Glucose and glutamine metabolism in pre-attachment cattle embryos in relation to sex and stage of development

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Summary. Individual Day-7 embryos (morulae to expanded blastocysts) were incubated with radiolabelled substrates and karyotyped to determine the sex. In Exp. 1, embryos were incubated for 3 h with D-[1-¹⁴C]glucose, as a measure of the activity of the pentose-phosphate pathway (PPP) and D-[5-³H]glucose, as a measure of total glucose metabolism. The labelled products ¹⁴CO₂ and ³H₂O were collected throughout the measurement period. Total glucose metabolism in male embryos was twice that in female embryos and increased between the morula and expanded-blastocyst stages. Relative to total glucose metabolism, PPP activity was four times greater in female than in male embryos. In Exp. 2, embryos were cultured with D-[1-¹⁴C]glucose, and L-[3,4-³H(N)]glutamine (a measure of Krebs cycle activity) in the presence of brilliant cresyl blue, a stimulator of the PPP. Glutamine metabolism increased from the morula to expanded-blastocyst stages. Relative to the metabolism of glutamine, the activity of the PPP was one-third greater in female than in male embryos.

Keywords: cattle; embryo; metabolism; development; sex

Introduction

The growth and development of early embryos requires considerable metabolic activity for the production of energy and for the synthesis of a variety of complex molecules. The relationship between development and a wide variety of aspects of metabolism has been studied in pre-implantation mouse and rabbit embryos (see Kaye, 1986). Apart from one study of pig embryos (Flood & Wiebold, 1988) and one of cattle embryos produced by in-vitro fertilization (Iwasaki *et al.*, 1989), we found no information about the relationship between metabolism and development of embryos of domestic animals.

The relationship between the sex and metabolic activity of early embryos has received little study in any species, but should be of major interest for two reasons. First, even before the development of the testis in mice, rats and humans, male embryos develop faster than females (see Mittwoch, 1989). This difference must, in some way, be related to differences in the activity of one or more metabolic pathways between male and female embryos. Second is the more specific effect of sex on the activity of the pentose-phosphate pathway (PPP) which produces two major biosynthetic substrates, ribose-5-phosphate, the precursor for the synthesis of nucleotides and nucleic acids, and NADPH, the major source of reducing equivalents for the synthesis of lipids and other complex molecules (Wood, 1985). The first and rate-limiting step of this pathway is catalysed by the enzyme glucose-6-phosphate dehydrogenase (G6PD) (Wood, 1985). The gene which codes for this enzyme is on the X chromosome (Chapman & Shows, 1976) and thus there are 2 copies in female cells but only one in male

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cells. This inequality has no effect on the biochemical activity of mature females, because early in embryonic development one of the X chromosomes is inactivated (Lyon, 1961). However, between the time of activation of the embryonic genome and the onset of X-inactivation, both X chromosomes in female embryos are potentially active and consequently the activity of several X-linked enzymes in broken-cell preparations from female mouse embryos has been shown to be twice that in male embryos (Monk, 1987; Monk & Handyside, 1988). In intact mouse embryos, total glucose uptake (Gardner & Leese, 1987) and reductive potential (Williams, 1986) tend to be higher in female embryos than in males.

We have measured the metabolic activity of individual cattle blastocysts using single radiolabelled glycolytic substrates and shown that the Krebs cycle, PPP and Embden–Meyerhof pathway (EMP) are active and that pyruvate kinase activity is blocked (Rieger & Guay, 1988). The technique has since been elaborated to permit the simultaneous measurement of any 2 of these pathways in individual embryos by using one ^{14}C -labelled substrate and one ^3H -labelled substrate together (Rieger *et al.*, 1989). The activity of the pathways can be related to each other, which is particularly important in studying the PPP because its activity can be related to total glucose metabolism, or to the activity of the Krebs cycle, which involves no known X-linked enzymes.

The study presented here used the double-label technique to determine the effect of sex and stage of development on the activities of the EMP and Krebs cycle and on the absolute and relative activity of the PPP in individual cattle embryos. In Exp. 1, the basal activities of the PPP and the EMP were measured. In Exp. 2, the activities of the PPP and the Krebs cycle were measured in the presence of brilliant cresyl blue (BCB), an oxidizing vital stain which stimulates the PPP (Williams, 1986). A preliminary report of some of the results presented here has been published as an abstract (Tiffin *et al.*, 1990).

