Uterine proteins and the activation of embryos from mice during delayed implantation

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Summary. Ovariectomy-induced delay of implantation was used to study the role of the uterine environment in controlling implantation in mice. Labelling studies *in vivo* showed that uterine protein synthesis and secretion is maximal 2–5 h and 24–30 h after the oestradiol injection which initiates implantation. Embryos removed from uteri 5, 12 or 30 h after oestradiol injection were able to transport and utilize precursors of nucleic acids and proteins in short-term cultures at the same rate as normal embryos, although ‘delayed’ embryos had low levels of activity. These results suggest that ‘delayed’ embryos are metabolically activated within 5 h of release from delay, perhaps because of the hormonally-induced changes in uterine proteins which occur at this time.

Introduction

During early development, the rodent uterus regulates embryonic metabolism through the two ovarian steroids, oestrogen and progesterone (Psychoyos, 1967). In the nonlactating mouse, the development of the blastocyst takes place in an oestrogen-sensitized uterus (Finn & Martin, 1972) and results in elevated levels of embryonic RNA synthesis (Daentl & Epstein, 1971), protein synthesis (Epstein & Smith, 1973), and other metabolic activities (McLaren, 1973). In a lactating rodent (McLaren, 1968), or one which has been ovariectomized before the endogenous ‘oestrogen surge’ and the pregnancy maintained with exogenous progesterone (rat: Psychoyos, 1967; mouse: Yoshinaga & Adams, 1966), the embryos enter an unsensitized uterus and become metabolically quiescent (McLaren, 1968; Weitlauf, 1973, 1974a).

This paper describes an analysis of the interaction, leading to implantation, between the embryo and its uterine environment using the experimentally induced delay of implantation in the mouse (Yoshinaga & Adams, 1966) as a model system.

Materials and Methods

Outbred mice of the CFLP strain were induced to superovulate with 10 i.u. PMSG (Folligon: Organon) and 10 i.u. HCG (Chorulon: Organon) 48 h later. CFLP males were used for mating and the day the vaginal plug was found was designated as Day 0·5 *post coitum* (*p.c.*). In these mice, embryos enter the uterus on the morning of Day 3·5 and the oestrogen surge occurs in the afternoon, after which the embryos expand and lose their zonae by the morning of Day 4·5. Implantation has begun by the evening of Day 4·5 (Gardner, 1972).

The ovaries were removed under Avertin (Winthrop) anaesthesia, leaving the oviducts intact, before 12.00 h on Day 3·5. Pregnancy was maintained by subsequent daily s.c. injections of 1 mg progesterone (Δ^4 -pregnen-3,20-dione) in arachis oil for 6 days. The implantation process was initiated with one s.c. injection of 50 ng oestradiol benzoate in arachis oil on the following day and would have been complete within 48 h had not various experiments been carried out as indicated in the ‘Results’.

A mixture of ^{14}C -labelled amino acids (sp. act. 57 mCi/milliatom of carbon: Radiochemical Centre, Amersham) or ^{35}S methionine (sp. act. 50 Ci/mmol: Radiochemical Centre, Amersham)

was injected into the lumen of each uterine horn from the ovarian end following exposure of the uterus by a dorsal incision under Avertin anaesthesia. The volume injected was restricted to 3 μ l/horn.

At autopsy, uteri were flushed with cold sterile phosphate-buffered saline and the fluid centrifuged (1000 g, 15 min), samples not containing embryos being discarded. Endometrial tissue was homogenized at 4°C in phosphate-buffered saline and cleared by centrifugation (5000 g, 30 min). When necessary, protein samples were concentrated by precipitation with ethanol, collected by centrifugation and dissolved in the appropriate buffer. Protein content was estimated using the method of Lowry, Rosebrough, Farr & Randall (1951). Radioactive protein was precipitated and counted as described by Davies & Hall (1969).

