Temporal expression and cellular localization of a gastrin-releasing peptide-related gene in ovine uterus during the oestrous cycle and pregnancy

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Abstract

Synthesis of both mRNA and peptide for gastrin-releasing peptide (GRP) has been demonstrated in the pregnant endometrium of sheep and women. However, it is not known whether GRP is synthesized in the sheep uterus during the oestrous cycle. Furthermore the cellular site of GRP mRNA synthesis in the uterus has not been determined. Therefore we examined the synthesis of GRP and determined the cellular location of GRP peptide and mRNA in sheep uterus taken at different times during the oestrous cycle (duration 17 days) and pregnancy (duration 145 days). Northern blot analysis of RNA isolated from ovine endometrium revealed low or no GRP mRNA at days 4, 10, 12 and 14 of the oestrous cycle and a 24-fold rise in GRP mRNA (normalized to glyceraldehyde-3-phosphate dehydrogenase (GAPDH) mRNA) between days 14 and 16. A similar pattern was observed during early pregnancy, with a 12-fold rise in GRP mRNA:GAPDH mRNA between days 17 and 20 of pregnancy. Levels of GRP peptide were determined by RIA and found to be low in endometrium isolated at days 4, 10, 12 and 14 of the oestrous cycle (1.0–1.6 pmol/g) and 4 to 5-fold higher at day 16. *In situ* hybridization localized GRP synthesis to the epithelial cells of the uterine glands at day 16 of the oestrous cycle and at days 17, 20, 40 and 50 of pregnancy. At day 140 of pregnancy diffuse hybrid-

ization to cells of the myometrium was also observed. Immunohistochemistry localized GRP peptide to the apical cytoplasm of uterine glandular epithelial cells at day 16 of the oestrous cycle. For samples obtained at day 20 of pregnancy, the area surrounding the glands also showed moderately strong staining. Further staining in the glandular lumen and the stromal tissue surrounding the glands was apparent at day 140 of pregnancy. No GRP immunoreactivity could be detected in the peripheral plasma during the oestrous cycle or the first 20 days of pregnancy. Sizing chromatography of GRP immunoreactivity extracted from endometrial tissue taken at day 10 of the oestrous cycle revealed two peaks that co-eluted with GRP(1–27) and GRP(18–27). However, during luteolysis and oestrus the major peak of GRP immunoreactivity extracted from endometrial tissue was larger than GRP(1–27) and similar to that seen previously in the gravid ovine endometrium. These studies demonstrate that a peptide similar to, but larger than, GRP is a major product of the glandular epithelium of the ovine uterus during the luteal regression phase of the oestrous cycle and post-blastocyst implantation in pregnancy and provide further evidence that GRP-related peptides have important regulatory roles in uterine function.

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Introduction

Gastrin-releasing peptide (GRP) is a mammalian homologue of amphibian bombesin and was named for its ability to stimulate the release of gastrin, hence the traditional view of it as a gastrointestinal regulatory peptide (see Bunnett 1994 for review). However, it has subsequently been shown that GRP is widely expressed in the mammalian brain, lung, pancreas and anterior pituitary

(Bunnett 1994) and more recently in the pregnant uterus of humans and sheep (Fraser *et al.* 1992, 1994, Giraud *et al.* 1993, 1994, Shulkes *et al.* 1996, Whitley *et al.* 1996, Xiao *et al.* 1996a).

Our previous studies have detected GRP immunoreactivity in the uteroplacental tissues (endometrium, myometrium, amnion, chorioallantois) and fluids (foetal lung fluid and urine, amniotic and allantoic fluids) of the pregnant ewe, as well as in the foetal and maternal

circulations throughout the last third of gestation (Giraud *et al.* 1993). Immunohistochemistry revealed GRP-like immunoreactivity in glandular epithelial cells, but not neuroendocrine cells of the pregnant endometrium. These were surprising findings considering the general role of GRP as a neurotransmitter, and suggested a hormonal role for the endometrial GRP-like peptide in foetal development. Northern blot analysis of GRP mRNA indicated that the sites of synthesis were the near-term pregnant endometrium and myometrium but not the amnion or chorioallantois (Shulkes *et al.* 1996). GRP mRNA was approximately 150-fold lower in the non-pregnant endometrium and not detected in the non-pregnant myometrium (Whitley *et al.* 1994). We have noticed a similar pattern of expression in the human uterus (Whitley *et al.* 1996).

