Gene expression in the mammary gland

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Introduction

The mammary gland is the specialized secretory organ that provides essential nourishment to mammalian young in the form of milk. Milk is primarily composed of water, fats, lactose and proteins; the major protein components are the various caseins and the whey proteins α -lactalbumin, β -lactoglobulin (ruminants) and whey acidic protein (rodents).

Mammary development and milk protein gene expression are regulated by a number of peptide and steroid hormones, as well as cell–cell and cell–substratum interactions within the gland (Topper & Freeman, 1980; Levine & Stockdale, 1985; Li *et al.*, 1987). During gestation the secretory capacity of the mammary gland increases due to cellular proliferation and differentiation; concomitantly milk protein gene expression is initiated in preparation for sustained milk production after parturition. In late lactation milk production declines due to decreased demand, and at weaning rapid mammary gland involution occurs to a state that approximates that of the mature virgin gland. Our present understanding of the relative contribution that hormones and morphological factors have on mammary gland development and milk protein gene expression results from investigations in the whole animal and *in vitro*, including mammary gland explants (Topper *et al.*, 1975) and mammary epithelial cell culture systems (Rosen *et al.*, 1986).

Man has traditionally exploited ruminant milk and its products as a source of dietary protein and in the food technology industry (see Fox, 1982). It has recently been proposed that manipulation of milk composition is a candidate for genetic manipulation in domestic dairy animals (Wagner & Murray, 1985; Lathe *et al.*, 1986; Clark *et al.*, 1987; Jimenez-Flores & Richardson, 1988). Several groups have successfully demonstrated the feasibility of targeting gene expression to the mammary gland of mice (Stewart *et al.*, 1984; Ross & Solter, 1985; Leder *et al.*, 1986; Andres *et al.*, 1987; Choi *et al.*, 1987; Gordon *et al.*, 1987; Simons *et al.*, 1987; Lee *et al.*, 1988; Muller *et al.*, 1988; Schonenberger *et al.*, 1988; Tsukamoto *et al.*, 1988; Bouchard *et al.*, 1989; Lee *et al.*, 1989; Tremblay *et al.*, 1989). In Edinburgh, transgenic sheep carrying hybrid gene constructs that express a human plasma protein in their milk have been generated (Simons *et al.*, 1988; Clark *et al.*, 1989). At present the levels of expression of the transgenic protein in the milk are low and this illustrates the need for a greater understanding of the structure and regulation of the milk protein genes used to target expression to the mammary gland (Clark *et al.*, 1989).

A number of genomic clones encoding milk proteins have been isolated and their organizations described (Yu-Lee & Rosen, 1983; Campbell *et al.*, 1984; Qasba & Safaya, 1984; Jones *et al.*, 1985; David-Inouye *et al.*, 1986; Yu-Lee *et al.*, 1986; Hall *et al.*, 1987; Kawara *et al.*, 1987; Vilotte *et al.*, 1987; Alexander *et al.*, 1988; Ali & Clark, 1988; Devinoy *et al.*, 1988; Gorodetsky *et al.*, 1988; Laird *et al.*, 1988). The identification of DNA sequence elements that contribute to regulated milk protein

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gene expression has been impeded by the absence of a suitable system in which to evaluate expression of manipulation DNA sequences. Intra- and inter-species DNA sequence comparisons have suggested common regulatory elements in the promoter region of a number of different caseins and whey proteins (Yu-Lee & Rosen, 1983; Jones *et al.*, 1985; Yu-Lee *et al.*, 1986; Bisbee & Rosen, 1986; Hall *et al.*, 1987; Devinoy *et al.*, 1988; Laird *et al.*, 1988). Furthermore, an in-vitro DNA-protein interaction has been proposed between part of this DNA sequence motif and nuclear protein(s) isolated from various tissues, including the mammary gland (Lubon & Hennighausen, 1987, 1988). A cell system has been described that allows characterization of the rat β -casein promoter covering this region (Ball *et al.*, 1988; Doppler *et al.*, 1989); the applicability of this cell system to other milk protein genes has yet to be demonstrated. While the significance *in vivo* of these observations remains unclear, the advent of methods for generating transgenic animals containing manipulated DNA sequences allows the investigation of milk protein gene expression in the whole animal, under the influence of all contributory factors.

