Unequal distribution of 16S mtrRNA at the 2-cell stage regulates cell lineage allocations in mouse embryos

Zhuxia Zheng1,2, Hongmei Li3, Qinfen Zhang3, Lele Yang1 and Huayu Qi1

1Key Laboratory of Regenerative Biology, Guangdong Provincial Key Laboratory of Stem Cell and Regenerative Medicine, South China Institute for Stem Cell Biology and Regenerative Medicine, Guangzhou Institutes of Biomedicine and Health, Chinese Academy of Sciences, A303, 190 Kaiyuan Boulevard, Science City, Huangpu District, Guangzhou, Guangdong 510630, China, 2School of Life Science, University of Science and Technology of China, Hefei 230026, China and 3State Key Lab for Bio-control, School of Life Sciences, Sun Yat-sen University, Guangzhou 510275, China

Correspondence should be addressed to H Qi; Email: qi_huayu@gibh.ac.cn

Abstract

Cell lineage determination during early embryogenesis has profound effects on adult animal development. Pre-patterning of embryos, such as that of Drosophila and Caenorhabditis elegans, is driven by asymmetrically localized maternal or zygotic factors, including mRNA species and RNA binding proteins. However, it is not clear how mammalian early embryogenesis is regulated and what the early cell fate determinants are. Here we show that, in mouse, mitochondrial ribosomal RNAs (mtrRNAs) are differentially distributed between 2-cell sister blastomeres. This distribution pattern is not related to the overall quantity or activity of mitochondria which appears equal between 2-cell sister blastomeres. Like in lower species, 16S mtrRNA is found to localize in the cytoplasm outside of mitochondria in mouse 2-cell embryos. Alterations of 16S mtrRNA levels in one of the 2-cell sister blastomeres via microinjection of either sense or anti-sense RNAs drive its progeny into different cell lineages in blastocyst. These results indicate that mtrRNAs are differentially distributed among embryonic cells at the beginning of embryogenesis in mouse and they are functionally involved in the regulation of cell lineage allocations in blastocyst, suggesting an underlying molecular mechanism that regulates pre-implantation embryogenesis in mouse.

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Introduction

Embryogenesis following fertilization lays down the foundation for future animal development. During early embryogenesis, descendent of the totipotent zygote (the 1-cell embryo that formed by sperm–egg fusion) gradually lose their developmental potential and acquire specific cell fate. Understanding the molecular inner works that underlie the regulation of cell fate determination during early embryogenesis has been one of the central issues in developmental biology (Johnson 2009, Rossant & Tam 2009, St Johnston & Ahringer 2010, Nance 2014, Du et al. 2015).

In lower species, such as Drosophila and Caenorhabditis elegans, embryogenesis is believed being pre-patterned. Unfertilized eggs and zygotes are highly polarized cells with distinctive cellular geometry, containing predisposed cell fate determinants that are asymmetrically localized in the cell, including various RNA species and RNA binding proteins (Johnstone & Lasko 2001, St Johnston & Ahringer 2010). Differentially segregated cell fate determinants can drive the inherited cells into specific cell lineages, such as the germ line cells (Johnstone & Lasko 2001, Kimble & Crittenden 2007). The early cellular polarity of eggs or zygotes can be translated into axes of developing embryos, such as the animal–vegetal and anterior–posterior axes, which are strongly correlated with the development of body axes in adult animals. Although the underpinning molecular mechanisms of these correlations remain to be fully elucidated, cell lineage formation during early embryogenesis in lower species is profoundly influenced by pre-existing intrinsic factors, mostly inherited from the maternal source.