Samples for electrophoresis were reduced with 1% β -mercaptoethanol and analysed on sodium dodecyl sulphate (SDS)-polyacrylamide gels calibrated with proteins of known molecular weight (Shapiro, Vinuela & Maizel, 1967). Gels containing radioactive proteins were cut into 2 mm slices, solubilized using the techniques of Young & Fulhorst (1965), and counted in Triton-toluene scintillation fluid (Patterson & Greene, 1965).

The handling and washing of embryos and extraction of labelled RNA were carried out as described by Daentl & Epstein (1971). For studies of uridine uptake and incorporation, embryos were cultured in Eagle's basic medium (BME: Flow Laboratories) supplemented with 5% heat-inactivated and dialysed fetal calf serum. The [5,6- 3 H]uridine (sp. act. 49 Ci/mmol: Radiochemical Centre, Amersham) was used at a concentration of 50 μ Ci/ml. The acid-insoluble radioactivity was completely solubilized by hydrolysis with NaOH or treatment with pancreatic RNAase (Daentl & Epstein, 1972), and was resistant to hydrolysis with RNAase-free DNAase (Worthington) in the presence of 10 mM-MgCl₂. All radioactivity detected in acid-insoluble material was therefore considered to be incorporated into RNA. For studies of protein synthesis, embryos were incubated with L-[4,5- 3 H]-leucine (sp. act. 55 Ci/mmol: Radiochemical Centre, Amersham) at a concentration of 100 μ Ci/ml in PBI medium containing 4 mg BSA/ml (Whittingham, 1971) and the acid-soluble and insoluble material was isolated as described by Brinster (1971).

Results

Uterine proteins

The total protein content of uterine flushings during normal pregnancy in mice is small (Table 1), but the increase after the oestrogen surge which occurs in the afternoon of Day 3.5 continues until Day 5.5 after implantation, when serum proteins are the main constituents of the uterine luminal proteins. Before attempting to relate changes in the luminal protein content to the 'activation' of the embryos, it was necessary to establish whether these proteins were different from those of serum. Plate 1 illustrates the patterns obtained when proteins from uterine secretions of mice with delayed implantation were analysed in SDS-polyacrylamide gels and compared with serum proteins. This procedure separates proteins on the basis of molecular weight and the mobility of a particular protein relative to the dye front is proportional to $\log_{10}$ molecular weight of that protein. Most of the proteins specific to the uterine lumen, i.e. not present in serum, migrate ahead of albumin and therefore have a molecular weight below 70×10^3 . This group of fast migrating proteins (P) was present in the uterine flushings of mice throughout delay of implantation and up to 30 h following administration of oestradiol (Pl. 1, Figs 1-4). By 12-15 h after termination of the delay of implantation with an oestradiol injection, two additional stained bands (A and C) with molecular weights of 120×10^3 and 65×10^3 respectively appeared in the uterine flushings and were still present at 30 h (Pl. 1, Figs 2-4), by which time embryo attachment had begun. These observations suggest that the luminal protein content of the pregnant mouse uterus depends qualitatively and quantitatively on its conditioning by ovarian steroids.

The synthesis and secretion of proteins into the uterine lumen was therefore examined systematically by pulse labelling uteri at various intervals after the oestradiol injection. The results in Table 2 demonstrate that there were two periods of synthesis and/or secretion of uterine proteins at

Figs 1-5. SDS-polyacrylamide gels of proteins in the pooled uterine flushings from 6 mice in delay of implantation at various times after initiating implantation with oestradiol treatment. The proteins were labelled with a mixture of ^{14}C -labelled amino acids ($0.5 \mu\text{Ci}/\text{horn}$) and each gel was loaded with $150 \mu\text{g}$ protein. The Bands A and C refer to regions incorporating radioactivity (see Text-fig. 1). P refers to the group of uterine proteins with molecular weights below 70×10^3 .

Fig. 1. Flushings from animals in delay (no oestradiol).

Fig. 2. Flushings from animals 2.5-5.0 h after oestradiol injection.

Fig. 3. Flushings from animals 12 h after oestradiol injection.

Fig. 4. Flushings from animals 30 h after oestradiol injection.