Generation of transgenic animals

There are currently three methods for incorporating exogenous DNA into the genome to produce transgenic animals. These are (1) retroviral vectors, (2) embryonic stem cells and (3) pronuclear microinjection (Palmiter & Brinster, 1986; Scangos & Bieberich, 1987; Simons & Land, 1987). At present most transgenic animals are produced by pronuclear microinjection of fertilized eggs (Fig. 1). Fertilized eggs are obtained from superovulated donor animals and tens to hundreds of DNA molecules are injected through a fine injection pipette into one of the pronuclei. Surviving eggs are implanted into foster mothers and proceed to term.

Fig. 1. Production of transgenic mice by pronuclear injection (see text for details).

Transgenic offspring are identified by molecular analysis of their DNA for the presence of the injected DNA molecule(s). The integrated DNA is frequently present in every cell at a single chromosomal location, usually as multiple copies in a tandem array. In mice up to 5% of injected eggs may develop to be transgenic. In domestic animals, such as sheep or pigs, the average success rate is approximately 1%. This may reflect differences in both reproductive physiology and the technical demands of microinjection into more fragile and almost opaque eggs (Simons *et al.*, 1988).

Ovine β -lactoglobulin

We have chosen to investigate expression of the gene encoding ovine β -lactoglobulin (BLG). β -Lactoglobulin is a whey protein of M_r 18 000 present in ruminant milk at a concentration of about 3 g/l. It is also found in the milk of horses, pigs, dogs and dolphins, but not in that of rodents and humans (Jenness, 1982; Pervaiz & Brew, 1985). While the protein has no recognized function it is related to a diverse family of other secretory proteins, many of which bind small lipophilic molecules (for references, see Ali & Clark, 1988; Ali, 1989). The genomic ovine BLG gene has been isolated and more than 7 kb of the DNA sequence encompassing the structural gene have been determined (Ali & Clark, 1988; Harris *et al.*, 1988). The DNA sequence suggests that a functional gene has been isolated, not a pseudogene, and that the pre-protein of 180 amino acids is encoded by 7 exons covering 4.9 kb of genomic DNA (Fig. 2). However, it remained to be demonstrated whether the isolated BLG gene contains the necessary *cis* regulatory sequences for efficient, regulated expression in the mammary gland.

Fig. 2. Organization of the ovine β -lactoglobulin clone SS1. **(a)** Bold line represents 5' and 3' flanking DNA while exon and intron sequences (not to scale) are represented by stippled and open boxes, respectively. **(b)** The extent of various regions of DNA used to generate transgenic mice by pronuclear injection: 7 and 45 correspond to lines of mice generated with the particular region illustrated (for results see Figs 3, 4 and 5) while ΔH , $\Delta A1$, $\Delta A2$ and ΔI are construct identifiers.

In the absence of a suitable in-vitro cell system in which to evaluate BLG expression we have investigated the ability of various BLG DNA constructs to direct the expression of BLG protein in milk of transgenic mice (Simons *et al.*, 1987; S. Harris, P. Simons & M. McClenaghan, unpublished

results). In parallel, we are also investigating the potential of transgenic sheep containing BLG-fusion genes as a source of authentic human protein in milk (Simons *et al.*, 1988; Clark *et al.*, 1989; A. Archibald, unpublished results). A mouse system would therefore be a rapid means of evaluating BLG-fusion constructs for their potential to direct the expression of a foreign protein in the milk of a domestic ruminant.

The mouse model system

To evaluate sheep BLG expression in mice, genomic DNA sequences must be correctly transcribed, mRNA translated, and the protein correctly processed and secreted into the milk, despite the absence of the BLG gene in mice. The ability of transgenes to direct the production of BLG protein or BLG-fusion gene products in a tissue-specific, regulated manner requires the presence of *cis*-acting DNA sequences that control BLG expression. In the absence of any information regarding the position of such sequences within the ovine clone SS1, the largest genomic DNA fragment from this clone was initially used to produce transgenic mice (Simons *et al.*, 1987). Additional lines of mice have been generated containing derivative BLG constructs to define more precisely those *cis*-acting DNA sequence elements important for tissue-specific, regulated expression (Simons *et al.*, 1987; Fig. 2).