In mammals, the establishment of distinguishable cell lineages following fertilization is thought to occur first at the blastocyst stage during pre-implantation embryogenesis (Rossant & Tam 2009). Mammalian blastocyst initially contains two cell-lineages: the inner cell mass (ICM) and the trophectoderm (TE). With development, ICM cells are further differentiated into epiblast (Epi) that will give rise to the embryo proper and primitive endoderm (PrE) that together with the TE will generate extra-embryonic tissues, including placenta. It is postulated that, in mouse, the blastocyst is bi-laterally symmetric and can be divided into embryonic half
First found in Drosophila, one of the maternal cell fate determinants that influence the formation of germ cells in invertebrates is the posterior mitochondrial ribosomal RNAs (mtrRNAs) that are localized outside of the organelle (Kobayashi et al. 1993, Iida & Kobayashi 1998). Examination of mtrRNAs in other species, including Xenopus and Sea urchin, also showed that they are localized asymmetrically and outside of mitochondria in early embryos, suggesting their possible functions as early cell fate determinants in these species (Kobayashi et al. 1998, Ogawa et al. 1999). Investigation of mtrRNAs in mouse pre-implantation embryos suggested that they are localized asymmetrically in the cytoplasm of MII oocytes toward the animal pole where the first polar body emits. Similar to that in lower species, 16S mtrRNA was also found in the cytoplasm outside of mitochondria in MII oocytes (Ninomiya & Ichinose 2007). However, it was suggested that mtrRNAs were absent in mouse 2-cell embryos and no difference was found in their distribution among blastomeres of pre-implantation embryos. Their functional roles during cell lineage specification in mouse have yet to be determined. In the present study, we re-examined the expression and distribution patterns of mtrRNAs in mouse pre-implantation embryos. Results showed that mtrRNAs are constantly expressed during pre-implantation embryogenesis. Both small and large mtrRNAs are unequally distributed between 2-cell sister blastomeres, suggesting a molecular variation between cells at the beginning of embryogenesis in mouse. Further functional analyses suggested that alterations of levels of large subunit 16S mtrRNA in 2-cell blastomeres functionally influence cell lineage allocations in the blastocyst.

Materials and methods

Animal handling and mouse embryos

CD1 mice were used for embryo isolations for most of the experiments, except that when testing the hormonal effects on embryos, C57BL/6 mice were also used. Adult female mice (1.5–to 2-month of age) were superovulated by i.p. injection of 10 IU of pregnant mare serum gonadotropin (PMSG) per mouse, followed by 10 IU of human chorionic gonadotropin (HCG) 48 h later. Injected female mice were placed with adult male mice overnight and checked for copulation plugs the following morning. Animals were sacrificed by cervical dislocation following CO₂ anesthetization. Embryos at different stages were isolated in pre-warmed M2 media (Sigma M7167) and washed in KSOM media (Specialty Media, MR-020P-D). Unfertilized eggs were isolated from females without mating 16–18 h post HCG injection. Zygotes, early 2-cell, middle 2-cell, late 2-cell, 4-cell, 8-cell, morula and blastocyst embryos were collected 24–26 h, 31–32 h, 45–46 h, 52–53 h, 56–60 h, 67–68 h, 77–78 h and 90–92 h post HCG injection. As controls, CD1 mice that were naturally mated and superovulated C57BL/6 mice were also used for the isolation of embryos. To isolate single blastomeres from 2-cell and 4-cell embryos, embryos were placed in Tyrodé's...
solution to remove ZP and blastomeres were washed and separated by pipetting in PBS/PVP-40 (4 mg/ml). Separated blastomeres were then transferred into PCR tubes. All animal husbandry and usage were carried out according to the guidelines of IACUC and approved by the Animal Care and Use Committee of Guangzhou Institutes of Biomedicine and Health, Chinese Academy of Sciences (Permit number: 2008023).

**cDNA cloning and real time quantitative RT-PCR**

cDNAs of 12S and 16S mtrRNA were cloned from mouse mitochondria genome extracted from CD1 mouse ovary using Mini Plasmid Extraction kit (Tiangen, Beijing, China). 12S mtrRNA cDNA was amplified using RT-PCR with a 5’-primer containing T7 promoter sequence for *in vitro* transcription. 16S mtrRNA cDNA was cloned into pcDNA expression vector (Promega) between SalI/NotI sites under the T7 promoter. Plasmid pEGFP-N2 containing EGFP coding sequence was used for PCR and *in vitro* transcription of Egfp mRNA.

For the examination of 16S mtrRNA and Actin expression in mouse pre-implantation embryos, 15 embryos at each pre-implantation stage were collected from superovulated CD1 mice. Embryos of each group were lysed in 10 μl lysis buffer (DEPC-H2O containing 0.1% Triton X-100, 0.1 M DTT and 1 μl RNase OUT Recombinant RNase inhibitor, 5000 U, Invitrogen). Same amount of *in vitro* transcribed Egfp mRNA was added to each sample as internal control. The RT reaction was carried out with an adapter–oligo–dT24 primer at 50°C for 1.5 h. Real time quantitative PCR was done with SYBR Green Mix (Takara, Japan) in optical 96-well reaction plates on a CFX96 Realtime System (Bio-Rad). PCRs were performed in triplicate and signals obtained were normalized against that of Egfp using the following equation: 2^((C_{Egfp} – C_{GeneX}).

Single-cell quantitative RT-PCR, expression of mitochondrial genes and the expression of 16S mtrRNA following micro-injection were examined in the similar fashion using endogenous Actin as control. PCRs were performed in triplicate and signals obtained were normalized against that of Actin using the following equation: 2^((C_{Actin} – C_{GeneX}).

