Regulatory roles for mitochondria in the peri-implantation mouse blastocyst: possible origins and developmental significance of differential $\Delta\Psi_m$

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Abstract

Studies of mitochondria in mouse and human oocytes and preimplantation stage embryos have focused primarily on their metabolic capacity to generate ATP. However, it is becoming increasingly apparent that mitochondria are also regulatory agents in other processes involved in the establishment of developmental competence, including calcium homeostasis and apoptosis. The magnitude of the inner mitochondrial membrane potential, or its polarity ($\Delta\Psi_m$), is a physiochemical property of mitochondria related to levels of organelle activity, and differences in the magnitude and spatial distribution of high- and low-polarized mitochondria have been suggested to influence oocyte and early embryo competence. Here, we investigated mitochondrial polarity in normal and diapausing peri-implantation-stage mouse blastocysts, and their corresponding out-growths, for indications of cell-type-specific regulatory functions or activities in which these organelles may be engaged. The results demonstrate that cell-type- and location-specific domains of differential $\Delta\Psi_m$ exist in the peri-implantation blastocyst and remain unchanged during blastocyst outgrowth and during delayed implantation, which for the latter, is accompanied by the suppression of mitochondrial oxidative phosphorylation. Our findings demonstrate that cell-type-specific $\Delta\Psi_m$ in the peri-implantation blastocyst is not an intrinsic property of the corresponding mitochondria but one that can be mediated by the dynamics of intercellular contact. Cells with high- or low-polarized mitochondria are differentially affected by photosensitization, with developmental consequences related to embryo behavior and outgrowth performance. Differences in polarity are discussed with respect to the participation of mitochondria in regulatory and morphogenetic processes in the normal peri-implantation embryo. The persistence of high $\Delta\Psi_m$ in the diapausing embryo is suggested to be associated with the regulation of levels of cytoplasmic free calcium and the ability of the embryo to reactivate development when delayed implantation terminates.


Introduction

It has long been known that mitochondria, by virtue of their ability to generate ATP, have a central role in the normality of early mammalian development (reviewed by Biggers & Borland 1976, Van Blerkom & Motta 1979). This conclusion is supported by recent studies (reviewed by Cummins 2002, Brenner 2004) in which metabolic, structural and numerical (organelle and mitochondria DNA copy number) defects have been be associated with maturational failure for the oocyte and premature arrest or abnormal development for the embryo (Van Blerkom et al. 1995, 2000, Müller-Höcker et al. 1996, Steuerwald et al. 2000, Reynier et al. 2001, Brenner 2004). However, whether mitochondria have a regulatory role in early development that is distinct from their metabolic contribution has only recently been considered with respect to other functions, such as their ability to sequester and release calcium, modify proteins or initiate apoptosis (reviewed by Van Blerkom 2004).

The magnitude of the potential difference (mitochondrial polarity) across the inner mitochondrial membrane ($\Delta\Psi_m$) is an electro-chemical property of these organelles that can be estimated in living cells with reporter stains whose fluorescent emission characteristics are potential (potentiometric) sensitive, such as JC-1 (Reers et al. 1995). Differences in $\Delta\Psi_m$ detected within and between cultured cells have been largely thought to reflect corresponding differences in mitochondrial function or levels of activity (Cossarizza et al. 1996, Salvioli et al. 1977, Dedov & Routogalis 1999, Dias et al. 1999). In the mouse
and human oocyte and early embryo, high- and low-polarized mitochondria have a distinct spatial distribution in the pericortical/subplasmalemmal and perinuclear domains respectively (Van Blerkom et al. 2002). Changes in spatial distribution or polarity (high to low, or vice versa) detected in fresh (Wilding et al. 2001, 2003, Van Blerkom et al. 2002, 2003, Acton et al. 2004) and thawed (Ahn et al. 2002, Jones et al. 2004) metaphase II oocytes and embryos have been associated with developmental incompetence or abnormality. These findings suggest that the analysis of $\Delta \Psi_m$ may be a means by which differential mitochondrial activity or regulatory function(s) can be investigated in living preimplantation-stage embryos, especially with respect to domains of high- and low-polarized organelles (Van Blerkom 2004).

Van Blerkom et al. (2002) reported that high-polarized mitochondria occur in the trophectoderm, but not the inner cell mass (ICM), of expanded mouse and human blastocysts. Here, mitochondrial polarity was examined in normal, diapausing and JC-1-photosensitized, hatched (peri-implantation) mouse blastocysts, and their respective normal, diapausing and JC-1-photosensitized, hatched blastocysts. We intended to investigate how location- and cell-type-specific $\Delta \Psi_m$ occurs and whether it indicates different regulatory functions for mitochondria. The results extend our earlier findings and show that the magnitude of $\Delta \Psi_m$ is determined by extrinsic forces and may reflect different regulatory roles for mitochondria during the peri-implantation stage.

Materials and Methods

Recovery of normal and delayed implanted mouse blastocysts

Superovulated mice were ovariectomized on day 2.5 of pregnancy (detection of vaginal plug = day 1) and given a single subcutaneous injection of Depo-Provera (1.0 mg). Chemical ovariectomy followed the protocol of MacLean-Hunter and Evans (1999), in which the estrogen antagonist/agonist tamoxifen (10 $\mu$g; Sigma) and Depo-Provera (1.0 mg; Pharmacia and Upjohn, Kalamazoo, MI, USA) were coadministered on day 2.5. Hatched blastocysts were recovered from untreated animals on day 4.5 and from treated animals on days 1–7 of delay by gently flushing the uterus with HEPES-buffered medium (M2 or HTF) supplemented with 1% BSA.

Blastocyst outgrowth

Outgrowths of normal and delayed implanted embryos occurred on glass cover slips in high-glucose-containing DMEM supplemented with essential amino acids, glutamine (GlutaMax) and 20% fetal bovine serum (Invitrogen), conditions previously shown to support activation from day 7 of delay and robust trophectoderm outgrowth and expansion in vitro (Van Blerkom et al. 1979, Van Blerkom & Chavez 1981). After 1.5 days of culture, the colonies resulting from normal and delayed implanted embryos were characterized by a highly flattened circumferential monolayer of outgrowing mural trophectoderm that surrounded a central mass of ICM cells. The ICM-containing central cores of the outgrowths were dissected from the mural trophectoderm with glass needles, exposed to trypsin (30 min, 0.25%) in calcium- and magnesium-free PBS with continual agitation and pipetting, and replated. A similar protocol was followed for JC-1-stained blastocysts illuminated in the FITC or RITC channels (see below). From three to five ICM masses were treated with trypsin, as described above, and replated. Proliferating ICM cells were restained with JC-1 at 6 h and on days 1–7 of culture, and examined by fluorescence microscopy. Undisturbed outgrowths were stained with JC-1 (see below) on days 1–5 of culture.