Materials and Methods

Embryo collection. Embryos were collected at Day 7 (Day 0 = day of first detected oestrous behaviour) from groups of 3–6 superovulated Holstein heifers, as described by Rieger *et al.* (1988). The embryos were isolated from the collection medium and classified according to the morphological criteria of Elsdon *et al.* (1978). Only embryos judged to be morphologically normal were used. Embryos classed as morulae included those uncompact and compacted, but not obviously retarded. Early blastocysts showed some indication of the appearance of a blastocoele, while those classed as midblastocyst stage had a clearly defined blastocoele and an inner-cell mass. Expanded blastocysts were larger in diameter and had thinner zonae pellucidae.

Culture medium. The basic culture medium was bicarbonate-buffered Ham's F-10 (Flow Laboratories, McLean, VA, USA), which contains 6.1 mM glucose and 1 mM glutamine in addition to a wide variety of other amino acids, vitamins, fatty acids and pyruvate. This was supplemented with 25 mM *N*-2-hydroxyethylpiperazine-*N'*-2-ethanesulphonic acid (Hepes: Flow Laboratories), 0.4% bovine serum albumin (BSA: Sigma, St Louis, MO, USA), 100 iu penicillin/ml and 100 µg streptomycin/ml (Gibco Laboratories, Grand Island, NY, USA). Before measuring their metabolism, the embryos were washed 4 times in culture medium. For Exp. 2, the final wash contained 0.02 mg BCB/ml (Sigma).

Radiolabelled substrates. For Exp. 1, D-[1- ^{14}C]glucose (sp. act. 56.6 mCi/mmol: New England Nuclear, Boston, USA) was mixed with D-[5- ^3H]glucose (21.1 Ci/mmol: Amersham International, Amersham, UK). For Exp. 2, D-[1- ^{14}C]glucose was mixed with L-[3,4- ^3H (N)]glutamine (52.2 Ci/mmol: Amersham International). The mixtures were dried under nitrogen and taken up in culture medium to give nominal concentrations of 0.25 µCi/µl for each labelled substrate.

Metabolic measurement. The culture apparatus consisted of inner wells cut from multiwell strips (Lux, 5250: Miles Laboratories, Naperville, IL, USA) placed in 400 µl of 25 mM- NaHCO_3 within wells of larger multiwell plates (Falcon 3047: Becton-Dickinson, Lincoln, NJ, USA) as described by Rieger & Guay (1988). Individual embryos were taken up in 2 µl of the final wash and placed in the inner well of the culture apparatus, with 2 µl of the appropriate mixture of radiolabelled substrates, to give a total culture volume of 4 µl. Three sham preparations, containing all reagents, but no embryo, were included for each set of treatments, and served as control for all nonspecific measurements of activity due to machine background, chemiluminescence, bacterial contamination or spontaneous breakdown of the labelled substrate. The plates were equilibrated under 5% CO_2 in air for 15 min before being sealed with self-adhesive polyester film (Falcon 3073: Becton-Dickinson) and incubated for 1 (Exp. 2) or 3 h (Exp. 1). At the end of the culture period, 50 µl of 1.0 M-NaOH was injected through the polyester film into the outer well via a 25-gauge

needle to convert the dissolved CO_2 and bicarbonate into carbonate. The film was then removed and 200 μl of the contents of the outer bath was aspirated, mixed with 13 ml scintillation fluid (HP/b: Beckman Instruments, Fullerton, CA, USA) and radioactivity was measured for 5 min in a liquid scintillation counter (Packard Instrument Co., Downers Grove, IL, USA) programmed for automatic dual-label correction to disintegrations per minute (d.p.m.). The total d.p.m. of labelled substrate was determined by mixing 2 μl of the labelled substrate solution with 400 μl 25mM- NaHCO_3 and counting in the same way.

Estimation of product recovery. The efficiency of recovery of the radiolabelled products was estimated as described by Rieger & Guay (1988). Briefly, known amounts of $\text{NaH}^{14}\text{CO}_3$ and $^3\text{H}_2\text{O}$ (New England Nuclear) were taken up in 4- μl droplets of culture medium and cultured for 0–3 h, as for the embryos. At the end of each period, the outer bath was aspirated, mixed with scintillation fluid and radioactivity was measured. The maximum proportions of both labels were recovered within 30 min, yielding recovery correction factors of 1.37 for $^3\text{H}_2\text{O}$ and 1.41 for $^{14}\text{CO}_2$.