Relative gene expression levels were calculated using Excel software. Primers used for cDNA cloning and qRT-PCR are summarized in Supplementary Table S1. Detailed experimental procedures are provided in Supplementary Materials and Methods.

**Whole-mount *in situ* hybridization of mouse embryos**

Briefly, digoxigenin (DIG)-labeled RNA probes were transcribed *in vitro* using T7 RNA polymerase (Takara) and DIG RNA Labeling Mix (Roche), and purified and stored in DEPC-H2O at −80°C. Collected embryos were fixed and stained with DIG-labeled cRNA antisense or sense probes (1 μg/ml in pre-hybridization solution). They were then washed and incubated with anti-DIG alkaline phosphatase (AP) conjugates (Roche) at 1:2000 dilution in 1% bovine serum albumin (BSA)/1× Phosphate Buffered Saline containing 0.1% Tween20 (PBST) for 2 h. They were further washed before transferred into Staining buffer containing BCIP/NBT solution. Stained embryos were then mounted onto glass slides in 50% glycerol/PBS and examined with an inverted microscope (Olympus IX71). ZP surrounding embryos were dissolved during hybridization. All procedures were carried out at room temperature unless indicated. Detailed experimental procedures are provided in Supplementary Materials and Methods.

Whole mount *in situ* hybridization (ISH) was carried out for the following genes: Actin, 12S mtrRNA, 16S mtrRNA, cytochrome b (Cytb), NADH dehydrogenase subunit 2 (Nd2), cytochrome c-1 (Cyc1), glyceraldehyde-3-phosphate dehydrogenase (Gapdh), hypoxanthine guanine phosphoribosyl transferase (Hprt), high mobility group box3 (Hmgb3), TATA box binding protein (TBP)-associated factor 9 (Taf9) and immediate early response 5 (Ier5). Primers used for amplifying probes are summarized in Supplementary Table S1.

**ISH at electron microscopic level**

ISH at the electron microscopic level (ISH-EM) was carried out according to the procedures described before (Yamashita et al. 2009), with some modifications. Briefly, isolated embryos were first fixed and hybridized with 16S mtrRNA long cRNA probes as described above. They were then processed for thin (100 nm) sections cut with an Ultramicrotome (Leica Instruments, Wetzlar, Germany) and mounted onto nickel grids. Grids were subsequently stained with primary mouse monoclonal anti-DIG antibody (1:200, Sigma) and secondary goat anti-mouse IgG conjugated with 12-nm colloidal gold (1:20, Jackson ImmunoResearch, West Grove, PA, USA). Stained samples were examined using an electron microscope (Tecnai G2 Spirit, FEI). Average numbers of colloidal gold particles on sections of stained embryos were counted and calculated for average from ten frames of electron micrographs at 16 800× magnification. All procedures were carried out at room temperature unless indicated. Transmission electron microscopy of thin (100 nm) embryo sections without ISH was also carried out according to the method described previously (Zhang et al. 2004). Detailed experimental procedures are provided in Supplementary Materials and Methods.

**Fluorescent ISH and immunofluorescent staining of mouse embryos**

Stellaris oligonucleotide probe sets (20-mers) that cover the entire lengths of 12S mtrRNA and 16S mtrRNA were synthesized and tagged with Quasar 570 Dye and Quasar 670 Dye (Biosearch Technologies, Petaluma, CA, USA) respectively. Collected mouse embryos were fixed and stained with 125 nM Stellaris probes, either singly or together. Stained embryos were then mounted onto glass slides via free access.
Live embryos were stained with MitoTracker Red (200 μM, Molecular Probes) for detecting mitochondria content and tetramethyl rhodamine methyl ester (TMRM, 25 μM, Molecular Probes) for detecting mitochondrial membrane potential, with or without pre-treatment of protonophore carbonyl cyanide p-trifluoromethoxyphenyl-hydrazone (FCCP). Hoechst 33342 (10 μg/ml) was used for nuclei staining for 30 min. In some cases, embryos stained with MitoTracker Red were further processed for fluorescent ISH (FISH) of mtrRNAs using Stellaris probes as described above. Embryos microinjected with 16S mtrRNA sense or anti-sense RNAs, together with Egfp mRNA, were also stained with TMRM in the same way 8 h post-injection. Stained embryos were examined with Confocal Laser Scanning Microscopy (CLSM, LSM 710 NLO, Carl Zeiss, Jena, Germany).