DNA and mitochondrial staining

Immediately after recovery, representative normal and delayed implanted blastocysts were fixed in PBS containing 3.7% formaldehyde for 15–20 min, and after washing in HTF supplemented with 1% BSA, they were exposed to the DNA-specific fluorescent stain DAPI (4',6-diamidino-2-phenylindole) 10 $\mu$g/ml for 15 min (Sigma). Living embryos were exposed to the following mitochondria-specific fluorescent probes (Molecular Probes, Eugene, OR, USA) (Van Blerkom et al. 2002, 2003, Van et al. 2004): rhodamine 123 (r123, 10 $\mu$g/ml, 30 min), MitoTracker Orange (MO, 200 nmol/l, 30 min) or JC-1 (5,5',6,6'-tetrachloro-1,1,3,3'-tetraethylbenzimidazolycarbocyanine iodide (1 $\mu$g/ml, 30 min). In other studies, normal and diapausing blastocysts were either mechanically collapsed by passage through a narrow-bore micropipette and stained with mitochondria-specific probes during re-expansion, or manipulated with glass needles such that a significant rent was made in the mural trophectoderm, directly exposing the ICM to the stains. Representative embryos and outgrowths were exposed to the $\Delta \Psi_m$-collapsing proton ionophore FCCP (carbonyl cyanide p-(trifluoromethoxy)phenyl-hydrazone (Sigma), 10 $\mu$g/ml, for 10 min before JC-1 staining. DNA analysis and mitochondrial fluorescence used conventional epifluorescence (EF) and scanning laser confocal microscopy (SLCM), as previously described (Van Blerkom et al. 2002, 2003, Jones et al. 2004).

BrdU staining

Normal and delayed implanted blastocysts and outgrowths were incubated in modified MEM containing 10 $\mu$M BrdU (5-bromo-2'-deoxyuridine) for a minimum of 4 h and a maximum of 24 h, followed by fixation and antibody staining, according to the manufacturer’s recommendations (In Situ Cell Proliferation Kit, FLUOS Assay; Roche). Briefly, intact embryos and outgrowths were washed in PBS, fixed at 4°C in a solution of 70% EtOH and 30% 50mM glycine (pH 7) for 30 min at room
temperature in PBS containing 0.1% Tween-20 (permeabilization buffer), and denatured in 4 M HCl for 10 min. Samples were washed several times in the incubation buffer (also provided with the kit) followed by 30-min incubation in the blocking solution provided with the kit. Washed samples were incubated in the presence of a fluorescein-conjugated, anti-BrdU monoclonal antibody for 45 min in a humidified chamber maintained at 37°C. Samples were washed in permeabilization buffer for 20 min and counterstained for 15 min in DAPI (10 μg/ml).

**Photosensitization**

Normal day-5 blastocysts were stained with JC-1 and exposed to ultraviolet illumination (100 W) in either the FITC or the RITC channel for up to 60 s with a × 40 fluorescence lens. After exposure, blastocysts were returned to culture and assessed for ATP content, BrdU incorporation, DNA fragmentation and oxidation, and outgrowth behavior and performance in vitro at timed intervals (see below). From preliminary studies, a standard exposure of 8 s was found to have differential effects on blastocyst activities that may be related to mitochondrial function.

**DNA oxidation and fragmentation analysis**

The detection of 8-deoxyoxygenanine has been used to assess quantitatively levels of oxidative damage to DNA, such as may result from the production of reactive oxygen species (e.g. superoxide) by structurally damaged or functionally compromised mitochondria (Struthers et al. 1998), that may occur after photosensitization. After illumination in the FITC or RITC channels, blastocysts were cultured for 24 h, washed in modified MEM, fixed in a PBS containing 3.7% formaldehyde (pH 7.4) for 20 min, and permeabilized and stained with Texas Red Avidin (1:200 dilution; Molecular Probes), as described by Radisky et al. (2005). Positive controls were generated by exposure to peroxide (100 μM/ml in PBS) for 30 min. The level of DNA fragmentation in normal, delayed implanted and photosensitized, JC-1-stained blastocysts was examined by whole-mount fluorescence microscopy after terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling (TUNEL), as previously described (Van Blerkom et al. 2001).

**ATP content analysis**

Measurements of net average ATP content followed a previously described protocol (Van Blerkom et al. 1995). Briefly, single blastocysts were rapidly frozen to −80°C in 200 μl ultrapure water. ATP levels were quantified by measuring the luminescence (Berthold LB 9501 luminometer, Berthold Technologies, Oak Ridge, TN, USA) generated in an ATP-dependent, luciferin-luciferase bioluminescence assay (Bioluminescence Somatic Cell Assay System; Sigma). A standard curve containing 14 ATP concentrations from 1 fmol to 5 pmol was generated for each series of analyses. Differences in cell number and ATP content were analyzed statistically by the unpaired t-test (two-sample) with values considered significant at P < 0.01.

**Time-lapse microscopy**

Hatched blastocysts that were untreated after recovery, recovered from the uterus on day 5 of diapause, and stained with JC-1 and illuminated in the FITC or RITC channel were cultured in outgrowth medium and imaged by time-lapse videomicroscopy at 5-min intervals. Cultures were maintained at 37°C, and the timed behavior of embryos was recorded on a laser disk, as previously described (Van Blerkom et al. 2001).

**Results**

**Nuclear and mitochondrial staining of normal and delayed implanted mouse blastocysts**

In preliminary studies, comparable patterns and intensities of fluorescent staining with DNA- and mitochondria-specific probes were observed in blastocysts on days 1, 3 and 5 of diapause with delayed implantation induced by surgical or chemical ovariectomy. Consequently, approximately 75% of the delayed implanted embryos examined in this study were obtained after chemical ovariectomy. Seventy day-4.5 blastocysts and a minimum of 30 embryos on days 1, 2, 3 and 5 of delayed implantation were stained with DNA- or mitochondria-specific fluorescent probes. Twenty-three and 12 embryos respectively were stained with JC-1 on days 6 and 7 of diapause.