The ability of various BLG constructs to direct the expression of BLG in mouse milk can generally be assessed by SDS polyacrylamide gel electrophoresis and Coomassie Blue staining of milk whey proteins as shown in Fig. 3(a). An additional protein of M_r 18 000 is present in milk from lactating transgenic mice, all other protein bands being present in milk from transgenic and control animals. This additional protein co-migrates with purified BLG, and western blotting of milk whey proteins confirms the identity of this protein as BLG (Fig. 3b). In abundantly expressing lines the level of BLG protein in mouse milk can be up to 5 times that present in sheep milk (e.g. line 7). This change in milk composition is not accompanied by any apparent deleterious side effects, either to the hemizygous transgenic females or their sucking offspring. Copy number-independent variations in BLG protein content are evident in different lines of mice containing the same BLG construct (e.g. lines 23 & 41), probably reflecting the influence of the site of integration on transgene expression in different lines (Palmiter & Brinster, 1986).

All the 5' and 3' flanking deletion constructs express similarly variable levels of BLG protein in the milk of transgenic mice (Fig. 2; S. Harris & M. McClenaghan, unpublished results). In contrast, the milk from three G0 females containing the intronless BLG construct (ΔI , Fig. 2) express a consistently low level of protein, detectable only by western blotting (S. Harris & M. McClenaghan unpublished results). This suggests a possible requirement for intron sequences for efficient BLG expression in transgenic mice. However, the ability of the BLG promoter region to direct significant levels of a foreign protein from a genomic DNA sequence containing introns (A. Archibald, unpublished results) suggests this may be a non-specific requirement, rather than a requirement for specific BLG intron sequences. It is also possible, although unlikely, that all 3 animals were either somatic mosaics or low level expressors, due to the site of integration. The dependence of gene expression on the presence of intron sequences has been noted by others (Brinster *et al.*, 1988). Further lines of mice would need to be analysed to clarify the mechanism(s) by which, intron sequences influence BLG expression in transgenic systems.

The tissue distribution of BLG expression has been examined in total RNA isolated from a variety of tissues and analysed after Northern blotting using BLG-specific cDNA probes (Simons *et al.*, 1987; Fig. 4). Mice expressing BLG protein in their milk contain a transcript present in RNA isolated from the lactating mammary gland that co-migrates with the equivalent sheep transcript of about 800 nucleotides (Gaye *et al.*, 1986). All BLG deletion constructs retain this strict tissue-specific expression pattern in transgenic mice, including the intronless gene (S. Harris, unpublished results). Furthermore, in all lines examined the RNA transcript initiates at the previously identified

Fig. 3. Analysis of whey proteins in the milk of mice transgenic for sheep β -lactoglobulin (BLG) (a) Milk samples were diluted 1/5 in distilled water and defatted by centrifugation. To prepare whey, the caseins were precipitated by the addition of 1 M-HCl to pH 4.5. Whey proteins (150 μ l) were precipitated in 5% TCA, washed in acetone, solubilized in 100 μ l loading buffer and electrophoresed on 15% polyacrylamide gels. Solubilized whey protein (2.0 μ l) from each transgenic mouse (as numbered) and one control mouse (CM) are shown electrophoresed alongside 10.0 μ l sheep whey protein (S), markers (M) and 2.5 μ g (a) and 10 μ g (b) of purified sheep BLG. (b) Western blot of whey proteins probed for sheep BLG. Samples from transgenic animals are in numbered lanes, from control mouse in lane CM and from sheep in lane S. Solubilized whey (1 μ l is equivalent to 0.3 μ l milk) was loaded as follows: 0.1 μ l for all transgenic samples except 7 (0.5 μ l); 23.5 and 23.9 (3 μ l); 8, 23 and 37 (5 μ l); 1 μ l for CM and S, 3 μ l. Standards are shown in lanes a, b, c and d and correspond to 100, 500, 2 and 100 ng purified BLG, respectively: note that a and b apply to the first 10 lanes, c to lanes 8, 23 and 37 and d to lanes 23.5 and 23.9. The high M_r band in tracks 8, 23 and 37 is due to non-specific binding of the antiserum. For further details see Simons *et al.* (1987). Part of figure reproduced by permission from Simons *et al.* (1987).

cap site (Ali, 1989; P. Brown & S. Harris, unpublished results). Moreover, when examined by immunohistochemical analysis, BLG protein expression is specific to the secretory epithelial cells within the mammary gland (S. Harris & M. McClenaghan, unpublished results).