The high abundance in the pregnant endometrium coupled with the known mitogenic effects of GRP implies an important growth-promoting role for a GRP-related peptide in the uterus. This peptide also may be involved in implantation of the blastocyst, as we have observed a 10-fold increase in GRP mRNA in the endometrium just before implantation at day 16/17 of pregnancy (Giraud *et al.* 1994). Preliminary characterization of the GRP-related peptide indicated that it is larger than the known forms of GRP including GRP(1–27), suggesting that it is a novel uterus-specific form of GRP (Giraud *et al.* 1996, Xiao *et al.* 1996b).

In sheep, cattle and pigs, the majority of protein production by uterine endometrium and transport of serum-derived proteins into the uterine lumen occurs during the luteal phase of the oestrous cycle, in pregnancy, or in response to exogenous progesterone. In the pregnant ewe, the embryo enters the uterus about 4 days after ovulation (Rowson & Moor 1966). After 8 days, the majority of embryos will have reached the blastocyst stage (characterized by a central cavity and presence of an embryonic disc) with a 1 mm diameter. After the 12th day, a period of rapid growth commences to form an elongated sac 12–33 mm in length. This rapid growth continues and by day 14–15 after fertilization the blastocyst is 70–100 mm in length. The first signs of attachment can be detected around day 16. Conceptus growth in the ewe is regulated in part by the maternal environment, possibly via paracrine secretion of endometrial factors (Lawson *et al.* 1983). These may be secreted into the uterine lumen and act directly on the conceptus or indirectly by controlling the release of secretory products from the uterine glands. Much is still unknown about the many factors released by the ewe endometrium during early pregnancy or about the regulation and interactions of these necessary factors for conceptus and uterine growth. Our previous studies have shown that a GRP-like peptide is expressed in early pregnancy (Giraud *et al.* 1994). However, it is not known whether GRP is expressed by the non-pregnant uterus, and if it is, whether the level of

expression correlates with the stage of the oestrous cycle. The cellular localization of the peptide in early pregnancy and during the cycle is also not known. The objective of the present study was to determine the temporal variation of uterine GRP during the ovine oestrous cycle and compare this with early pregnancy. Northern blot analysis and RIA were used to determine the synthesis, storage and secretion of uterine GRP, and *in situ* hybridization and immunohistochemistry were used to ascertain the cellular sites of synthesis and storage. The nature of the uterine GRP synthesized during the oestrous cycle was characterized by sizing chromatography.

Materials and Methods

Animals

Sheep were prepared by one of two methods. In protocol 1, the oestrous cycles of parous Corriedale ewes were synchronized by treatment for 12 days with intravaginal inserts containing progesterone (CIDR-G; EAZI-breed, Riverina Breeders Pty Ltd, Albury, Vic., Australia) and an injection of the prostaglandin analogue cloprostenol (125 µg Estrumate; Pitman-Moore, North Ryde, NSW, Australia) 1 day before withdrawal of the progesterone insert. Ewes were mated at the subsequent oestrus with either vasectomized or intact rams bearing a marking harness as described previously (Salamonsen *et al.* 1986). On the evening before surgery, animals were penned, and food and water were withheld. On days 4, 10, 12, 14 and 16 after oestrus (day 0) and on days 4, 7, 8, 9, 10, 12, 14, 15, 16, 17 and 20 of pregnancy, the animals (three to six in each group) were anaesthetized, and hysterectomy performed as described previously (Salamonsen *et al.* 1986). Pregnancy was confirmed by the presence of an embryo in the lumen, either removed by flushing with 20 ml saline (days 4–16) or by visualization after hysterectomy (days 17 and 20). Endometrium was collected and either frozen at -70°C before extraction of RNA and peptide or frozen in embedding compound (OCT; Miles Inc, Elkhart, IN, USA) for analysis by *in situ* hybridization and immunohistochemistry.

Blood samples were collected from a separate group of control ewes mated with either vasectomized or intact rams for another study (Vogiagis *et al.* 1997). Frozen plasma samples (-70°C for 10 months) from days 4, 10, 12, 14 ($n=4$) and 16 ($n=3$) of the oestrous cycle and early pregnancy were thawed, 1 ml was removed and EDTA was added to 1 mM to inhibit proteases during the RIA. The samples were stored again at -70°C until RIA 4 days later.