Primary antibodies used were: rabbit polyclonal anti-E-cadherin (Abcam, 1:50); mouse monoclonal anti-CDX2 (Abcam, ab115595, 1:50); rabbit polyclonal anti-NANOG (Abcam, ab80892, 1:100) and TRITC-Phalloidin (Sigma, 1:5000). Respective secondary antibodies used were: goat-anti-rabbit FITC conjugate (Molecular Probes, 1:1000); Goat-anti-mouse Alexa Fluor568 (Molecular Probes, 1:1000) or goat-anti-rabbit Alexa Fluor656 (Molecular Probes, 1:1000). Detailed experimental procedures are provided in Supplementary Materials and Methods.

**RNA microinjection of 2-cell embryos and cell allocation analysis**

mRNAs of Egfp were transcribed in vitro using mMessage mMachine SP6/T7 Ultra Kit with Capping analogous (Ambion) from linearized pCS2 vector containing EGFP open reading frame (ORF) with human serum albumin 5' and 3'-UTRs. Full-length 16S mtrRNA sense or antisense RNAs were transcribed in vitro using T7 promoter with the same kit without Capping analogous. Transcribed RNAs were purified using RNase Mini Kit (Qiagen), re-suspended in DEPC-H2O and stored at −80 °C. Ten fragments (100–200 nt in length) of anti-sense RNAs, spanning the entire 16S mtrRNA sequence were prepared and mixed together for microinjection. Single blastomere of a 2-cell embryo was microinjected with either Egfp mRNA (0.8 μg/μl) alone, or together with 16S mtrRNA sense (1.5 μg/μl, 2:1) or anti-sense RNAs (1.5 μg/μl, 2:1) respectively.

Injected embryos were cultured in vitro in KSOM for 46–48 h till blastocyst stage in a humidified incubator containing 5% CO2, 37 °C. Embryos were then fixed in 4% PFA/PBST and stained with DAPI (0.5 μg/ml). Each embryo containing EGFP-labeled (EGFP+) cells was scanned at 2-μm intervals using CLSM. Scanned images from each embryo were composed together and examined at three dimensions using ZEN 2010 software (Carl Zeiss). Total number of cells and positions of individual cells within an embryo were determined by their nuclear DAPI staining, in relation to the surrounding cells.

Using ortho module of the software, cells that were located on the surface of the embryo along either one of the three axes (X, Y and Z) were designated as TE cells, whereas cells that are located on the inside of the embryo and surrounded by other cells along all three axes were assigned as ICM cells. Positions of EGFP+ cells were determined by superimposing the green fluorescence over DAPI staining. Total number of cells with either EGFP signal or CDX2 staining or both were counted in the same way. Alternatively, distributions of EGFP+ progeny cells in either embryonic, abembryonic or both sides within blastocysts were examined using 3-D confocal images. Embryos were grouped into three different patterns according to the allocations of EGFP+ cells within them. Pattern 1 contains EGFP+ cells in the embryonic side only; pattern 2 contains EGFP+ cells in both embryonic and abembryonic sides and pattern 3 contains EGFP+ cells in the abembryonic side only. Total numbers of cells with or without EGFP in either ICM or TE and patterns of embryos following microinjection were counted and calculated using Excel software. Cell numbers surveyed were summarized in Supplementary Tables, see section on supplementary data given at the end of this article.

**Statistical analyses**

Signal intensities of whole-mount ISH were measured against background using single blastomere from light micrographs using Image-Pro 6 software. Fluorescent signals, including that of FISH, were quantified from entire stacks of confocal sections for each sister blastomere using ImageJ software. Ratios of signal intensities (R=OD1/OD2, OD1> OD2, OD: optical density measured for each sister blastomere) between 2-cell sister blastomeres were then calculated using Excel. Statistical significances of signal variances between experimental and control groups were assessed using non-parametric Mann–Whitney test for independent samples. Alternatively, significance of variation was also examined using z-test for two samples' means. One-way ANOVA and post-hoc Turkey test were performed to assess the significance of differences when there were more than two groups of data in the experiment using QIMacros software. χ2 test was performed for assessing the significance of changes of embryonic patterns following microinjection. Significance was set as P<0.05. Data are presented as mean ± s.d.

**Results**

**Expression of mtrRNAs in mouse pre-implantation embryos**

The expression of 16S mtrRNA in mouse oocytes and pre-implantation embryos following fertilization was first examined using quantitative RT-PCR. In order to have an internal control with constant quantity, same amount of in vitro transcribed Egfp mRNA was added to each RNA sample extracted from embryos at various stages before RT reaction. Quantitative RT-PCR was then used to examine the expression of 16S mtrRNA. It was found that levels of 16S mtrRNA remained constant in MI oocytes, zygotes and 2-cell embryos and increased dramatically after 4-cell stage, similar to the expression pattern of Actin control (Fig. 1D). No apparent decrease of 16S mtrRNA was found in 2-cell embryos.