Fig. 5. Normal mouse serum.

(Facing p. 2)

Table 1. The mean $\pm$ S.E.M. protein content of uterine flushings from non-lactating pregnant mice (at least 5/group)

Days post coitum	Hours	Protein content (μ g/mouse)
Normal		
2.5	20.00	33 ± 6
3.5	20.00	18 ± 5
4.5	12.00	28 ± 4
4.5	20.00	54 ± 6
5.5	09.00	125 ± 13
'Delay'		
10.0	09.00	16 ± 5

Table 2. The incorporation of [35 S]methionine (2 μ Ci intraluminally 2 h before death) into proteins of uterine flushings of mice in delay of implantation at various times after administration of oestradiol

Time after oestradiol injection (h)	0	2	4	15	20	24	30	40
Acid-insoluble [35 S]methionine (ct/min $\times 10^{-2}$ per mg protein)	510	1400	60	20	50	100	1080	1110

Each value represents the mean for 3 animals.

approximately 2 and 30 h after injection of oestradiol. Since methionine is a relatively infrequent amino acid, the experiment (see legend to Text-fig. 1) was repeated using 14 C-labelled amino acids and similar results were obtained. When a short labelling period (2 h) was used, most of the uterine proteins synthesized and/or secreted into the uterine lumen during delay of implantation had a molecular weight of 70×10^3 and above (see Text-figs 1a and 1b, no oestradiol). The group of uterine proteins with molecular weights below 70×10^3 (P) did not incorporate label under these conditions. The major change in the profile of radioactive uterine proteins occurred within 5 h of administering oestradiol and involved radioactive peaks with the mobilities of Bands A and C (Plate 1). The Band C protein(s) is apparently only synthesized during the first 12 h after release from delay while Band A protein(s) continues to incorporate label for up to 30 h. Apart from these proteins, another prominent band of radioactivity (Band B: mol. wt approximately 90×10^3) was observed in the endometrium but this disappeared after oestradiol treatment. The proteins in Bands A, B and C were shown to be of uterine rather than embryonic origin by their appearance in the uteri of ovariectomized pseudopregnant mice subjected to a similar hormonal regimen.

Metabolism of embryos

Because release from delay of implantation is associated with changes in the synthesis of uterine proteins, it is possible that these proteins are involved in the activation of the quiescent embryos. Attempts were therefore made to relate some aspects of the metabolism of 'delayed', activated and normal embryos to their respective uterine environments.

Embryos in 'delay' are probably reversibly arrested in the G-1 phase of the cell cycle (Sherman & Barlow, 1972), and as such exist in an analogous physiological state to cells maintained *in vitro* under conditions of nutritional deprivation. These embryos might therefore be expected to exhibit a typical pleiotypic response to activation for implantation, e.g. changes in amino acid and nucleoside transport and incorporation (Kram, Mamont & Tomkins, 1973) followed 24–30 h later by DNA synthesis and mitosis (Todaro, Lazar & Green, 1965; Sander & Pardee, 1973). Consequently, these parameters were the ones selected to compare the metabolic activities of the different types of embryos.

A medium which is known to support a degree of postimplantation development was used (Spindle & Pederson, 1974) and incorporation of [3 H]uridine into RNA by normal (4.5 days *p.c.*)

Text-fig. 1. Radioactivity scans of (a) pooled uterine flushings (see Plate 1) and (b) endometrial homogenates of mice in delay of implantation and treated with oestradiol at various times before death. Each pool is represented by 6 mice in (a) and 3 mice in (b). The arrows indicate the mobility of albumin in each gel. O = origin; F = front.