In protocol 2, mature Corriedale ewes with normal oestrous cycles were used during the breeding season and mated with a vasectomized ram. At 10–12 days after mating, luteolysis was induced by injection (i.m.) with cloprostenol (Estrumate). Previous data have shown that

ewes treated in this way normally have a preovulatory luteinizing hormone surge within 40–70 h of a cloprostenol injection (Tsonis *et al.* 1984). Endometrial tissue was taken from sheep culled at day 8 (before cloprostenol injection), 30 h after cloprostenol injection (considered to be equivalent to the luteal regression phase) and 48 h after cloprostenol injection (when ewes are sexually receptive to the ram and therefore in oestrus).

Tissues taken from sheep treated as described in protocol 2 were used for characterization of uterine GRP by sizing chromatography. Tissues taken from sheep treated as described in protocol 1 were used for all other studies. All animal experimentation was approved by the animal ethics committees at the Animal Research Institute, Werribee and Monash Medical Centre, Clayton, Victoria.

Isolation of RNA and Northern blot analysis for GRP mRNA and glyceraldehyde-3-phosphate dehydrogenase (GAPDH) mRNA

Total RNA was isolated as reported previously (Whitley *et al.* 1996) from the endometrial tissues taken on days 4, 10, 12, 14 and 16 of the oestrous cycle and on days 4, 7, 8, 9, 10, 12, 14, 15, 16, 17 and 20 of pregnancy. A 20 µg portion of each was separated by agarose gel electrophoresis before transfer to nylon membranes (Hybond N; Amersham, Melbourne, Vic., Australia). Northern blot analysis for GAPDH mRNA and GRP mRNA using radiolabelled ovine cDNAs as probes was performed essentially as described previously (Giraud *et al.* 1994). GRP mRNA was detected after autoradiography for approximately 24 h using two intensifying screens. GRP mRNA was quantified by laser densitometry (LKB Ultrosan XL Enhanced Laser Densitometer, LKB/AMRAD, Kew, Vic, Australia) and normalized to the amount of GAPDH mRNA.

In situ hybridization

Riboprobe preparation ³²P-labelled antisense and sense cRNA probes were prepared by *in vitro* transcription from cloned ovine GRP cDNA. Approximately 1 µg linearized DNA was included in a reaction mixture consisting of 40 mM Tris-HCl (pH 7.9), 6 mM MgCl₂, 2 mM spermidine, 10 mM NaCl, 10 mM dithiothreitol, 0.5 mM GTP, ATP and CTP, 40 U RNasin (Promega, Madison, WI, USA), 2.8 µM [³²P]UTP (1000–3000 Ci/mmol), 7.2 µM unlabelled UTP and 10 U T3 or T7 RNA polymerase (Promega). After incubation at 37 °C for 1 h, 0.5 µg RNase-free DNase I (Boehringer-Mannheim, Sydney, NSW, Australia) was added and incubation continued for 15 min at 37 °C. The reaction volume was made up to 50 µl, and radiolabelled probe was purified using a ProbeQuant G-50 Micro column (Pharmacia-Amrad, Kew, Vic, Australia) according to the manufacturer's instructions.

Hybridization conditions Sections (5 µm) of uterus frozen in embedding compound (OCT) were thawed on to silane coated glass slides and baked at 50 °C for 2 h. Mounted sections were rehydrated by immersion in 100% ethanol (2 × 2 min), 95% ethanol (1 × 2 min), 70% ethanol (1 × 2 min) and then water (brief rinse). They were then immersed in proteinase K buffer (50 mM Tris-HCl, pH 7.5) (5 min), fresh proteinase K buffer containing 10 µg/ml proteinase K (10 min; 37 °C), PBS containing 2 mg/ml glycine (2 min), PBS (2 × 1 min), 4% paraformaldehyde (10 min), PBS (2 × 1 min), acetylation buffer (0.1 M triethanolamine, 0.25% acetic anhydride) (10 min), PBS (2 × 1 min) and finally dehydrated in 70, 95 and 100% ethanol (20 s each). The sections were then hybridized overnight at 50 °C in the presence of 50 µl hybridization buffer (50% deionized formamide, 5 × SSC (1 × SSC=0.015 M NaCl and 0.015 M sodium citrate, pH 7.0), 0.1 M NaH₂PO₄ (pH 7.0), 10 mM dithiothreitol, 100 µg/ml tRNA, 100 µg/ml single-stranded sheared salmon sperm DNA, 1 × Denhardt's solution) containing 50 000 d.p.m. sense or antisense probe. After hybridization, the slides were rinsed in 50% formamide/5 × SSC (60 min; 50 °C), followed by RNase A digestion buffer (0.3 M NaCl, 10 mM Tris-HCl, pH 8.0, 5 mM EDTA) (3 × 5 min; 37 °C) and then fresh RNase A digestion buffer containing 20 µg/ml RNase A (Sigma) for 30 min at 37 °C. The slides were then washed under conditions of increasing stringency using 2 × SSC and finally 0.5 × SSC (30 min each; 65 °C) and then dehydrated (70, 95 and 100% ethanol, 2 × 1 min each). The slides were exposed to XAR-5 film (Kodak, Coburg, Vic., Australia) for 6 days. After development of the film, the slides were coated with NTB-2 photographic emulsion according to the manufacturer's instructions (Kodak), stored for 7 days and then developed, stained and photographed.