Calculation of substrate metabolism. The amount of each substrate metabolized by each embryo was calculated in a fashion similar to that previously described for single-labelled substrates (Rieger & Guay, 1988). For each labelled substrate, the mean d.p.m. for the sham preparations was subtracted from the d.p.m. for each embryo. The difference was divided by the total d.p.m. of labelled substrate added and multiplied by the total quantity of substrate in 4 μl of medium and by the appropriate recovery correction factor.

Cytogenetic analysis. Immediately after the metabolic measurement, the embryos were cultured individually in 0.5 ml of medium containing 0.05 μg Colcemid (Gibco)/ml for 4–8 h and then fixed individually on slides as described by King *et al.* (1979). The slides were stained in a 4% Giemsa solution for 3 min and the metaphase spreads located and counted under a compound microscope at low power. The chromosomes were examined at $\times 1000$ magnification and the sex was diagnosed as female by the presence of two X chromosomes in at least one intact metaphase, or as male by the observation of a Y chromosome. Preparations for which a definite conclusion regarding the sex of the embryo could not be reached were further stained for centromeric heterochromatin (Sumner, 1972) and examined again.

Statistical analysis. The distributions of embryos by sex (male, female, not determined) and stage of development (morula and early, mid-, and expanded blastocyst) in each experiment were evaluated by χ^2 analysis. Logarithmic values of the metabolic measurements were first evaluated by two-way analyses of variance (sex by stage of development) and group means across each factor compared by Duncan's multiple-range tests.

Results

In Exp. 1, the classifications by sex and stage of development were unrelated (Table 1, $P = 0.29$). The relationship between sex and stage approached statistical significance ($P = 0.06$) in Exp. 2, but this was entirely due to the distribution of embryos of undetermined sex because there was no relationship between classifications when only the sexed embryos were considered ($P = 0.62$).

Table 1. Numbers of cattle embryos, by sex and stage of development, in Exps 1 and 2

Developmental stage	Sex			Total
	Male	Female	Not determined	
Exp. 1				
Morula	10 (8.3)	5 (4.1)	36 (29.8)	51 (42.1)
Early blastocyst	3 (2.5)	3 (2.5)	15 (12.4)	21 (17.4)
Midblastocyst	1 (0.8)	5 (4.1)	19 (15.7)	25 (20.7)
Expanded blastocyst	2 (1.7)	1 (0.8)	21 (17.4)	24 (19.8)
Total	16 (13.2)	14 (11.6)	91 (75.2)	121 (100)
Exp. 2				
Morula	3 (5.6)	2 (3.7)	9 (16.7)	14 (25.9)
Early blastocyst	7 (13.0)	8 (14.8)	3 (5.6)	18 (33.3)
Midblastocyst	5 (9.3)	2 (3.7)	5 (9.3)	12 (22.2)
Expanded blastocyst	1 (1.9)	2 (3.7)	7 (13.0)	10 (18.5)
Total	16 (29.6)	14 (25.9)	24 (44.4)	54 (100)

Numbers in parentheses are percentages of total number of embryos used in each experiment.

For Exp. 1 (Table 2) analysis of variance for the metabolism of D-[1-¹⁴C]glucose indicated that the overall effect of sex approached significance ($P = 0.06$), but there was no effect of stage of development ($P = 0.29$) or interaction between sex and stage ($P = 0.35$) and no significant differences among individual group means. For the metabolism of D-[5-³H]glucose, there were highly significant ($P < 0.01$) effects of both sex and stage, but no effect of interaction ($P = 0.51$). The mean metabolism of D-[5-³H]glucose by male embryos was significantly greater than that by female embryos and increased with development from the morula to expanded blastocyst stage. There was a significant effect of sex ($P = 0.02$) on the ratio of D-[1-¹⁴C]glucose to D-[5-³H]glucose metabolism and a strong tendency toward a significant effect of stage ($P = 0.07$), but no effect of interaction ($P = 0.85$). The mean ratio was not different among the developmental stages, but, although not statistically different, was 4 times greater in females than in males.

