm diameter capable of responding to hCG/LH, whereas by the follicular phase all the follicles committed to ovulate have been selected so that the probability of observing a genotype difference on a very small population of follicles over a short window of time is low.

It has been shown in rat granulosa cells, that BMP15 inhibits FSH-induced expression of LHR (Shimasaki et al. 2004). The present study showed no difference in the FSHR characteristics nor the cAMP responsiveness of granulosa cells to endogenous FSH between 1+ and + + animals. However, it is possible that FSH and...
BMP15 crosstalk between the two different signalling pathways, downstream of the production of cAMP. Lowering the amount of bioactive BMP15, as is the case in I+ animals, may alter this cross talk resulting in the attainment of the LHR in an increased proportion of follicles from I+ animals.

In summary, from FSH-receptor binding analyses and using a highly purified oFSH preparation to assess granulosa cell cAMP responses from individual follicles, no evidence was found to suggest that heterozygous Inverdale ewes have a higher ovulation rate as a consequence of enhanced follicular sensitivity to FSH. By contrast, using hCG as a surrogate for LH, a higher proportion of follicles in heterozygous Inverdale ewes contained granulosa cells capable of producing high levels of cAMP compared to the wild-type. It is concluded that the mutation in the BMP15 gene in heterozygous Inverdale ewes leads to an earlier acquisition of LH responsiveness in a greater proportion of follicles and this accounts for the small but significantly higher ovulation rate in these animals.

Materials and Methods

Animals

All experiments were performed with approval of the Animal Ethics Committee of the Wallaceville Animal Research Centre in accordance with the Animal Welfare Act Regulations of New Zealand. All animals had access to pasture and water *ad libitum*. The animals in this study were 5–9 year old heterozygous Inverdale ewes derived from mating known carrier Romney rams with control Romney ewes (I+), whereas the wild-types (+ +) were generated by mating control Romney rams and ewes. The time between recovery of ovaries and initiation of follicular dissection was < 30 min.

In the first study, where the aim was to measure FSH binding characteristics in granulosa cells, ovaries of 55 I+ and 52 + + ewes were recovered from the slaughterhouse. In the second experiment, ovaries were recovered from 21 I+ and 18 + + ewes during anoestrus (November–December), 10 I+ and 10 + + ewes from days 10 to 12 of a PGF2α-induced luteal phase during the breeding season (June) and in 10 I+ and 10 + + ewes at 24 h after a PGF2α-induced follicular phase during the breeding season (June–July).

FSH and LH reagents

An in-house highly purified oFSH preparation (oFSH-Wal) was used for all studies herein. The oFSH was purified from ovine pituitary glands using tryazine-dye chromatography, hydrophobic interaction chromatography and gel filtration as described elsewhere (Moore et al. 1997, Fidler et al. 2003). The oFSH preparation was > 90% pure as determined by gel exclusion chromatography, PAGE and HPLC ion exchange chromatography with a bioactivity of 1.4 × USDA-oFSH-19-SIAFP RP2 or 33 000 IU/mg when the second human FSH International Reference Preparation 78/549 was used as a standard in a radioreceptor assay. The level of LH contamination was < 0.002% as determined by bioassay.

The LH preparation used in all studies was the hCG preparation (CR121; 13 450 IU/mg; NICHD, Bethesda, MD, USA).

Ovarian collection for the FSH binding studies

The method used for the recovery of granulosa cells were similar to that described elsewhere (McNatty et al. 1989). Briefly, ovaries were collected and dissected in saline (0.9%) containing 20 mM Hepes buffer (pH 7.4). Thereafter, all follicles ≥ 2.5 mm diameter were dissected and the granulosa cells removed with a platinum loop. The granulosa cells recovered from all isolated and selected follicles were pooled with respect to genotype, centrifuged at 450 g at 4–6 °C for 20 min and the pellets resuspended in Tris buffer (i.e. 0.05 M-Tris–HCl buffer containing 0.02 M-sucrose and 5 mM-MgCl2 (pH 7.5) to a final concentration of 25 × 10⁶ cells/ml. These cell suspensions were then added in 0.2 ml aliquots to assay tubes, capped and stored at −70 °C until the binding studies were undertaken. For each FSHR binding study, 15–20 animals per genotype were required to obtain sufficient number of cells and the collections were replicated for each genotype thrice.