RIA

GRP immunoreactivity was measured by a previously described RIA (Giraud *et al.* 1994) using polyclonal antiserum R40 raised against [Lys³]-bombesin (kindly donated by Professor B Jarrott, Department of Pharmacology, Monash University, Melbourne, Vic, Australia). The RIA uses ¹²⁵I-[Tyr⁴]-bombesin as tracer. The assay diluent was 0.7 ml 25 mM phosphate buffer containing 0.1% BSA and 2 µM NaN₃, to which were added 100 µl label (5000 c.p.m.), 100 µl antibody R40 (1:100 000 final concentration) and 100 µl sample or standard. Tubes were incubated for 48 h at 4 °C, and bound counts were separated from free by the addition of 300 µl of a cocktail consisting of equal volumes of goat anti-serum IgG (plasma diluted 1:60 in assay buffer), non-immune rabbit plasma (1:600) and polyethylene glycol (10%), followed by centrifugation (2000 g; 30 min). Non-specific binding was 1.5 ± 0.09% (n=47). The RIA standard was bombesin (3.9–1000 fmol/ml); 50% inhibition of label binding

occurred at 125 ± 7 fmol/ml ($n=48$). Inter- and intra-assay coefficients of variation were 14 and 4% respectively. The antiserum is directed to the amidated C-terminus and cross-reacts 100% with bombesin, [Tyr⁴]-bombesin, GRP(18–27) and GRP(1–27) and less than 0.1% with neuromedin B and substance P. The detection limit per assay tube was 4 fmol/ml.

Immunohistochemistry

Sections (10 µm) of whole uterus, fixed for 4–6 h in fresh 4% paraformaldehyde and embedded in paraffin were stained using an indirect peroxidase technique and an avidin–biotin kit (Vectalabs, Burlingame, CA, USA). The primary antiserum AB 11–6 was raised in a rabbit immunized with [Lys⁴]-bombesin coupled to key-hole limpet haemocyanin conjugate and was used at 1:500–1:1000 dilution. Method specificity was tested by incubation with normal rabbit serum, and antiserum specificity by incubation with bombesin or GRP(18–27).

Chromatography

Endometrial tissue samples were taken from sheep ($n=3$) that had been prepared as described in protocol 2. Endometrium (11 g) was boiled in 10 volumes 3% glacial acetic acid at 95 °C for 5 min, centrifuged and the supernatant subjected to solid phase extraction on C₁₈ Sep-Pak (Waters Australia Pty Ltd, Rydalmere, NSW, Australia). The subsequent eluate was dried, reconstituted in 3% glacial acetic acid, and chromatographed on a column (1.5 × 90 cm) of Sephadex G-50 with 3% glacial acetic acid as the eluent. Fractions (100 × 1.4 ml) were collected, lyophilized, and assayed for GRP immunoreactivity by RIA. The column was calibrated with BSA (void volume), ovine GRP(1–27) and ovine GRP(18–27).

Statistical analysis

The data for GRP immunoreactivity in tissue samples were analysed by descriptive statistics, whereas the data for GRP mRNA synthesis (normalized to GAPDH mRNA) in endometrium isolated during the oestrous cycle or pregnancy were analysed by one-way ANOVA followed by the Tukey test for all pairwise multiple comparisons. Comparison between the oestrous cycle and pregnancy for the amount of GRP mRNA synthesis in endometrial tissue was made using two-way ANOVA followed by the Tukey test for pairwise multiple comparisons. These analyses were performed using SigmaStat statistical software (Jandel Scientific, San Rafael, CA, USA). $P < 0.05$ was accepted as significant. Data are presented as the mean $\pm$ S.E.M.