Cell numbers in normal (Fig. 1A) and delayed implanted blastocysts (e.g. day 3; Fig. 1G) were derived from nuclear counts in representative embryos stained with DAPI, and were analyzed with pseudocolor imaging by scanning laser confocal microscopy (SLCM). Differences in average cell numbers were evident when normal day-4.5 blastocysts (119 ± 11; n = 29) were compared with embryos on days 1 (127 ± 9; n = 27) and 3 (131 ± 7; n = 31) of delay, and these differences were significant. There were no further increases in cell numbers beyond day 3. These findings agree with previous cell number determinations (Chavez & Van Blerkom 1979).

**Mitochondrial staining of normal and delayed implanted blastocysts**

Representative hatched blastocysts from cohorts recovered on day 4.5 of pregnancy and on days 1, 2, 3, 5 and 6 of delay were examined by conventional epifluorescence (EF) or SLCM after staining with MO (n = 16; e.g. Fig. 1B, normal day 4.5) or r123 (n = 28; e.g. Fig. 1C, day-5 delay, shown in the FITC channel). Mitochondrial staining was clearly detectable in the ICM and trophectoderm of all normal and diapausing embryos. Figure 1E and F show the typical pattern of JC-1 fluorescence in a day-4.5 blastocyst.
Figure 1 (Panels A and I) Fully compiled, pseudocolor, scanning laser confocal images (SLCM) of DAPI fluorescent nuclei in a normal day-4.5 mouse embryo and a diapausing blastocyst on day 3 of delayed implantation respectively. The following panels show mitochondria staining with MitoTracker Orange (panel B), rhodamine 123 (panel C) and JC-1 fluorescence in the FITC (panels E, H, K, N and Q) and RITC channels (panels F, I, L and O) in normal (panel D) and delayed implanted, diapausing blastocysts (panels J and M). Perinuclear (N) mitochondrial (M) JC-1, J-aggregate fluorescence in a single mural trophectodermal cell is shown by SLCM in panel P. (Panels D, J and M) Light microscopic images of the blastocysts from which the corresponding patterns of JC-1 fluorescence noted above were obtained. Differences in the distribution of mitochondrial JC-1 and J-aggregate fluorescence (panel S) in a mouse blastocyst outgrowth on day 1.5 (panel R) indicate cells containing high-polarized red-fluorescent and low-polarized green-fluorescent mitochondria. ICM: inner cell mass; pTR: polar trophectoderm; mTR: mural trophectoderm.
JC-1 staining of blastocyst outgrowths

Comparable patterns and extent of trophectoderm outgrowth (OG) were observed for day-4.5 blastocysts (n = 57) and diapausing blastocysts reactivated in vitro on days 3 (n = 26), 5 (n = 16) and 6 (n = 19) of delayed implantation. As in previous findings (Van Blerkom et al. 1979), a circumferential layer of highly flattened, mural trophectodermal cells surrounded a central core composed primarily of ICM cells (e.g. OG day 1, Fig. 1R; OG day 1.5, Fig. 2A). A characteristic spatial and cell-type-specific pattern of mitochondrial polarity was evident during the early stages of outgrowth (Figs 1S and 2B); low-polarized mitochondria (e.g. LP, Fig. 2B) occurred in the centrally located, multilayered mass of ICM cells, while high-polarized mitochondria occurred throughout the cytoplasm of trophectodermal cells (arrow, Fig. 1S; HP, Fig. 2B).

Cell-type-specific differences in mitochondrial polarity persisted between the ICM and trophectodermal compartments of undisturbed blastocyst outgrowths after cytokinesis and expansion of the trophectoderm ceased, around OG days 3–4 (OG day 5, Fig. 2C). However, continued culture, the spatial distribution that characterized high- and low-polarized mitochondria in trophectodermal cells became less distinct. While high-polarized mitochondria were confined to the margins of these cells (red arrows, Fig. 2D), perinuclear mitochondria were largely low polarized (white arrows, Fig. 2D), unlike the situation observed during earlier phases of active outgrowth and expansion, where perinuclear mitochondria were primarily high polarized (Fig. 2B). The behavior, outgrowth characteristics and differential pattern of JC-1 and J-aggregate fluorescence of the ICM and trophectoderm were identical in day-4.5 blastocysts (Fig. 1R and S) and diapausing embryos activated in vitro (day 5 of diapause, Fig. 2A and B).

To determine whether the low mitochondrial polarity detected in the ICM of intact blastocysts and undisturbed outgrowths was an intrinsic feature of these cells, ICM cores from day-1.5 outgrowths (asterisk, Fig. 3A; n = 21) were isolated, disaggregated and replated. During the early stages of disaggregation, the ICM cores appeared as clusters of round cells that were loosely connected at the periphery and more densely associated in the interior (Fig. 3B). When stained with JC-1 and imaged by fluorescence microscopy 1.5 h after disaggregation, some of the loosely associated peripheral cells showed apical J-aggregate fluorescence (red arrows, Fig. 3C), while those in the interior fluoresced green. From three to five disaggregated ICM cores were combined, replated and cultured. The resulting colonies of cells were uniformly small, mononucleated and phenotypically distinct (n = 12; e.g. Fig. 3D) from the highly flattened and occasionally multinucleated trophectodermal cells typical of undisturbed blastocyst outgrowths (e.g. Figs 2A and 4G).

ICM-derived cells showed robust proliferation during 7 days of monitored culture (e.g. day 4 of culture; Fig. 3D). After staining with JC-1, J-aggregate-positive cells were observed throughout the cultures (Fig. 3E), with the exception of high-density portions of the monolayer (asterisk, Fig. 3D), whose fluorescence characteristics indicated the presence of low-polarized mitochondria (asterisk, Fig. 3E). At higher magnifications, cells loosely associated in small clusters (such as cells indicated by a red arrow in Fig. 3E)
contained both high- (red arrow, Fig. 3F) and low-polarized mitochondria (green arrow, Fig. 3F). By contrast, isolated cells and cells in low-density regions of the monolayer whose margins were largely free (e.g. blue arrow, Fig. 3E) were uniformly J-aggregate positive (Fig. 3G), indicating that the corresponding population of mitochondria was high polarized.