Table 3. The uptake (mean $\pm$ S.E.M.) and incorporation of [3 H]uridine or [3 H]leucine into normal, 'delayed' and activated mouse embryos

Embryos	Incubation time (h)	[3 H]uridine*		[3 H]leucine†	
		Total uptake (pmol/embryo)	Incorporation (pmol/embryo)	Total uptake (pmol/embryo)	Incorporation (pmol/embryo)
Normal	4	0.327 $\pm$ 0.005	0.0361	0.150 $\pm$ 0.009	0.0765
	24	0.633 $\pm$ 0.008	0.0776	0.235 $\pm$ 0.008	0.1090
Delayed (untreated)	4	0.159 $\pm$ 0.009	0.0079	0.012 $\pm$ 0.001	0.0010
	24	0.749 $\pm$ 0.060	0.0821	0.130 $\pm$ 0.005	0.0600
Delayed (activated)					
	5 h	0.296 $\pm$ 0.006	0.0267	0.210 $\pm$ 0.002	0.0018
	12 h	0.272 $\pm$ 0.020	0.0270	0.248 $\pm$ 0.030	0.0012
	30 h	0.279 $\pm$ 0.030	0.0320	0.299 $\pm$ 0.040	0.0027

Mice with delay of implantation (10 day *p.c.*) were given either 1 mg progesterone (untreated group) 5 h before death or 1 mg progesterone + 50 ng oestradiol (activated group) 5, 12 or 30 h before being killed and the embryos removed. Embryos (normal) were also obtained from intact mice on the morning of Day 4.5 *p.c.* Values were obtained from three experiments in which embryos were assayed in duplicate in groups of 5–10.

* 0.1 pmol uridine represents 1.9×10^3 ct/min detected.

† 0.1 pmol leucine represents 2.4×10^3 ct/min detected.

embryos was linear for up to 6 h and was independent of total uptake at the concentration of precursor used (1.3 μ M). Table 3 demonstrates that after 4 h in culture the total uptake of uridine and its incorporation into RNA by normal embryos was substantially greater than for 'delayed implanting' embryos. However, after 24 h of culture the activity of the two types of embryo was similar. When [3 H]leucine uptake and incorporation into protein by 'delayed' and normal embryos was compared, the 'delayed' embryos were substantially less active than normal 4.5 day embryos (Table 3), but their activity increased after prolonged incubation, indicating that any repressive uterine influence on the delayed embryos had been overcome and resulted in activation of nucleic acid and protein synthesis similar to that of normal embryos. Short-term incubations are therefore necessary to detect metabolic differences between embryos.

To observe the effects of initiating implantation *in vivo*, embryos were removed from the uteri of mice in delay at 5, 12 and 30 h after injecting the mother with oestradiol and their metabolic activity was compared with that of unstimulated 'delayed' embryos and normal embryos (Table 3). The activated embryos behaved similarly regardless of the time spent in the uterus before incubation. The uptake and incorporation of leucine and uridine approached that of normal 4.5 day *p.c.* embryos within 5 h of the oestradiol injection.

Discussion

These experiments have confirmed the existence of proteins specific to the uterine lumen of the mouse (Mintz, 1970) and demonstrated changes in these macromolecules following the induction of implantation. Changes in uterine components during early pregnancy and the oestrous cycle have been observed in other species including the rabbit (Daniel, 1968), pig (Murray, Bazer, Wallace & Warnick, 1972), and man (Shirai, Iizuka & Notake, 1972). A steroid-binding function has been suggested for the progesterone-dependent rabbit protein, uteroglobin (Arthur, Cowan & Daniel, 1972; Fowler, Johnson, Walters & Pratt, 1976), but the function of the other uterine proteins and their possible roles in maintaining the development of the embryos remain obscure. It has been shown that uterine control is not operating at the level of availability or restriction of simple nutrients during this period (Weitlauf, 1968, 1971) and therefore the proteins that appear in the flushings after the initiation of implantation could be involved in controlling embryonic growth.