Results

Northern blot analysis for GRP mRNA during the oestrous cycle and early pregnancy

A 1 kb mRNA transcript was detected using ovine GRP cDNA as a probe. There was variable and low detection of GRP mRNA in endometrium isolated at days 4, 10, 12 and 14 of the oestrous cycle (Fig. 1a). Synthesis of GRP mRNA was normalized to the synthesis of GAPDH mRNA which was observed to decline slightly at day 16 of the oestrous cycle and moderately after day 14 of pregnancy; however, correction for GAPDH expression did not affect the overall trends in variation of GRP mRNA reported. The ratio of GRP mRNA to GAPDH mRNA was seen to increase markedly at day 16 of the oestrous cycle with a 24-fold rise compared with day 14 (Fig. 1c). For endometrium isolated during pregnancy (Fig. 1b), there was low or no detection of GRP mRNA at days 4, 7 and 8 ($n=4, 3, 3$ respectively, Fig. 1c) and variable and low detection at days 9 and 10 ($n=6, 5$ respectively, Fig. 1c). GRP mRNA was readily detected from day 12 of pregnancy (Fig. 1b), and a marked increase (7.8-fold) in GRP mRNA:GAPDH mRNA was observed between days 12 and 14. This level was maintained on days 16 and 17 and was followed by a further increase at day 20 (12-fold). This pattern observed during pregnancy is similar to that reported previously in a different group of animals (Giraud *et al.* 1994). GRP mRNA:GAPDH mRNA on day 14 of the oestrous cycle was significantly lower than on day 14 of pregnancy (Fig. 1c). Conversely, GRP mRNA:GAPDH mRNA was significantly higher on day 16 of the oestrous cycle compared with day 16 of pregnancy (Fig. 1c).

Cellular distribution of GRP mRNA

Hybridization of the ovine GRP antisense riboprobe to the ovine uterus was compared for days 10 and 16 of the oestrous cycle and for days 9, 17, 20, 40, 50 and 140 of pregnancy (Fig. 2; data not shown for days 9, 17 and 50 of pregnancy). Under the conditions used, hybridization was not apparent at day 10 of the oestrous cycle (Fig. 2A) or day 9 of pregnancy (not shown). These observations are consistent with the Northern blot analysis (Fig. 1). At day 16 of the oestrous cycle (Fig. 2C) and day 17 of pregnancy (data not shown), specific hybridization was observed in the glandular epithelium only. No signal was detected in the luminal epithelium, intercaruncular stroma, caruncular stroma, blood vessels or myometrium. A gradation of hybridization was observed, with strongest expression of GRP mRNA in glands closest to the lumen (superficial glandular epithelium) and little or no expression at glands furthest from the lumen (deep glandular epithelium). For uterine tissue taken at days 20 and 40 of pregnancy (Figs 2E and G) there was a further increase in hybridization to

Figure 1 Expression of GRP mRNA (upper panels) and GAPDH mRNA (lower panels) in (a) endometrium obtained from sheep during the oestrous cycle (days 4, 10, 12, 14 and 16 as indicated) and in (b) endometrium obtained from sheep during early pregnancy (days 12, 14, 16, 17 and 20 as indicated; days 4, 7, 8, 9 and 10 are not shown). A 20 µg portion of total RNA from each sample was fractionated by denaturing gel electrophoresis, transferred to nylon membrane and subjected to hybridization using radiolabelled ovine GRP cDNA as probe. (c) Densitometric representation of GRP mRNA relative to GAPDH mRNA in endometrium isolated from sheep during the oestrous cycle and early pregnancy. Data are plotted as the mean $\pm$ S.E.M. * $P < 0.05$ comparing oestrous cycle and pregnancy.

GRP antisense riboprobe by epithelial cells of glands closest to the lumen. Gradation of hybridization for superficial glands and deep glands continued to be observed. Intense hybridization of glandular epithelium

was maintained for day 50 pregnant uterine tissue, with every gland responding and no evidence of hybridization to cells of any other region (data not shown). In near-term uterine tissue (day 140 of pregnancy), strong hybridization was observed in the greatly expanded uterine glands (Fig. 2I). Diffuse hybridization to cells of the myometrium was also observed (data not shown). *In situ* hybridization for uterine tissue collected at days 10 and 16 of the oestrous cycle and days 9, 17, 20, 40, 50 and 140 of pregnancy yielded no response when a sense ovine GRP riboprobe was used (data for day 40 pregnancy is shown in Fig. 2H).