Table 2. Glucose metabolism by Day-7 cattle embryos during a 3-h incubation at 37°C (Exp. 1)

Group	No. of embryos	Substrate metabolized (pmoles/embryo/3 h)		Ratio D-[1- ¹⁴ C]glucose: D-[5- ³ H]glucose
		D-[1- ¹⁴ C]glucose	D-[5- ³ H]glucose	
By sex				
Male	16	7.02 ± 1.15	35.00 ± 3.93 ^a	0.25 ± 0.06
Female	14	5.55 ± 0.97	15.40 ± 3.28 ^{ab}	0.99 ± 0.45 ^f
Not determined	91	4.10 ± 0.31	28.75 ± 1.98 ^b	0.21 ± 0.03 ^f
By stage				
Morula	51	4.34 ± 0.51	22.32 ± 2.07 ^{cd}	0.30 ± 0.05
Early blastocyst	21	3.68 ± 0.53	29.62 ± 3.00 ^c	0.21 ± 0.06
Midblastocyst	25	5.28 ± 0.74	27.26 ± 3.72 ^e	0.55 ± 0.27
Expanded blastocyst	24	5.53 ± 0.70	39.56 ± 4.82 ^{de}	0.18 ± 0.03
Total	121	4.66 ± 0.31	28.06 ± 1.67	0.31 ± 0.06

All values are means ± s.e.m.
^{abcdef}Values with the same superscript letter are significantly different ($P \leq 0.05$).

The effect of BCB on the activity of the PPP can be appreciated by comparing the metabolism of D-[1-¹⁴C]glucose in the presence (Exp. 2, Table 3) and absence (Exp. 1) of the stimulator. In Exp. 2, the overall mean metabolism of D-[1-¹⁴C]glucose was more than 5 times greater than in Exp. 1 (8.69 pmol/embryo per h vs. 1.55 pmol/embryo per h).

Analysis of variance of D-[1-¹⁴C]glucose metabolism in Exp. 2 indicated that there were no overall effects of sex ($P = 0.53$), stage of development ($P = 0.36$), or interaction ($P = 0.51$) and no significant differences among individual group means. There were also no effects of sex or interaction on the metabolism of L-[3,4-³H(N)]glutamine, but there was a highly significant effect of stage of development ($P < 0.01$). The mean metabolism of this substrate did not differ among the morula, early, and midblastocyst stages, but was significantly greater in expanded blastocysts than in the earlier stages. For the ratio of D-[1-¹⁴C]glucose to L-[3,4-³H(N)]glutamine metabolism, there was, similarly, no overall effect of sex ($P = 0.20$) or interaction ($P = 0.21$), but there was a significant effect of stage ($P = 0.04$). The ratio decreased significantly between the morula and expanded blastocyst stages, was almost twice as great in female embryos than in those of undetermined sex, and more than one-third greater in females than in males, although the last difference was not statistically significant.

The inability to determine the sex of the embryo by karyotyping was usually due to a lack of cells in metaphase and, less frequently, to chromosomes that were poorly spread or otherwise unanalysable. The success rate for sex determination more than doubled from 24.7% in Exp. 1 to 55.6% in Exp. 2, but it is unclear whether this was due to an improvement in technique, or to the

Table 3. Glucose and glutamine metabolism by Day-7 cattle embryos during a 1-h incubation at 37°C in the presence of 0.01 mg brilliant cresyl blue/ml (Exp. 2)

Group	No. of embryos	Substrate metabolized (pmol/embryo per h)		Ratio D-[1- ¹⁴ C]glucose: L-[3,4- ³ H(N)]glutamine
		D-[1- ¹⁴ C]glucose	L-[3,4- ³ H(N)]glutamine	
Sex				
Male	16	9.40 ± 1.52	0.55 ± 0.11	29.54 ± 9.75
Female	14	8.47 ± 0.86	0.41 ± 0.07 ^a	40.39 ± 11.95 ^e
Not determined	24	8.35 ± 0.57	1.44 ± 0.48 ^a	21.31 ± 4.82 ^e
Stage				
Morula	14	10.07 ± 1.66	0.55 ± 0.09 ^b	30.24 ± 11.10 ^f
Early blastocyst	18	7.41 ± 0.57	0.48 ± 0.09 ^c	33.68 ± 9.52 ^g
Midblastocyst	12	8.37 ± 0.70	0.51 ± 0.14 ^d	28.93 ± 5.74 ^h
Expanded blastocyst	10	9.46 ± 1.29	2.66 ± 1.04 ^{bcd}	17.26 ± 7.04 ^{fgh}
Total	54	8.69 ± 0.55	0.91 ± 0.22	28.69 ± 4.61

All values are means ± s.e.m.