Since the cells of a diapausing embryo are probably arrested in the G-1 phase of the cell cycle (Sherman & Barlow, 1972) as are other non-dividing cells (Costlow & Baserga, 1973), it is possible

that the stimulus to implant is associated with a transition to the S phase (DNA synthesis). This is likely to result in increased transport and incorporation of nucleosides and amino acids similar to those induced by mitogenic stimulation or serum repletion in a variety of cell types (Sander & Pardee, 1973). Although the absence of data on endogenous pool sizes makes it difficult to compare the different types of embryos, previous autoradiographic studies of normal and 'delayed' embryos have confirmed *in-vitro* observations to be a true indication of the activity of such embryos *in vivo* (Weitlauf, 1971). In addition, the extent of uptake and incorporation of precursors into RNA and protein by normal preimplantation embryos is comparable to the values reported in previous studies using similar assay conditions (Brinster, 1971; Daentl & Epstein, 1972). Differences in incorporation of leucine and uridine *in vitro* after short culture periods are therefore taken to be indicative of a true difference in metabolic activity between diapausing and normal embryos. Activity is expressed on a per embryo basis in this paper, though mitosis may continue in some cells of 'delayed' embryos, resulting in an increase of approximately 20% over the total cell number in normal embryos (McLaren, 1968; Sherman & Barlow, 1972).

To understand the control of implantation, it must be determined whether embryonic activity is regulated directly by the ovarian steroids and, if not, how their effect is mediated through the uterus. The experiments described here do not completely exclude the possibility that steroids may be having a direct effect on embryos *in vivo* or *in vitro* (Smith & Smith, 1972; Lau, Davis & Chang, 1973). However, for the *in-vitro* experiments, steroid contamination was kept as low and as uniform as possible by washing all embryos three times in flushing medium and using a single batch of fetal calf serum after extensive dialysis. Under these conditions, 'delayed' embryos required more than 4 h incubation before uptake and incorporation of uridine increased to normal 4.5 day *p.c.* levels (Table 3). This lag for optimal RNA synthesis *in vitro* only occurred in 'delayed' embryos and has been reported by others (Psychoyos & Bitton-Casimiri, 1969; Weitlauf, 1974b). It is also associated with increased CO₂ production (Menke & McLaren, 1970) and the ability to resume proliferation (Gwatkin, 1966). All these activities are dependent upon a non-dialysable component(s) of fetal calf serum and its omission from the medium leads to resumed quiescence of the embryos. It therefore appears that after the oestrogen surge the uterus provides the same growth-enhancing environment as a serum-containing medium, and maintenance of the 'delayed' condition is due to absence of these growth stimuli.

The results of the experiments involving activation of embryos *in vivo* support this idea. Permeability and synthetic changes are established within 5 h of priming with oestradiol (Table 3). The incorporation of uridine into embryonic RNA, detected *in utero* by autoradiography, increases within 1 h of administering oestradiol to 'delayed implanting' rats and is maintained for up to 18 h (Dass, Mohla & Prasad, 1969). The increase in metabolic activity observed *in vitro* occurs without the 4 h lag period observed for 'delayed' embryos (Table 3). This interval could represent the 4 h pre-activation period demonstrated in the experiments of Weitlauf (1973, 1974b), and the absence of such a lag period for embryos activated *in vivo* suggests that the necessary metabolic changes take place in response to the oestradiol treatment before removal of the embryos from the uterus.

Factors responsible for the activation of preimplantation embryos could be steroids (probably in the form of steroid-protein complexes) or proteins or glycoproteins acting in a similar manner to that of mitogenic lectins or polypeptide hormones. It is difficult to exclude the influence of steroids in *in-vivo* or *in-vitro* experiments. However, the observation that 'delayed' embryos undergo similar metabolic changes after activation *in vivo* or *in vitro* suggests that the factors responsible are growth-promoting proteins or glycoproteins similar to those already isolated from serum (Holley, 1974). Since the metabolic activation of 'delayed' embryos 5 h after oestrogen administration *in vivo* occurs in association with a major change in the profile of uterine proteins, it seems worthwhile investigating whether these uterine proteins (either alone or as steroid-protein complexes) have any direct regulatory influence on embryonic growth.

This work was supported by a grant from the Sybil Eastwood Memorial Trust, Cambridge, and a grant from the Ford Foundation to Professor C. R. Austin. I thank Dr A. Surani and Dr M. H. Johnson for helpful discussions.

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Received 29 May 1976