Cellular distribution of GRP immunoreactivity

Immunohistochemical techniques were used for the cellular localization of GRP peptide in sheep uterus obtained during the oestrous cycle and pregnancy. No peptide could be detected at day 10 of the oestrous cycle (Fig. 2B) in agreement with the lack of GRP mRNA (Figs 1a and 2A). At day 16 of the oestrous cycle, the apical cytoplasm of uterine gland cells stained for GRP immunoreactivity (Fig. 2D). At day 20 of pregnancy, the cytoplasm of uterine gland cells and the area immediately surrounding the glands showed moderately strong staining (Fig. 2F). In near-term uterus (day 140), intense immunohistochemical localization of GRP was observed in the apical cytoplasm of gland cells with immunoreactive material also apparent in the glandular lumen and the stromal tissue surrounding the glands (Fig. 2J). The immunohistochemistry generally reflected the changes seen with *in situ* hybridization except that no GRP mRNA was detected in stromal tissue.

GRP immunoreactivity in endometrial tissue during the oestrous cycle

GRP immunoreactivity was low (around 1.2 pmol/g) in extracts of endometrium obtained from ewes at days 4, 10, 12 and 14 of the oestrous cycle (Table 1). A significant increase (4-fold; 5.4 pmol/g) was observed at day 16.

GRP immunoreactivity in plasma obtained during the oestrous cycle and early pregnancy

No GRP immunoreactivity could be detected in plasma derived from the peripheral circulation and obtained from sheep at days 4, 10, 12, 14 and 16 of the oestrous cycle or pregnancy.

Chromatography

Endometrial tissue was taken from sheep at the mid-luteal phase of the oestrous cycle and 30 and 48 h after injection of a prostaglandin analogue (equivalent to luteal regression and oestrus respectively), and the molecular forms of GRP immunoreactivity were partially characterized by gel filtration chromatography (Figs 3A, B and C respectively).

Table 1 GRP immunoreactivity in ovine endometrial tissue isolated on different days of the oestrous cycle. Results are mean $\pm$ S.E.M. (n in parentheses)

Day of oestrous cycle	GRP (pmol/g)
4	1.6 $\pm$ 0.4 (4)
10	1.2 $\pm$ 0.2 (4)
12	1.4 $\pm$ 0.2 (4)
14	1.0 $\pm$ 0.2 (4)
16	5.4 $\pm$ 0.6 (3)

The extract from the mid-luteal phase endometrium displayed two molecular forms in approximately equimolar concentrations (Fig. 3A). These two forms were co-eluted with GRP(1–27) and GRP(18–27). The molecular profiles from tissue taken later in the oestrous cycle (Fig. 3B and C) showed three molecular mass forms; two of these were co-eluted with GRP(1–27) and GRP(18–27) and the third and predominant form was eluted before (and partly obscured by) GRP(1–27). This higher molecular mass peptide corresponds in elution position to the pregnancy-associated GRP-like peptide, which we have previously described in the gravid ovine endometrium (Giraud *et al.* 1996) and shown to be larger than human transforming growth factor- α (5.6 kDa).

Discussion

In this study we have shown that a peptide similar to, but larger than, GRP is temporally expressed in the ovine uterus during the oestrous cycle and that GRP mRNA is localized to the glandular epithelial cells during luteolysis, oestrus and pregnancy.

Northern blot analysis, *in situ* hybridization, RIA and immunohistochemistry revealed that expression of the GRP gene in the ovine uterus varied markedly during the oestrous cycle suggesting a regulated phenomenon. Previous data from our laboratory, and results from this study have suggested that the synthesis of uterine GRP is associated with the presence of a conceptus in the uterus, and is further enhanced after implantation (Giraud *et al.* 1994). Nevertheless, the present observations of substantial GRP expression in endometrium during the luteal regression phase of the normal oestrous cycle shows that GRP induction is a cyclic phenomenon which is initiated before

Figure 3 Gel filtration chromatography (Sephadex G-50) of extracts from 11 g of endometrial tissue isolated from non-pregnant sheep in (A) mid-luteal phase, (B) luteal regression and (C) oestrus. The elution positions of synthetic ovine GRP(1–27) and GRP(18–27) standards are indicated.

conception, and implies that GRP may participate in the cellular changes that occur in the uterus in preparation for pregnancy.