Values with the same superscript letter are significantly different ($P \leq 0.05$).

shorter measurement period in Exp. 2. The metabolism of D-[1-¹⁴C]glucose and D-[5-³H]glucose by embryos in which the sex could not be determined was not different from that of either the male or the female embryos. However, in Exp. 2, the metabolism of L-[3,4-³H(N)]glutamine by the embryos of undetermined sex was 3 times greater than that for sexed embryos.

Discussion

We have previously shown that much less ¹⁴CO₂ is released from D-[6-¹⁴C]glucose than from D-[1-¹⁴C]glucose by Day-7 cattle embryos in Ham's F-10 medium immediately after collection (Rieger *et al.*, 1989), or after 24 h of culture to the blastocyst stage (Rieger & Guay, 1988). When such a relationship exists, Larrabee (1989) has suggested that the production of ¹⁴CO₂ from D-[1-¹⁴C]glucose can be considered to arise exclusively from processing through the PPP. Similarly, O'Fallon & Wright (1987) have shown that ¹⁴CO₂ release from D-[6-¹⁴C]glucose by mouse embryos is minimal in the presence of lactate or pyruvate and have taken the production of ¹⁴CO₂ from D-[1-¹⁴C]glucose as a direct measure of PPP activity under those conditions. O'Fallon & Wright (1987) suggested that the production of ³H₂O from D-[5-³H]glucose reflects only the metabolism of glucose through the EMP and thus have calculated total glucose metabolism as the sum of this activity and the activity of the PPP. However, that portion of D-[5-³H]glucose-6-phosphate that passes through the PPP and is not recycled will re-enter the EMP as [2-³H]-glyceraldehyde phosphate and the tritium is ultimately released as ³H₂O. The very small quantities of ¹⁴CO₂ released from D-[6-¹⁴C]glucose also indicate that recycling is limited. Consequently, we have taken the production of ³H₂O from D-[5-³H]glucose as a measure of total glucose metabolism and calculated the proportion of glucose metabolism by the PPP as the ratio of the production of ¹⁴CO₂ from D-[1-¹⁴C]glucose to the production of ³H₂O from D-[5-³H]glucose.

In Exp. 1, metabolic activity of the embryos was measured in the absence of any metabolic stimulators and these measurements may then be considered to represent basal metabolic rates and be compared with studies of the embryos of other species. The results of this experiment indicate that the absolute activity of the PPP, evaluated by the metabolism of D-[1-¹⁴C]glucose alone, or relative to D-[5-³H]glucose metabolism, does not change significantly during development of cattle embryos from the morula to expanded blastocyst stages. This is similar to the pattern observed in mouse embryos (O'Fallon & Wright, 1986), but differs markedly from that in pig embryos (Flood & Wiebold, 1988), where the metabolism of D-[1-¹⁴C]glucose increases significantly between the

compacted morula and blastocyst stages. Using a histochemical technique, Iwasaki *et al.* (1989) were unable to detect G6PD in cattle morulae, which could simply reflect the relative insensitivity of this approach, although they did find significant G6PD activity up to the 8-cell stage.

The total metabolism of glucose, as measured by the metabolism of D-[5-³H]glucose, increased significantly between the morula and expanded blastocyst stages, as has been shown for mouse (O'Fallon & Wright, 1986) and pig (Flood & Wiebold, 1988) embryos. Glucose uptake has similarly been shown to increase markedly between the 8-cell or morula and blastocyst stages in mouse (Gardner & Leese, 1986, 1988) and human (Gott *et al.*, 1990) embryos.

Total glucose metabolism was significantly greater in male embryos than in female embryos, which is, to our knowledge, a novel observation. The biochemical mechanism responsible for this difference is unknown, but it may have a physiological significance related to the different rates of development of male and female embryos. Within groups of Day-7 embryos collected from individual superovulated cattle, Avery (1989) found that embryos of the least advanced developmental stages are predominantly female, those of the most advanced stages are predominantly male, and the sexes are equally represented among those of intermediate stages. In a study of cattle embryos produced by in-vitro fertilization, those which cleaved early were predominantly male and those which cleaved late were predominantly female (B. Yadav, W. A. King & K. J. Betteridge, unpublished observations). Although it is likely that the greater rate of total glucose metabolism by male embryos is related to their more rapid development, it is unclear which is the cause and which is the effect.

Gardner & Leese (1987) have observed that glucose uptake (measured as disappearance from the medium) by female mouse embryos is ~10% greater than that by male embryos. However, this observation is not necessarily contradictory to our own, because, whereas we measured only glucose metabolized through the PPP and the EMP, measurement of uptake would also include glucose which was incorporated into glycogen and other macromolecules. Ménéz & Khatchadourian (1990) have suggested that the accumulation of glycogen is related to the 'two-cell block' to development in cultured mouse embryos. If the slower development of female embryos can be considered a milder form of impeded development, it would be interesting to know whether glycogen accumulation is greater in female than in male embryos.