**Photosensitization of JC-1-stained embryos**

The extent to which mitochondrial polarity may be related to cell function and behavior was investigated by JC-1 photosensitization of day-4.5 blastocysts, which were illuminated in the FITC or RITC channels and then returned to culture. Preliminary experiments demonstrated that 8-s exposure had differential and cell-type-specific sublethal effects of developmental significance, and this duration became standard. Normal patterns of BrdU incorporation (which reflects the ability of the cells to replicate DNA) detected by fluorescence microscopy in day-4.5 blastocysts (Fig. 4A; n = 20) are shown in Fig. 4B, with nuclear staining observed in all embryonic compartments (ICM, mural (MT) and polar trophectoderm (PT)). Blastocyst outgrowths from these embryos (e.g. Fig. 4C, OG day 2) composed of a characteristic ICM core (asterisk) and a highly flattened peripheral trophectodermal monolayer (arrow), routinely showed nuclear BrdU staining throughout the culture (Fig. 4D; n = 26). Exposure of JC-1-stained embryos (Fig. 4E) in the FITC channel (n = 40) had no discernible effect on BrdU incorporation (Fig. 4F), the timing or extent of outgrowth (OG day 2, Fig. 4G), or BrdU incorporation (Fig. 4H). In contrast, exposure in the RITC channel (Fig. 4I) resulted in no detectable BrdU incorporation in the mural trophectoderm, but normal levels of incorporation in the ICM and polar trophectoderm (Fig. 4J; n = 55). Comparable frequencies of outgrowth (>95%) were observed for control blastocysts (47/50) and JC-1-stained blastocysts illuminated in the FITC channel (40/42; e.g. Fig. 4G).

In the first study of embryos illuminated in the RITC channel, 40% (18/44) initiated outgrowth during the first 24–36 h of culture and when observed after 48 h, the trophectodermal component contained few cells (Fig. 4K).
Figure 3 Differences in mitochondrial polarity in cells derived from the ICM component of blastocyst outgrowths are shown in these fluorescent images of JC-1-stained cells. (Panel A) Typical distribution of cells containing high- (trophectoderm, TR) and low-polarized ICM indicated by asterisk) mitochondria in an undisturbed outgrowth from a normal day-4.5 embryo after 1.5 days of culture. (Panel B) Representative example of an ICM core from a nascent outgrowth that was mechanically dissected from the outgrowth (asterisk, panel A), and stained with JC-1 during dissociation (panel B). Peripheral cells showing apical J-aggregate fluorescence are indicated by red arrows in panel C. (Panel D) Four-day-old culture of pooled ICM cores composed entirely of cells that are relatively small, uniform in size and mononucleated. Differences in mitochondrial polarity are cell-contact dependent with those in dense regions containing low-polarized organelles (asterisk, panels D and E). For cells with free margins, the corresponding mitochondrial polarity was either high (blue arrow, panel E, and at higher magnification in panel G) or a mixture of high-and low-polarized organelles (red arrow, panel E, and at higher magnification in panel F). The apparent balance between high- and low-polarized mitochondria in the less dense regions of the ICM-derived cultures was related to the extent of intercellular association.
Figure 4 The typical pattern of BrdU incorporation into the ICM, mural (MT) and polar (PT) trophectoderm in a normal (panel A) day-4.5 mouse blastocyst, and a representative example of incorporation into a typical nascent blastocyst outgrowth (panel D), are shown in these light and fluorescent microscopic images. The ICM is indicated by an asterisk and trophectodermal cells by an arrow. Panels E–M show the effects of photosensitization on BrdU incorporation in JC-1-stained, day-4.5 blastocysts (panels E and I), and representative outgrowths from similar embryos (panels G and K) after illumination in the FITC (panels F and H) or RITC (panels J and L) channels (see text for details). The arrow in panel M shows DAPI-stained nuclei in small mononucleated cells that emerged from outgrowths after several days of culture of JC-1-stained blastocysts illuminated in the RITC channel. Panel O shows a single, TUNEL-positive nucleus in a normal day-4.5 blastocyst (panel N), and its apparent location within the embryo after DAPI staining is indicated by an arrow in panel P. Panel Q is an RITC-illuminated, JC-1-stained blastocyst that remained intact for 4 days after exposure. A single fluorescent nucleus indicating guanosine oxidation was detected after staining with Texas Red Avidin. All nuclei were fluorescent after exposure to peroxide (panel S). In comparison to RITC-channel-exposed, JC-1-stained blastocysts that initiated outgrowth on day 1 after exposure (e.g. panel K), similar embryos that remained intact for several days (e.g. panel Q) showed a relatively normal pattern of outgrowth on day 1 (panel T), although the intensity and distribution of J-aggregate-positive cells was reduced when compared with untreated embryos (e.g. Figures 1S and 2A). However, by day 2, the appearance of the cultures was characteristic of blastocyst outgrowths (panel V).
and BrdU incorporation was confined to the ICM core (Fig. 4L). However, small, mononucleated cells (arrow, Fig. 4K) first appeared around the ICM core on OG day 3 (DAPI-stained nuclei, white arrow, Fig. 4M) and, by light microscopy, were phenotypically comparable to cells derived from the disaggregated ICM cores and their progeny, as described above (Fig. 3D). A second series of exposures was undertaken to confirm these findings with a larger number of embryos; approximately 60% (59/105) remained intact and unattached for as many as 5 days of culture (e.g. day 4, Fig. 4Q), and, during this time, showed no apparent increase in trophodermal or ICM cell numbers (nuclear staining comparable to Fig. 1G). However, approximately 60% (36/59) of these embryos initiated outgrowth at days 3-4 of culture (e.g. Fig. 4T, OG day 1), but, unlike their siblings, which outgrew earlier (e.g. OG day 1, Fig. 4K), trophodermal cells increased in number during time-lapse-monitored culture (OG day 2, Fig. 4V), and the pattern of BrdU staining was comparable to those observed days earlier in outgrowths from their FITC-channel-exposed siblings (comparable to Fig. 4H). Like early outgrowths from untreated blastocysts, outgrowths from blastocysts illuminated in the RITC channel showed J-aggregate fluorescence in trophodermal, but not ICM, cells (Fig. 4U). Initially, the J-aggregate signal was punctate and confined to apical regions of outgrowing cells. However, with continued outgrowth (e.g. Fig. 4V), the intensity and distribution of J-aggregate fluorescence were comparable to their normal counterparts (e.g. the pattern shown in Fig. 2B). About 40% (23/59) of these blastocysts failed to outgrow after 5 days of time-lapse-monitored culture (see below).