To confirm the expression of mtrRNAs in pre-implantation embryos, ISH of both small (12S) and Supplementary Materials and Methods.

mRNAs of Egfp were transcribed in vitro using mMessage mMachine SP6/T7 Ultra Kit with Capping analogous (Ambion) from linearized pCS2 vector containing EGFP open reading frame (ORF) with human serum albumin 5' and 3'-UTRs. Full-length 16S mtrRNA sense or antisense RNAs were transcribed in vitro using T7 promoter with the same kit without Capping analogous. Transcribed RNAs were purified using RNase Mini Kit (Qiagen), re-suspended in DEPC-H2O and stored at −80 °C. Ten fragments (100–200 nt in length) of anti-sense RNAs, spanning the entire 16S mtrRNA sequence were prepared and mixed together for microinjection. Single blastomere of a 2-cell embryo was microinjected with either Egfp mRNA (0.8 μg/μl) alone, or together with 16S mtrRNA sense (1.5 μg/μl, 2:1) or anti-sense RNAs (1.5 μg/μl, 2:1) respectively.

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To confirm the expression of mtrRNAs in pre-implantation embryos, ISH of both small (12S) and...
16S mRNA regulates cell lineage allocations

A  
MII oocyte  Zygote  Early 2-cell  Mid 2-cell  Late 2-cell

16S mRNA

3-cell  4-cell  8-cell  Morula  Blastocyst

B  
MII oocyte  Zygote  Early 2-cell  Mid 2-cell  Late 2-cell

16S mRNA

3-cell  4-cell  8-cell  Morula  Blastocyst

C  
MII oocyte  Zygote  Early 2-cell  Mid 2-cell  Late 2-cell

Actin

3-cell  4-cell  8-cell  Morula  Blastocyst

D  

E  
Zygote  2-cell  4-cell

12S mRNA/16S mRNA
large (16S) mtrRNAs was applied using long cRNA probes containing DIG labeled uridine, coupled with anti-DIG-AP conjugates. Strong signals of both 12S and 16S mtrRNAs were seen in embryos at all pre-implantation stages (Fig. 1A and B). In consistency with previous report, mtrRNAs were found to concentrate in the area toward the animal pole where the first polar body emits in MII oocytes (Ninomiya & Ichinose 2007), whereas they appeared to distribute in the cytoplasm of zygotes surrounding pro-nuclei. Following compaction after 8-cell stage, higher levels of mtrRNAs were found in inner cells of morula and blastocysts, comparing with the outer cells of embryos. In contrast, Actin control appeared to be evenly distributed in all embryonic cells (Fig. 1C). Using short oligonucleotide probes (20-mers) encompassing the entire lengths of mtrRNAs, FISH was further performed to determine the expression of mtrRNAs in mouse embryos. These Stellaris oligonucleotide probes were tagged with either Quasar 570 or Quasar 670 Dye for 12S and 16S mtrRNAs respectively. Similarly, signals of mtrRNAs were seen in pre-implantation embryos at all stages, including 2-cell and 4-cell embryos (Fig. 1E). These results indicate that mtrRNAs are continuously expressed in mouse embryos across all pre-implantation stages.

**mtrRNAs are differentially distributed between mouse 2-cell blastomeres**

Although no clear asymmetric distribution of mtrRNAs was found in zygotes, one of the 2-cell sister blastomeres in some embryos appeared to contain more mtrRNAs than the other (Fig. 1A and B, late 2-cells). To better quantify and define the differences of mtrRNAs between sister blastomeres, 2-cell embryos were separated into early, middle and late stages and hybridized with DIG-labeled long cRNA probes (Fig. 2A). The optical densities of ISH signals in sister blastomeres were then measured in pair for each 2-cell embryo outlined from light micrographs. Ratios of signal intensity between two sister blastomeres (the higher vs the lower one, \( R = \text{OD1}/\text{OD2}, \text{OD1} > \text{OD2} \)) were calculated as an indicator for differences of RNA levels. Average ratios were obtained from groups of embryos at each 2-cell stage for each RNA species examined (Fig. 1B and C, Supplementary Table S2). It was found that the ratios of Actin mRNA signals were around 1 between sister blastomeres at all three 2-cell stages, suggesting an equal amount of Actin mRNA between sister blastomeres. In contrast, ratios of signals were much higher for both 12S mtrRNA (\( R = 1.4 \pm 0.2, n = 24 \), late stage) and 16S mtrRNA (\( R = 1.7 \pm 0.6, n = 30 \), late stage) between 2-cell sister blastomeres (Fig. 2B and C). The signal differences of 12S and 16S mtrRNAs appeared to be significantly varied when their ratios were compared with those of Actin control at the same stage (\( P < 0.05 \), Mann–Whitney test). Furthermore, variations of mtrRNA levels between sister blastomeres increased with the development of 2-cell embryos, comparing to that of Actin (\( P < 0.05 \), one-way ANOVA, \( \alpha = 0.05 \)) (Supplementary Table S3). The difference of 16S mtrRNA between 2-cell sister blastomeres could reach up to threefold in some of the embryos examined (Supplementary Table S2).