FSHR binding studies

The equilibrium binding studies were similar to those reported by McNatty et al. (1989) except that the in-house highly purified FSH preparation oFSH-Wal was used as the ligand. The incubations of 125I-oFSH (45 μCi/μg) with granulosa cells were performed at 37 °C for 75 min to achieve equilibrium. The maximum bindability of the radiolabelled FSH to the granulosa cell preparations was ∼ 15%. Duplicate measurements were made at each binding point. Non-specific binding was determined using excess unlabelled Gonadotrophin FSH (Paines & Byrne Ltd, Greenford, UK). The amount of specifically bound 125I-oFSH in c.p.m. was calculated by subtracting non-specific binding from the total amount of bound FSH. Woof plots (i.e. free hormone/bound hormone versus free hormone; Keightley & Cressie 1980) were generated by using a standard amount of 125I-oFSH (∼ 3 ng) and increasing amounts of unlabelled oFSH (i.e doubling amounts from 2 to 128 ng). The binding capacity Bmax was derived from the slope of the fitted line (i.e. slope = 1/Bmax) and the equilibrium Ka calculated from the Bmax × intercept.

Recovery and preparation of granulosa cells for cAMP studies and assay

The ovarian weights and numbers and weights of CL were recorded. Thereafter, all follicles > 1.0 mm were dissected at room temperature in DMEM with 20 mM Hepes buffer, 0.2 mM 3-isobutyl-1-methylxanthine (Sigma Chemical Co.) and 0.1% (w/v) BSA (> 97% pure; ImmunoChemical Products Ltd, Auckland, New Zealand). Nonatretic follicles were those defined as having a vascularised theca interna, no debris in
folicular fluid, $\geq 25\%$ of the maximum number of granulosa cells for a given size and an oocyte of healthy appearance (McNatty et al. 1986). An atretic follicle was so defined if one or more of the above criteria were not satisfied. Once recovered, the cells from individual follicles were centrifuged at 450 g at 4–6 °C for 5 min and the pellets resuspended in the aforementioned medium so that a final concentration of 60 000 cells per culture was achieved. As many follicles between $>1.0–2.5$ mm diameter had insufficient granulosa cells for evaluating their cAMP responses to both FSH and hCG, the results from here onwards are described only for individual follicles $>2.5$ mm diameter. The cells from these follicles were then incubated in 48-well culture plates with or without FSH (10, 100 or 1000 ng/ml) or hCG (1000 ng/ml) in a final volume of 600 µl at 37 °C in a water bath for 45 min. Subsequently, the cultures were heated at 80 °C for 15 min. Samples were stored at $-20$ °C until assayed for cAMP. The cAMP assay was similar to that described by Jolly et al. (1997) except that an in-house rabbit anti 0, 2-monosuccinyl-adenosine-3’,5’-cyclic monophosphate antibody was employed and separation of bound from free cAMP with an in-house sheep anti-rabbit second antibody was followed by the addition of 2.5 volumes of 14% (w/v) polyethylene glycol 8000 (Union Carbide Corp., Danbury, CN, USA). The in-house primary antibody to cAMP cross-reacted 9% with dibutyryl cAMP and <0.0014% with cGMP and $\leq 0.0001\%$ with AMP, ADP or ATP. The detection limit was 0.2 pmol/million cells and the intra- and interassay coefficients of variation were both $<9\%$ respectively.

**Statistical procedures**

The proportions of follicles with granulosa cells expressing high ($\geq 5$ pmol/million cells) or low levels of cAMP ($<5$ pmol/million cells) in response to the highest dose of FSH or hCG during anoestrus, the luteal or follicular phase or overall were examined by $\chi^2$ analyses. cAMP responses of cells to FSH or hCG from each ewe were examined with respect to genotype, FSH dose, reproductive status or follicular size and potential interactions by ANOVA after all cAMP data were first normalised by Ln transformation. Where appropriate, the post hoc comparisons were made using the Bonferroni test.

**Declaration of interest**

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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