Analysis of DNA fragmentation and oxidation

As noted previously, photosensitization with mitochondria-specific fluorescent probes has been associated with the production of oxidative free radicals that are thought to induce nuclear DNA fragmentation (detected by TUNEL staining) and eventual apoptotic cell death. To determine whether exposure of JC-1-stained blastocysts in the RITC channel has similar effects, TUNEL and BrdU staining was performed to detect DNA fragmentation and guanosine oxidation respectively. Approximately 30% (11/37) of day-4.5 blastocysts (Fig. 4N) showed one or two TUNEL-positive nuclei (arrow, Fig. 4O), but avidin fluorescence indicative of guanosine oxidation was not detected (image not shown). However, with continued outgrowth (e.g. Fig. 4V), the intensity and distribution of J-aggregate fluorescence were comparable to their normal counterparts (e.g. the pattern shown in Fig. 2B). About 40% (23/59) of these blastocysts failed to outgrow after 5 days of time-lapse-monitored culture (see below).

Analysis of ATP contents in normal and delayed implanted blastocysts

The average net ATP content (±22) of hatched blastocysts on day 4.5 of pregnancy (n = 106) was 127 fmol, and 187 and 196 fmol respectively on days 3 (n = 61) and 5 (n = 61) of diapause. The differences in ATP content between normal and delayed implanted embryos were significant (P = 0.01). The ATP content of JC-1-stained blastocysts illuminated in the FITC (n = 81) and RITC channels (n = 81) was measured after 20–24 h of culture. The average ATP content was 131 fmol (FITC, ±20) and 90 fmol (RITC, ±20) respectively. The average ATP content of blastocysts illuminated in the RITC channel that did not outgrow after 2 (n = 33), 3 (n = 26), 4 (n = 29) or 5 (n = 19) days of culture was 85 fmol (±18), and this lower level was significant (P ≤ 0.01) with respect to untreated blastocysts and JC-1-stained embryos illuminated in the FITC channel. In contrast, the net ATP content of RITC-channel-exposed embryos that showed indications of incipient outgrowth (follicular elaboration of filopodia and bulbous projections; see below) after 3 (n = 25) or 4 (n = 21) days of time-lapse-monitored culture was 138 (±21), a level comparable to the one measured in normal day-4.5 embryos.

Time-lapse analysis

The behavior of hatched blastocysts recovered from the uterus on day 4.5 (n = 25) of pregnancy was investigated by time-lapse microscopy. After 8-10 h of culture (Fig. 5A), small (asterisk), highly motile filopodial projections (arrow) emerged from the abembryonic region of the mural trophoderm. Expanded diapauing blastocysts (n = 31) cultured immediately after removal from the uterus on days 3 (n = 20) or 5 (n = 16) of delayed implantation showed no obvious morphodynamic activity during the first 5 h in vitro (Fig. 5B). However, around 6 h, filopodia (arrow, Fig. 5C) emerged from the abembryonic portion of the mural trophoblast. The dynamic nature of these projections was indicated in time lapse by cycles of evolution and resorption that increased in frequency by 16 h in control blastocysts, and by 12 h of culture in blastocysts released from diapause in vitro (arrow, Fig. 5D). In all instances, focal filopodial formation preceded the establishment of robust outgrowths, and in some time-lapse sequences, filopodia were observed to detach from the embryo and ‘crawl’ on the glass culture surface for several hours (see below).
Figure 5 Static images of morphodynamic processes taken from time-lapse monitoring of normal day-4.5 blastocysts (panel A), blastocysts reactivated from diapause in vitro (panels B–D), and JC-1-stained blastocysts photosensitized in the FITC (left) and RITC channels (right) (panels E and F) are shown. For normal embryos, the emergence of bulbous projections (asterisk, panel A) and filopodia (arrow, panels A, F and G) from the abembryonic portion of the mural trophectoderm preceded robust trophectodermal outgrowth and expansion (TOG, panels I and J). Their appearance in reactivated embryos (panel B) occurred some hours later than in normally progressing embryos. Spatial and temporal aspects of their occurrence in FITC-channel-illuminated embryos (arrows, right-hand embryo, panel F) were normal. In contrast, for RITC-channel exposed, JC-1-stained blastocysts, they often appeared after 2 or more days in culture (arrow, panel J). In these instances, filopodia appeared suddenly (e.g. from 76 h 5 min to 76 h 20 min of culture; panels K and L). Some filopodia detached (panel M) and were observed to ‘crawl’ around the surface of the culture dish for several hours. As with untreated embryos, the elaboration of numerous filopodia preceded outgrowth for these embryos. The time of observation (hours) in panels E–J is indicated in the upper or lower right-hand portion of each image.
For time-lapse studies of photosensitized blastocysts, embryos obtained from the same animal were cultured in pairs, with the FITC- and RITC-exposed embryos placed on the left and right respectively (Fig. 5E–J). Morphodynamic activities of blastocysts illuminated in the FITC were temporally and spatially identical to their unexposed control and reactivated counterparts, and exhibited robust outgrowth, as shown in Fig. 5G–J, which are representative images of the first 48 h of culture. The arrow in Fig. 5F denotes filopodia that had developed about 2 h earlier in the abembryonic trophectoderm. In contrast, most blastocysts illuminated in the RITC channel showed no apparent morphodynamically activity during 3–5 days of culture (n = 38), and while some remained expanded, over 80% (31/38) were in a persistent state of ‘collapse’, which initially occurred with a spontaneous discharge of blastocelic fluid around h 10 of culture (Fig. 5F–J).

In comparison to control blastocysts or embryos reactivated from diapause in vitro, outgrowth for most JC-1-stained blastocysts illuminated in the RITC channel began between days 3 and 5, including those cocultured with siblings illuminated in the FITC channel (see above). Outgrowth was also preceded by the elaboration of filopodia (arrows, Fig. 5K–M), which appeared rather suddenly in embryos that seemed morphodynamically ‘inert’ prior to onset of this activity. In Fig. 5I, the region of the trophectoderm from which filopodia emerged is indicated by an arrow. It is assumed that the abembryonic portion of the mural trophoblast was involved owing to the absence of embryo movement during time-lapse-monitored culture. As described previously, trophectodermal outgrowths from embryos illuminated in the RITC channel were generally less robust if they occurred within the first day or two of culture, but were largely typical of this process if initiated on or after culture day 3.