To confirm that the relative quantification of ISH signals indeed reflects different levels of RNA species between 2-cell sister blastomeres, additional control experiments were performed on house-keeping genes (Gapdh and Hprt) and genes that have been shown to highly express at 2-cell stage (Hmgb3, Tal9 and Ier5) (Zeng et al. 2004). Unlike mtrRNAs, ratios of ISH signals for all genes examined were similar to that of Actin (\( R \approx 1.1–1.2, n = 29–52 \)), indicating that their mRNAs are evenly distributed between 2-cell sister blastomeres (Supplementary Figure S1A and B, see section on supplementary data given at the end of this article). This also suggested that relative comparison of ISH signals could reflect the differences of RNA levels in-between cells and the mtrRNAs are indeed distributed unevenly between sister blastomeres of 2-cell embryos. Although the differences of mtrRNAs between 2-cell sister blastomeres can be as high as threefold in some embryos, the average differences of their signals between sister blastomeres are about 10–50% when their ratios of intensities were subtracted with those of Actin controls at the same stage (13 ± 18% for 12S mtrRNA and 58 ± 67% for 16S mtrRNA at late 2-cell stage respectively). These differences (less than twofold) and large variations among samples could be difficult to reveal using PCR method due to the exponential amplification effect (Weaver et al. 2010). Nevertheless, single-cell quantitative RT-PCR of dissected blastomeres suggested similar

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**Figure 1** Expression of mitochondrial ribosomal RNAs in mouse pre-implantation embryos. (A) In situ hybridization of 12S mtrRNA in mouse embryos at different stages. (B) In situ hybridization of 16S mtrRNA in mouse embryos at different stages. (C) In situ hybridization of Actin in mouse embryos at different stages. MII oocytes were also included. Stars indicate the position of the first polar body. Note the difference in signal intensities between late 2-cell blastomeres in (A) and (B). (D) Quantitative RT-PCR of 16S mtrRNA in pre-implantation embryos. The expression of 16S mtrRNA remained at constant level before 2-cell stage and greatly increased afterwards (black dashed line, three independent experiments). The expression of Actin also increased after 2-cell stage (red dashed line). Same amount of Egfp mRNA was added to each sample and used as internal control (see Materials and methods). PCR without RT was included as negative control. 50 embryos at each stage were used for cell lysate preparation and RT reaction. (E) Fluorescent in situ hybridization (FISH) of mouse pre-implantation embryos. Embryos were hybridized with Stellaris oligonucleotide probes tagged with Quasar 570 (for 12S mtrRNA) and Quasar 670 dye (for 16S mtrRNA) simultaneously. 16S mtrRNA signals were pseudo-colored in green. Shown are merged confocal images.
high variations and differences of mtrRNAs between sister blastomeres (Supplementary Figure S1G). We further quantified mtrRNA levels more directly using the fluorescent signals following FISH of mtrRNAs. Both 12S and 16S mtrRNAs could be readily seen in the cytoplasm of blastomeres when 2-cells were hybridized with short fluorescent probes (Fig. 2D and E). Quantification of the fluorescent signals in paired sister blastomeres (Supplementary Figure S1G). We further quantified mtrRNA levels more directly using the fluorescent signals following FISH of mtrRNAs. Both 12S and 16S mtrRNAs could be readily seen in the cytoplasm of blastomeres when 2-cells were hybridized with short fluorescent probes (Fig. 2D and E). Quantification of the fluorescent signals in paired sister blastomeres. (A) In situ hybridization of mouse 2-cell embryos. Mouse 2-cell embryos were separated into early, middle and late stages and hybridized with long cRNA probes for 12S and 16S mtrRNAs respectively. Actin was used as control. No signals above background were detected when embryos were hybridized with 16S mtrRNA sense probes. (B and C) Ratios of in situ hybridization signal intensities for Actin and mtrRNAs between 2-cell sister blastomeres (R = OD1/OD2, OD1 > OD2). Embryonic stages, number of embryos measured and average ratios of signal intensities are shown in tables (upper panels). Bar graphs are displayed in lower panels. Significances of the differences between control and experimental groups were assessed using Mann–Whitney test for two independent samples. (D and E) Fluorescent in situ hybridization (FISH) of 12S and 16S mtrRNAs in mouse 2-cell embryos using Stellaris oligonucleotide probes (20-mers). Probes were tagged with Quasar 570 and Quasar 670 dyes for 12S and 16S mtrRNAs respectively. 16S mtrRNA signal was pseudo-colored as green. (F) Quantification of FISH signals between 2-cell sister blastomeres. Comparing to DAPI staining, both mtrRNAs appeared higher in one blastomere than the other. Number of embryos measured and the average ratios of signal intensity are shown in table (upper panel). Bar graph is shown in lower panel. Significances of the differences between control and experimental groups were assessed using Mann–Whitney test. Error bars: s.d.
blastomeres showed that 2-cell embryos contained differential levels of mtrRNAs in sister blastomeres (about 10–20% at the examined mid-2-cell stage), comparing with the equally distributed nuclear DAPI signals (n=20, 23; P=0.048, 0.0003 for 12S and 16S mtrRNAs respectively, Mann–Whitney test) (Fig. 2F).