Figure 2 Cell-specific localization of GRP mRNA or peptide in uteri obtained from sheep in the oestrous cycle or pregnancy using *in situ* hybridization (A, C, E, G, H, I) or immunohistochemistry (B, D, F, J) respectively. The black colour in panels A, C, E, G, H and I indicates hybridization of the GRP antisense riboprobe, and the brown colour in panels B, D, F and J indicates the presence of immunoreactivity. (A) GRP mRNA (magnification $\times$ 110) and (B) GRP peptide (magnification $\times$ 500) at day 10 of the oestrous cycle; (C) GRP mRNA (magnification $\times$ 75) and (D) GRP peptide (magnification $\times$ 480) at day 16 of the oestrous cycle; (E) GRP mRNA (magnification $\times$ 160) and (F) GRP peptide (magnification $\times$ 500) at day 20 of pregnancy; *in situ* hybridization using GRP anti-sense riboprobe (G) (magnification $\times$ 16) and GRP sense probe (H) (magnification $\times$ 85) at day 40 of pregnancy; (I) GRP mRNA (magnification $\times$ 90) and (J) GRP peptide (magnification $\times$ 220) at day 140 of pregnancy.

A comparison of the changes in GRP mRNA during the oestrous cycle and pregnancy suggests that regulation of GRP gene expression may be related to ovarian steroid release. GRP mRNA and peptide levels are maximal at day 16 of the oestrous cycle when progesterone is low. However, both GRP mRNA and peptide are elevated at day 20 of pregnancy when progesterone is high. This observation of conflicting relationships between progesterone and GRP suggests that, if progesterone does have a role in regulating GRP gene expression, it can be overcome or modulated by other, probably local, factors. Synthesis of GRP at day 16 of the oestrous cycle correlates with the pre-oestrus surge of oestrogen and also with the concentrations of oestrogen receptors in the superficial and deep glands of the sheep endometrium (Cherny *et al.* 1991, Wathes & Hamon 1993). This could suggest that GRP is directly under the influence of oestrogen and acts as a mediator of the actions of oestrogen (i.e. as an oestromedin) on the uterus similar to growth factors such as epidermal growth factor and insulin-like growth factor (Brigstock 1991).

The site of GRP mRNA synthesis in uterine tissue during the luteal regression phase of the oestrous cycle was found to be the epithelial cells of the glands closest to the uterine lumen. However, GRP immunoreactivity was not confined to these cells but also found in the surrounding stromal tissue. This is consistent with rapid secretion of GRP peptide from glandular epithelial cells into the surrounding tissue and ultimately the uterine lumen and circulation and may explain why the increase in GRP peptide (5-fold) between days 14 and 16 of the oestrous cycle was not as marked as the increase observed for GRP mRNA (24-fold). Alternatively the differential between increases in GRP immunoreactivity and mRNA could reflect increased lability of the peptide or less-than-full cross-reactivity of our C-terminally directed antiserum with the endometrially derived GRP. GRP immunoreactivity in the stroma could represent intact peptide or products produced by the action of neutral endopeptidase, which is known to degrade bombesin and is present in the ovine endometrium (Riley *et al.* 1995). This remains to be determined.

In the ovine uterus, two types of cell were found to synthesize GRP mRNA: glandular epithelium during luteolysis and pregnancy and myometrial cells at the end of pregnancy. An important function of the glandular epithelial cells is production and secretion of the factors that comprise the histotroph such as glycogen, uterine milk proteins and growth factors. In many species, accessory placental structures known as areolae or chorionic vesicles develop opposite the uterine glands and are active in absorption of glandular secretions (King 1993). We propose that a peptide related to, but larger than, GRP be added to the list of components of uterine glandular secretions. The particular functions of this peptide remain to be determined, but, in view of its localization and

temporal synthesis, it is likely to have a function in nutrition and/or growth within the ovine utero-foetal axis. The continuous expression from early in gestation until term suggests another possible role in maintenance of the corpus luteum. GRP mRNA detected in myometrium at late gestation could signify a role for GRP in uterine contractions and possibly parturition. Nerve fibres containing GRP immunoreactivity have been observed in the myometrium of rat, guinea pig and pig (Happola & Lakomy 1989), and GRP and bombesin have been shown to stimulate concentration-dependent contraction of smooth muscle in isolated rat uterus (Stjernquist *et al.* 1986) mediated via a signal-transduction mechanism coupling a G-coupled seven-transmembrane domain receptor to phospholipase C activation (Amiot *et al.* 1993).