Fig. 4. Tissue-specific expression of the sheep BLG gene in the mammary gland of transgenic mice. Northern blot analysis of 5 µg total RNA from 11-day lactating transgenic mice (numbered) and a control animal C57BL/6 (CM). The tissues analysed are mammary gland (Ma), liver (Li), kidney (Ki), spleen (Sp), salivary gland (Sa), and lachrymal gland (La). Lane S = RNA from the mammary gland of a lactating ewe. Duplicate transgenic mice give essentially identical results. For further experimental details see Simons *et al.* (1987). Reprinted by permission from Simons *et al.* (1987).

Occasionally a low abundance RNA transcript could be seen in other tissues after a long exposure, particularly in the salivary gland. This is most likely to be due to contamination of this tissue during dissection by thoracic mammary gland tissue because (1) other endogenous murine mammary specific transcripts that are known not to be expressed in the salivary gland are also present (e.g. β -casein), and (2) the quantity of salivary gland BLG transcript present in a given RNA sample fluctuates for a given transgenic line.

We have demonstrated abundant tissue-specific expression of the ovine BLG gene in transgenic mice. However, for the mouse model to be a useful representation of BLG gene expression in sheep it is also important that gene expression is temporally and developmentally regulated rather than constitutive. To examine this aspect of regulation in transgenic mice total RNA was isolated from virgin mice and other animals killed at specific times during gestation, lactation and weaning. Using RNA dot-blot analysis the level of BLG-specific RNA was determined for each time point (Fig. 5; S. Harris, unpublished).

Fig. 5. Expression of the ovine β -lactoglobulin gene during gestation in transgenic mice. Total RNA values isolated from mammary gland tissue at different days during gestation are shown after hybridization to 32 P-labelled DNA probes. RNA samples were transferred to filters through a manifold. Control mouse (C57BL/6), control sheep and 11-day post-parturient RNA samples were diluted 10-fold compared to gestation samples (line 7, see Fig. 1). The ovine β -lactoglobulin probe was derived from p931 (Gaye *et al.*, 1986); the murine β -casein probe was from pCas51 (Mehta *et al.*, 1981); the murine WAP probe from pmWAP1 (Hennighausen & Sippel, 1982) and the *Xenopus* ribosomal rDNA probe was from pMB9 (Sollner-Webb & Reeder, 1979).

When the expression profile for BLG is compared to that of RNA encoding other endogenous murine milk proteins, BLG is expressed with a developmental profile similar to that of endogenous murine β -casein, but different from that of the murine whey acidic protein. This BLG expression profile in mice resembles that established for BLG in sheep when equivalent time points are examined (Ali, 1989). Similar results are also obtained for deletion constructs over a limited time course (S. Harris, unpublished results). These observations, that the BLG transgene is temporally and developmentally regulated in transgenic mice, suggest that all the essential *cis*-acting DNA sequences required for regulated expression of the ovine BLG gene are represented within the deletion construct Δ A2 (Fig. 1).

Summary and conclusions

We have demonstrated that the ovine genomic clone SS1 can be used to generate transgenic mice that produce significant quantities of BLG protein in milk. The smallest BLG construct so far examined that retains the ability to direct BLG to mouse milk encompasses approximately 7.3 kb of genomic DNA, of which about 0.8 kb is derived from the promoter region. Gene expression is tissue-specific and regulated in a temporal and developmental fashion that is similar to that reported for sheep. We conclude, therefore, that the *cis*-acting sequences determining mammary expression of the ovine BLG gene are correctly interpreted in mice, despite the absence of an equivalent gene in this species, and that conclusions drawn from future work on BLG expression in the mammary gland of transgenic mice will be broadly applicable in sheep and other ruminant species.

Work is currently in progress to define other sequences within the promoter of BLG that are required for regulated expression in transgenic mice. These and other studies into the DNA-protein

interactions within the promoter which are required for efficient tissue-specific, regulated expression should lead to a greater understanding of milk protein gene expression in the mammary gland. Furthermore, in the current absence of a suitable in-vitro system, the mouse will be most useful for evaluating the expression of further constructs designed to express foreign proteins in milk of domestic ruminants.

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