It has been suggested that hormone treatment during superovulation or different mouse strains would influence the quality of embryos obtained, therefore casting environmental effects on development (Sanfins et al. 2003). To eliminate the possibility that these artificial treatments will deviate the distribution of mtrRNAs between 2-cell sister blastomeres, mid-2-cell embryos isolated from CD1 mice that were naturally mated or from C57BL/6 mice that were superovulated was hybridized with DIG-labeled long cRNA probes. Comparing with the Actin controls that were processed at the same time, expression of 16S mtrRNA was found to be more differential between 2-cell sister blastomeres from these mice. The average ratios of ISH signal intensity for 16S mtrRNA were significantly more variable (1.4 ± 0.3 for CD1 mice, n=23; 1.4 ± 0.4 for C57BL/6, n=20) comparing with those of Actin (P<0.05, z-test for two samples’ means) (Supplementary Figure S1C, D, E and F). This suggested that the uneven distribution of 16S mtrRNA between 2-cell sister blastomeres was not affected by hormone injection or different strains of mice. Taken together, these results indicated that mtrRNAs are differentially distributed between mouse 2-cell sister blastomeres.

To examine whether the asymmetric distribution of mtrRNAs would persist with further cell divisions, levels of mtrRNAs among blastomeres in 4-cell embryos were examined using ISH. To better visualize 4-cell sister blastomeres, they were first manually separated into single cells from each embryo (Fig. 3A). Groups of four sister blastomeres of the same embryo were then hybridized with the same long cRNA probes together. Signals of mtrRNAs in individual blastomeres were hybridized with DIG-labeled long cRNA probes. Comparing with the sister blastomere with the highest intensity for Actin, signal intensities for Actin and 12S mtrRNA among sister blastomeres were significantly varied (P<0.05, n=22 and 16 respectively), comparing with those of Actin (P=0.438, n=15, one-way ANOVA, α=0.05) (Supplementary Tables S4 and S5). The high variations and differences of mtrRNA levels among 4-cell sister blastomeres were also seen using single cell quantitative RT-PCR (Supplementary Figure S1H). These results suggested that the asymmetric distribution of mtrRNAs could be propagated with further cell divisions. Recent studies showed that mouse 4-cell blastomeres contain differentially expressed epigenetic

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Figure 3 Distribution of mtrRNAs among sister blastomeres of mouse 4-cell embryos. (A) In situ hybridization of mouse 4-cell embryos. Sister blastomeres of 4-cell embryos were separated manually by micromanipulation before being hybridized together with indicated probes. Shown are representative light micrographs. Images of in situ hybridization with 16S mtrRNA sense probe from a separate experiment was included as control. (B) Ratios of in situ hybridization signal intensities for Actin and 12S mtrRNA among 4-cell sister blastomeres. Comparing to the sister blastomere with the highest signal intensity (set as 1), ratios of Actin and 12S mtrRNA signals for the two lowest blastomeres are 0.93 ± 0.03 (n=16) and 0.82 ± 0.11, 0.73 ± 0.1 (n=22), respectively. (C) Ratios of in situ hybridization signal intensities for Actin and 16S mtrRNA among 4-cell sister blastomeres. Comparing with the sister blastomere with the highest signal intensity (set as 1), ratios of Actin and 16S mtrRNA signals for the two lowest blastomeres are 0.95 ± 0.03, 0.92 ± 0.04 (n=15) and 0.83 ± 0.07, 0.75 ± 0.07 (n=16) respectively. Significances of the differences between control and experimental groups were assessed using Mann–Whitney test (see also Supplementary Tables S4 and S5). Error bars: s.d.
modifiers that influence the destination of respective progeny cells in blastocyst (Torres-Padilla et al. 2007, Burton et al. 2013). Although it is not clear at this stage whether the differentially distributed mtrRNAs in mouse 2-cell embryos are related to the differential gene expression at later stages, the results nevertheless suggest that molecular asymmetry occurs in mouse 2-cell embryos, earlier than previously found embryonic stage.