No GRP immunoreactivity could be detected in peripheral plasma obtained from sheep during the oestrous cycle or early pregnancy in contrast with plasma obtained from mid- and late-term sheep (Giraud *et al.* 1993) when GRP immunoreactivity is readily detected. This does not mean, however, that GRP is not secreted by the glandular epithelial cells. In a previous study (Giraud *et al.* 1996), we detected GRP immunoreactivity in uterine flushings during early pregnancy, suggesting that endometrial epithelial GRP is secreted apically. However, as the mass of the endometrium during the oestrous cycle and early pregnancy is relatively small compared with the mid-late-term pregnant uterus, the total amount of GRP reaching the circulation would probably be too low to allow detection.

A novel GRP was identified in the ovine endometrium during luteal regression and oestrus. Gel filtration chromatography suggests that this GRP peptide is larger than GRP(1–27) and GRP(18–27) and similar to the larger GRP previously described in the late-term pregnant uterus (Giraud *et al.* 1996). Larger forms of GRP mRNA were observed at day 16 of the oestrous cycle and days 14 and 20 of pregnancy. These are similar to the 1.9 kb and 4.3 kb transcripts that we observed in RNA prepared from late-term ovine endometrium (Shulkes *et al.* 1996) and others observed in RNA prepared from pulmonary carcinoma tumour, 18- and 22-week human foetal lung and forebrain (Spindel *et al.* 1987) and human small cell lung carcinoma cell lines (Sausville *et al.* 1986) and probably represent GRP mRNA precursors. The larger GRP peptide is unlikely to result from a different transcript arising from an alternative first exon, alternative splicing sites or a second GRP gene since two GRP cDNA clones prepared from pregnant sheep endometrium cDNA library in another laboratory (Fraser *et al.* 1994) and five GRP cDNA clones prepared in our laboratory (S C Whitley, unpublished observation) are all identical. Predicted translation of the GRP cDNA structure suggests a preproprotein that, if processed similarly to the human and rat preproGRP (Reeve *et al.* 1988), would yield a signal peptide, GRP(1–27), with an amidated C-terminus (and therefore recognized by the antibody used in our RIA) and

a C-terminal extension peptide. Further processing of the 27-amino acid peptide would yield a decapeptide identical with human GRP(18–27). The larger than expected GRP observed in the ovine endometrium during luteolysis and late pregnancy may suggest an as yet undescribed processing pathway for preproGRP.

Four subtypes of receptor with affinity for GRP have been identified (Nagalla *et al.* 1995): GRP-R with greatest affinity for GRP; NMB-R with low affinity for GRP and high affinity for the related peptide neuromedin B; BB4 with greatest affinity for [Phe¹³]-bombesin; and BRS-3 with moderate affinity for GRP and as yet no identified ligand. Bombesin receptor subtype 3 (BRS-3) mRNA has been detected using reverse transcription PCR in the pregnant myometrium of guinea pigs (Gorbulev *et al.* 1992) and women (Fathi *et al.* 1993, Gorbulev *et al.* 1994) but not in the non-pregnant myometrium of these species. Our own studies found BRS-3 mRNA to be expressed in the pregnant endometrium, myometrium, amnion, chorion and placenta isolated from three women at term delivery by caesarean section (Whitley *et al.* 1996). Thus BRS-3 could be the receptor that mediates the actions of the larger form of GRP in the ovine uterus. This possibility is made particularly attractive by the fact that the natural ligand for this receptor has not yet been discovered. Alternatively other GRP receptors may exist. At present, nothing is known of the expression of these receptors in the sheep uterus.

The original discovery of a larger-than-expected GRP in the pregnant sheep uterus (Giraud *et al.* 1996) was the impetus for the suggestion of a novel role for this misleadingly named peptide. The present study shows that GRP expression in the ovine uterus varies during the oestrous cycle and pregnancy, suggesting temporally dependent regulation at different sites. Cellular localization as deduced by *in situ* hybridization and immunohistochemistry indicates that GRP is a major transcription and translation product of the epithelial cells of the superficial glands during luteal regression late in the oestrous cycle and also during blastocyst elongation. The additional rise in expression of the GRP gene at the time of blastocyst implantation may indicate a positive feedback mechanism whereby secreted products of the implanting conceptus stimulate further expression of the uterine GRP gene.

In conclusion, our data have shown that a peptide similar to, but larger than, GRP is a major product of the ovine uterus, supporting the hypothesis that a GRP-related peptide has paracrine and/or autocrine roles in this tissue during the oestrous cycle and pregnancy.

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