The mean proportion of glucose metabolized through the PPP was much greater in female embryos than in males, which supports the hypothesis that G6PD activity is greater in female embryos because of the double gene dose. In 2- to 8-cell cattle embryos, Iwasaki *et al.* (1989) found no difference in G6PD activity between males and females, and no consistent relationship between G6PD activity and the number of X chromosomes in polyploids. However, the embryonic genome in cattle is probably not expressed before the 16-cell stage (see Telford *et al.*, 1990) and therefore the G6PD measured by Iwasaki *et al.* (1989) was almost certainly of maternal origin and would not have reflected the embryonic X chromosome dose.

We performed a number of small preliminary experiments (data not shown) to stimulate the PPP, in an effort to overcome any intracellular control mechanisms which might mask the difference between males and females. For example, we have previously shown that phenazine ethosulphate stimulates the activity of the PPP (Rieger & Guay, 1988), but we have been unable to find an effective concentration that is not lethal to cattle embryos. Conversely, exposure of embryos to 0.01 mg BCB/ml significantly increased the metabolism of D-[1-¹⁴C]glucose while having no effect on the metabolism of L-[3,4-³H(N)]glutamine and was not lethal to the embryos.

Although BCB clearly stimulated the metabolism of D-[1-¹⁴C]glucose (compared with Exp. 1), the stimulation was equivalent in both sexes of embryos. In contrast to total glucose metabolism in Exp. 1, the metabolism of L-[3,4-³H(N)]glutamine by unsexed embryos was significantly greater than that by sexed embryos. The reason for this is not evident, but we speculate that it may reflect a reduced viability of the unsexed embryos because it resembles the direct relationship that we have observed between glutamine metabolism and cell death in cryopreserved Day-6.5 horse embryos (Rieger *et al.*, in press).

In the present study, L-[3,4-³H(N)]glutamine metabolism was also significantly greater in expanded blastocysts than at earlier stages. The fact that total glucose metabolism was also greater in expanded blastocysts (in Exp. 1) suggests that there is a greater energy requirement at this stage. In rabbit embryos, development of the blastocoele is accompanied by marked increases in the energy requirements for Na⁺/K⁺ ATPase (Benos & Balaban, 1980), ouabain binding (Benos, 1981) and the content of Na⁺/K⁺ ATPase α -subunit and α -subunit mRNA (Gardiner *et al.*, 1990). There is no information available on Na⁺/K⁺ ATPase in cattle embryos, but it is likely that much of the increased metabolic activity in expanded blastocysts is to provide the ATP required by this ion pump, for expansion of the blastocoele.

The activity of the PPP, in this case compared with the metabolism of L-[3,4-³H(N)]glutamine by the Krebs cycle, was greater in female embryos than in males, but, as in Exp. 1, the difference was not statistically significant. We have suggested (Rieger, 1984) that it should be possible to determine the sex of an intact embryo by measuring the activity of an X-linked enzyme during the period between the activation of the embryonic genome and X inactivation. The enzymes in the embryos of the stages used in this study almost certainly arise from transcription and translation of the embryonic genome, rather than being of maternal origin (see Telford *et al.*, 1990). X inactivation begins at the early blastocyst stage in mouse embryos (West, 1982) and ~2 days after the trophoctoderm has differentiated in horse embryos (Romagnano *et al.*, 1987). The time of X inactivation in cattle embryos is unknown, but we have observed cells having 2 active X chromosomes as late as Day 13 (W. A. King, unpublished).

In conclusion, we have shown that total glucose metabolism is greater in male Day-7 cattle embryos than in females, which may be related to more rapid development of male embryos. Total glucose metabolism and glutamine metabolism increased with development from the morula to the expanded blastocyst stage, which probably reflects the increased requirement for energy for Na⁺/K⁺ ATPase activity. Although not statistically significant, the activity of the PPP tended to be greater in female embryos than in males, which supports the concept of a greater content of the X-linked enzyme G6PD in female embryos due to the double dose of the gene. Exposure of embryos to 0.01 mg BCB/ml increased the relative activity of the PPP without being lethal, indicating that BCB can be used effectively in the study of embryo metabolism.

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