Discussion

The primary role of mitochondria in the preimplantation-stage embryo has traditionally been considered a metabolic one, that is, to provide ATP by means of oxidative phosphorylation, and adverse effects of numerical or structural defects that occur naturally (Müller-Höcker et al. 1996, Van Blerkom et al. 2000) or are experimentally induced (Eichenlaub-Ritter et al. 2004, Thouas et al. 2004; reviewed by Cummins 2002, Brenner 2004) are usually considered in this context. Mitochondrial polarity is a physiochemical property that results from the outward pumping of protons across the inner mitochondrial membrane, creating a proton gradient with two components, an inner membrane potential (ΔΨm) and a pH gradient; the energy stored in either component drives the conversion of ADP to ATP by respiratory chain enzymes. Differences in the relative magnitude of the membrane potential detected with ΔΨm-specific probes have been suggested to reflect corresponding differences in mitochondrial activity or function that affect the competence of mature oocytes and early embryos in the mouse and man (Wilding et al. 2001, 2002, 2003, Van Blerkom et al. 2002, Acton et al. 2004, Jones et al. 2004, Van Blerkom 2004). For these studies, JC-1 has been the probe of choice owing to the specificity with which it reports high and low ΔΨm in living cells (Salvioli et al. 1977). Comparatively low-polarized mitochondria (under ~140 mV) fluoresce green, reflecting accumulation of the JC-1 monomer in the mitochondrial matrix, while comparatively high-polarized mitochondria (over ~140 mV) fluoresce orange to red, as a result of monomer multimerization into so-called J-aggregates that shift in the emission spectrum to longer wavelengths (for review of JC-1, see Reers et al. 1995).

Differences in the spatial distribution of high- and low-polarized mitochondria in mouse and human oocytes and cleavage-stage embryos have been suggested to represent discrete regions of differential mitochondrial activity with distinct or focal regulatory functions (e.g. ATP production, calcium regulation, post-translation modification; Van Blerkom et al. 2002, Van Blerkom 2004). For blastocyst-stage embryos, Van Blerkom et al. (2002) detected high-polarized mitochondria in the trophectoderm, but not the ICM, and proposed that different states of mitochondrial polarization may reflect corresponding differences in organelle activity or function in the two lineage-determining (progenitor) cell types of the mammalian embryo. Here, the physiologic and possible developmental significance of ΔΨm during the terminal stage of the preimplantation period was investigated further.

ΔΨm in normal mouse blastocysts

The present results confirm and extend our previous findings (Van Blerkom et al. 2002) by demonstrating that the occurrence and intensity of high- and low-polarized mitochondria are location- and cell-type specific in the hatched mouse blastocyst as follows:

1. the ICM was consistently J-aggregate negative, indicating that the corresponding mitochondria are low polarized
2. polar trophectodermal cells overlying the ICM exhibited no detectable J-aggregate fluorescence or an intracellular signal that was scant, indicating that high-polarized mitochondria are largely absent in this region
3. the mural trophectoderm exhibited regional differences in fluorescent intensity that extended from the abembryonic (high) to the peripolar regions (low).

The specificity of JC-1 labeling was confirmed by the absence of J-aggregate fluorescence after treatment with ΔΨm-collapsing proton ionophore FCCP, and by the detection in mural trophectodermal cells of J-aggregate-positive, rod-shaped elements that were equivalent in size and distribution to mitochondria identified with other organelle-specific fluorescent probes. Comparable
patterns, distributions and intensities of J-aggregate fluorescence detected under different staining conditions and after mechanical manipulations of embryos support our previous contention (Van Blerkom et al. 2002) that JC-1 reports actual location- and cell-type-specific differences in embryo mitochondrial polarity. However, the possibility that the magnitude of $\Delta \Psi_m$ is related to mitochondrial fine structure needs to be considered, because, unlike mitochondria, in the oocyte and early cleavage-stage embryo (Van Blerkom et al. 2002), differences in matrix density, the number of lamellar cristae, and the degree of organelle elongation distinguish ICM and polar trophodermal mitochondria from their mural trophodermal counterparts at the late blastocyst stage (Van Blerkom & Motta 1979, Van Blerkom et al. 1979). At present, these differences are not thought to be related to the magnitude of $\Delta \Psi_m$, because the undeveloped pericortical mitochondria in the oocyte and early cleavage-stage embryo, which are small, spherical organelles with a few, short cristae that surround, but rarely penetrate, a matrix of very high-density, are J-aggregate positive (Van Blerkom et al. 2002). The occurrence of J-aggregate-positive cells in ICM out-growths (see below) is a further indication that low $\Delta \Psi_m$ is not an intrinsic property of their mitochondria.

$\Delta \Psi_m$ in delayed implanted mouse blastocysts

Aerobic metabolism with glucose as the primary metabolic substrate is the major pathway of energy production in the mouse blastocyst, and respiration from fully developed mitochondria accounts for over 85% of all ATP produced (Benos & Balaban 1983, Trimarchi et al. 2000). The facultative delayed implantation mouse model was used to study the relationship between ATP production and $\Delta \Psi_m$ because mitochondria are metabolically suppressed during the diapause. The decreased energy requirement for diapausing mouse blastocysts results from significantly reduced levels of transcription, translation and protein phosphorylation, and the cessation of DNA replication and cell division (Van Blerkom et al. 1979, Weitlaf & Kiessling 1980, Hamatani et al. 2004, Lopes et al. 2004). Nieder and Weitlaf (1984) showed that diminished mitochondrial respiratory activity in diapausing mouse blastocysts is associated with significant elevation in cytoplasmic ATP content, and proposed that a constitutively high ATP content caused allosteric inhibition of phosphofructokinase and coincident reduction in glucose oxidation. Increased mitochondrial metabolism detected some hours after blastocyst reactivation could be a consequence of the reversal of this inhibition for the existing enzyme combined with translation from nascent phosphofructokinase transcripts (Hamatani et al. 2004). Our finding of elevated levels of ATP in diapausing blastocysts supports the notion of allosteric inhibition advanced by Nieder and Weitlaf (1984).

Changes in $\Delta \Psi_m$ (high to low) have been reported for early mouse and human embryos that develop abnormally or arrest prematurely in vitro (Wilding et al. 2002, 2003, Acton et al. 2004), are developmentally compromised after photosensitization of mitochondria stained with r123 (Thouas et al. 2004), or undergo iatrogenically induced reductions in polarity under certain culture conditions (Van Blerkom et al. 2003) or after cryopreservation (Ahn et al. 2002, Jones et al. 2004). These changes are largely considered in the context of a coincident reduction in mitochondrial metabolic activity, and for diapausing embryos, we had anticipated a downward shift in $\Delta \Psi_m$ owing to the suppression of mitochondrial respiratory activity. However, during the 7 days of delayed implantation examined in the present study, the location, cell-type specificity and relative intensity of J-aggregate fluorescence remained unchanged in embryos with elevated cytoplasmic ATP contents. A rapid upregulation of mitochondrial metabolic activity during JC-1 staining is an unlikely explanation for this finding because the normal pattern of J-aggregate fluorescence was detected within the first 5 min of staining in medium devoid of metabolic substrates. The translation of functional respiratory enzymes from nascent transcripts seems equally unlikely to occur during this time (Hamatani et al. 2004); we suggest that the detection of J-aggregate fluorescence reflects the in vivo state and that high polarity in mural trophodermal mitochondria is maintained during the diapause. Because ATP is required for a potential difference to exist across the inner mitochondrial membrane, the relatively high ATP content of diapausing mouse blastocysts may be sufficient simultaneously to suppress mitochondrial metabolism and support high $\Delta \Psi_m$. A similar situation exists in p$^0$ cells, whose mitochondria have neither DNA nor a functional respiratory chain cells, yet consume about 13% of the ATP produced by glycolysis to maintain a $\Delta \Psi_m$ of ~110 mV (Appleby et al. 1999). Possible reasons why high polarity is retained are discussed below.

$\Delta \Psi_m$ and the regulation of development during diapause

Our results raise several questions concerning how mitochondria may be involved in the regulation of development during the peri-implantation period; in particular, whether high-polarized mitochondria have specialized functions in this regard, especially during the diapause. One possibility under investigation is related to mitochondrial regulation of calcium homeostasis. Calcium is a ubiquitous regulator of numerous cell functions, including control of the cell cycle, cytoskeletal/cytoplasmic remodeling, levels of mitochondrial respiration, morphodynamic movements, and, through various signal transduction pathways, differential gene expression and cell death (Duchen 2000, Rutter & Rizzuto 2000, Bertridge et al. 2003, Webb & Miller 2003). All of these processes are either downregulated or arrested during the diapause (Van Blerkom et al. 1979, Hamatani et al. 2004, Lopes et al. 2004). Also well known is the...

To test this possibility, we measured in preliminary studies changes in levels of cytoplasmic free calcium after mitochondrial depolarization was chemically induced in normal day-4.5 embryos and on day 5 of delayed implantation (unpublished). Embryos were preloaded with the fluorescent calcium reporter Fluo-4 AM in medium free of calcium, magnesium and metabolic substrates (Van Blerkom et al. 2003, Jones et al. 2004), and then exposed to the proton ionophore FCCP at concentrations shown to dissipate \( \Delta \Psi_m \) abruptly and induce a significant increase of intracellular free calcium in mature mouse oocytes (Van Blerkom et al. 2002, 2003). Changes in cytoplasmic free calcium levels were determined quantitatively by measuring the increase in relative fluorescence intensity (above the pre-FCCP baseline) by scanning laser confocal microscopy. After mitochondrial depolarization, the increase in free calcium in diapausing embryos was 3-4 times higher than in their normal counterparts. If these initial findings are confirmed, it could suggest that high polarity is maintained during the diapause in order to sequester calcium in mitochondria such that cytoplasmic levels are reduced, but can return to ‘normal’ (baseline) when delayed implantation ends. Although speculative, mitochondrial calcium sequestration and release in diapausing/reactivated embryos may be associated with the down- and upregulation respectively of regulatory proteins whose function or activity is calcium dependent (Hamatani et al. 2004). In this regard, it will be of interest to determine whether changes in free calcium occur after reactivation and are temporally correlated with the upregulation or reinitiation of biosynthetic, cell-cycle (Van Blerkom 1979, Weitlauf & Kiessling 1980, Lopes et al. 2004) and morphodynamic activities (see below).

**Differential effects of mitochondrial photosensitization**

Photosensitization of mitochondria stained with organelle-specific fluorescent dyes has been widely used to study the cellular and developmental consequences of 1. perturbing electron transport and reducing metabolic activity, and 2. inducing the generation of reactive oxidative species at levels that cause structural damage to mitochondria and DNA, or the release of proapoptotic factors. For these studies, preloading with a fluorescent probe is usually followed by an acute exposure to high-intensity ultraviolet illumination at appropriate excitation and emission frequencies, for times sufficient to have detectable pathophysiologic effects. When the developmental sequelae are considered in the context of the central role mitochondria have in cell function, it is not surprising that the downstream effects of altered organelle function in oocytes and early embryos (Lane & Gardner 2005) induced by photosensitization (Thouas et al. 2004) are largely dose-dependent and often developmentally lethal.

Here, we used photosensitization to derive a sublethal exposure for JC-1-stained blastocysts to determine whether cells containing high- or low-polarized mitochondria are differentially affected. Illumination in the FITC channel had no discernible effects, while exposure in the RITC channel reduced the net ATP content, inhibited BrdU incorporation in the (high-polarized) mural trophoderm, and delayed or precluded outgrowth. However, outgrowth-arrested embryos showed no increase in the frequency of nuclei exhibiting fragmented DNA or indications of guanosine oxidation, indicating that this protocol did not induce apoptosis or other forms of lethal cytopathology. We suggest that putative fluorescence resonance of the J-aggregate multimers in high-polarized mitochondria may cause an acute, but not irreversible, reduction in mitochondrial activity that appears to include ATP generation. Whether the magnitude of \( \Delta \Psi_m \) was affected in high-polarized mitochondria could not be determined under the present conditions, because, once formed, J-aggregates are metastable structures that continue to fluoresce, albeit at reduced intensity, even if \( \Delta \Psi_m \) is dissipated by FCCP or fixation (Van Blerkom et al. 2002). Likewise, it is unknown whether the arrest of cell division in the ICM of embryos in outgrowth-delay for several days indicates a direct effect of photosensitization on low-polarized mitochondria. However, photosensitization does seem to induce a developmental ‘quiescence’ with cellular and phenotypic similarities to the natural diapause. Whether the reduced cytoplasmic ATP content measured during delayed outgrowth is sufficient to down-regulate critical cellular activities (such as DNA synthesis and cytokinesis) that could induce a diapause-like state remains to be determined.

Our studies show that mitochondria have a central regulatory function during the peri-implantation stage that mediates the ability of mouse embryo to develop progressively. Cellular domains containing high-polarized mitochondria may be particularly important in this regard owing to their spatial association with morphodynamic processes that precede implantation. For example, the initial stages of implantation in the mouse involve dynamic activities localized to abembryonic mural trophoderm that are characterized by the elaboration of focal clusters of filopodia and bulboous protrusions. The filopodia are thought to establish the first stable contacts with the endometrial epithelium, and they later physically participate in the implantation process (Potts 1968, Bergstrom & Nilsson 1976). Trophoblast giant cells, which become a terminally differentiated, nondividing (polyploid) cluster of cells involved in the invasive phase of
implantation (Maris et al. 1988), are first identified in the day-5 mouse embryo as ‘bulbous projections’ from the abembryonic trophoderm (Kaufman 1983). Here, time-lapse analysis showed the elaboration of filopodia and bulbous projections from this region in normal (on day 5 in vitro), reactivated, and FITC-channel-exposed, JC-1-stained blastocysts. In all instances, active filopodial formation preceded robust trophodermal outgrowth. In contrast, most JC-1-stained blastocysts illuminated in the RITC channel showed no apparent morphodynamic activity after several days of coculture with normally out-growing siblings that had been illuminated in the FITC channel. However, when ‘delayed’ outgrowth was initiated, it, too, was always preceded by the appearance of filopodia and bulbous projections. Owing to the retention of residual J-aggregate fluorescence, we could not determine quantitatively whether this morphodynamic activity is preceded by a focal change in $\Delta \Psi_{m}$. For embryos in which comparatively normal outgrowth occurred after a delay of several day, the results indicate that DNA replication, cell division, filopodial formation and outgrowth may await the restoration of ‘normal’ mitochondrial function. This interpretation was supported by cytoplasmic ATP contents at levels comparable to day 4.5 in these embryos, where the appearance of filopodia indicated outgrowth was imminent. Restoration of mitochondrial activity or function to near normal levels may be especially relevant to the expression of morphodynamic activities in the abembryonic region, which typically displays higher relative intensities of J-aggregate fluorescence.

**Cell-type-specific $\Delta \Psi_{m}$ in the peri-implantation blastocyst may be determined by extrinsic factors**

Our previous studies suggest that within oocytes and blastomeres, spatial differences in mitochondrial polarity are cell-contact associated rather than a reflection of differential metabolic metabolism (Van Blerkom et al. 2002, 2003). The present results indicate that a similar situation may exist at the blastocyst stage. Trimarchi et al. (2000) found oxygen consumption to be uniform along the entire circumference of the hatched mouse blastocyst, and concluded that regional differences in levels of oxidative metabolism probably do not exist. If this accurately reflects underlying metabolic activity, cellular domains of high and low $\Delta \Psi_{m}$ would appear to be unrelated to corresponding differences in mitochondrial oxidative phosphorylation, and this appears to be the case during the diapause as well.

A nonmetabolic origin for differential polarity in the peri-implantation mouse embryo is suggested by the finding that within cells, high or low polarity is related to the extent and nature of intercellular contact and communication. Diaz et al. (1999) showed for several differentiated cell lines that high-polarized (J-aggregate-positive) mitochondria occurred in the subplasmalemmal cytoplasm at the free margins of cells, whereas low-polarized (J-aggregate-negative) mitochondria occurred in the interior of the cytoplasm and in subplasmalemmal zones where intercellular contacts existed. The state of polarity was found to be a dynamic one in which the loss of intercellular contact was associated with a shift from low to high $\Delta \Psi_{m}$, while the opposite occurred when such contacts were established. Because the relative magnitude of $\Delta \Psi_{m}$ in subplasmalemmal mitochondria can be influenced by ionic and electrical fluxes within and between cells (Ichas et al. 1997, Diaz et al. 1999), the occurrence of high-polarized mitochondria at free margins may reflect a need to ‘work harder’ to regulate calcium homeostasis than their counterparts deeper within the cytoplasm or where cells membranes are in direct contact and ionic homeostasis is facilitated by specialized (gap) junctions (Diaz et al. 1999).

A similar relationship with mitochondrial polarity has been proposed for the oocyte and cleavage-stage embryo (Van Blerkom et al. 2003, Van Blerkom 2004); high-polarized mitochondria occur at the free margins and low-polarized mitochondria are located internally; however, for the cleavage-stage embryo, low-polarized mitochondria also occur in the subplasmalemmal cytoplasm subjacent to zones of contact (Van Blerkom et al. 2002), where intercellular communication involves gap junctions (Van Blerkom & Motta 1979). The dynamic nature of this spatial pattern was demonstrated by disaggregating two- and four-cell mouse embryos and repositioning and realigning blastomeres such that formerly J-aggregate-negative cortical domains became positive, and vice versa (Van Blerkom et al. 2002).

The same types of intercellular interactions that influence the magnitude of $\Delta \Psi_{m}$ in somatic cell cultures exist in the hatched blastocyst. For example, the elongated apical and basal surfaces of the abembryonic mural trophectoderm are free, but toward the peripolar region, focal contacts with early migrating endodermal cells are established (Van Blerkom et al. 1979). The basal surface of the polar trophectoderm fully associates with the ICM, whose cells are largely in close contact throughout their margins (Van Blerkom & Motta 1979, Makabe & Van Blerkom 2006). The occurrence and relative intensity of high-polarized trophodermal mitochondria coincides with the presence or absence of subjacent cells and where present, the extent of association. Similar to the situation observed in cultured cells, gap junctions that regulate the flow of ions between cells exist between ICM cells and between the polar trophectoderm and ICM (Van Blerkom & Motta 1979). In this respect, gap-junction functionality and density, combined with the degree of intercellular contact, may largely determine cell-type-specific $\Delta \Psi_{m}$ within the peri-implantation blastocyst. The present studies of undisturbed blastocyst outgrowths, disaggregated ICM cores and proliferating ICM-derived cells, demonstrate that low polarity is not an intrinsic property, but one related to the extent of intercellular contact. We suggest that cell-
location-specific mitochondrial polarity at the blastocyst stage is determined by the normal intercellular morphodynamics of cavitation and blastocoele expansion. Motosugi et al. (2005) recently proposed that these same intercellular dynamics alone lead to the specification of the first developmental polarity in the mouse embryo, the establishment of the embryonic–abembryonic axis. It will be of interest to determine whether other regulatory functions of mitochondria (e.g., oxygen sensing) whose activity may be \( \Delta \Psi_m \) associated are involved in the molecular specification of cell types, or influence their corresponding state of differentiation (trophoderm) or totipotency (ICM) at the blastocyst stage.

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