**Mouse 2-cell embryos contain cytoplasmic 16S mtrRNA outside of mitochondria**

Since mtrRNAs are essential components of mitochondrial ribosomes, whether the differential localization of mtrRNAs reflects differences in mitochondria distribution or mitochondrial activity between 2-cell sister blastomeres was examined next. Whole-mount immunofluorescent staining of embryos with Mito-Tracker Red showed overall even distribution of mitochondria in the cytoplasm of 2-cell blastomeres. Quantification of MitoTracker Red signals in paired sister blastomeres of 2-cell embryos from confocal micrographs gave rise to the ratio of 1.1 ± 0.08 between sister blastomeres, similar to that of nuclear DAPI staining (R = 1.1 ± 0.07, n = 30), indicating that 2-cell sister blastomeres contain similar amounts of mitochondria (Fig. 4A and B). Interestingly, in contrast to apparent higher levels of mtrRNAs in inner cells of morula and blastocyst (Fig. 1A and B), fluorescent signals of MitoTracker Red in the inner cells appeared weaker than those of outer cells (Supplementary Figure S2A, see section on supplementary data given at the end of this article), suggesting a probable higher mitochondria contents in cells at the outer layer of morula and blastocysts.

Next, the membrane potential of mitochondria was examined using TMRM staining of live 2-cell embryos. Comparing to embryos treated with FCCP (the membrane potential un-coupler), live embryos were readily stained by the dye (Fig. 4C). Little difference in the intensity of TMRM staining was found between 2-cell sister blastomeres (R = 1.1 ± 0.08, n = 40), similar to that of nuclear DNA staining (R = 1.08 ± 0.12, Fig. 4D), suggesting that 2-cell blastomeres also have similar mitochondrial activity. Thus, neither the quantity nor the activity of mitochondria is differentially distributed between sister blastomeres of 2-cell embryos.

One possibility for un-evenly distributed mtrRNAs seen in 2-cell embryos is the incomplete hybridization of long anti-sense cRNA probes due to their difficulty to penetrate mitochondrial membranes, as previously suggested (Ninomiya & Ichinose 2007). To test this, the expression of two mitochondria encoded genes (Nd2 and Cytb) and a nuclear gene (Cyc1) was analyzed using respective long cRNA probes. Quantitative RT-PCR first showed that both Nd2 and Cytb were expressed at levels higher than that of 16S mtrRNA at most pre-implantation.

![Figure 4](https://www.reproduction-online.org/151/351-367/Reproduction_2016_151_351-367)

**Figure 4** Mouse 2-cell embryos contain overall equal amount and activity of mitochondria. (A) MitoTracker Red staining of mouse 2-cell embryos. Nuclei were stained with DAPI. Shown are representative confocal images. (B) Ratio of signal intensities of MitoTracker Red between 2-cell sister blastomeres. Similar ratios of mitochondria (R = 1.09 ± 0.09) and DAPI (R = 1.09 ± 0.07, n = 30) indicate equal amounts of mitochondria between sister blastomeres. (C) TMRM staining of live 2-cell embryos. As control, FCCP was used to un-couple mitochondrial membrane potential before TMRM staining (lower panels). Nuclei were stained with Hoechst 33342. Shown are representative confocal images. (D) Ratio of TMRM signal intensities between 2-cell blastomeres. Similar ratios of TMRM (R = 1.12 ± 0.08) and Hoechst (R = 1.08 ± 0.12, n = 40) indicate the equal mitochondrial activity between sister blastomeres.
embryonic stages, whereas Cyc1 was expressed at much lower level (Fig. 5A). However, both mRNAs of Nd2 and Cytb that are known to be inside of mitochondria were not detected by ISH, whereas Cyc1 mRNA was readily seen in morula and blastocysts using the same method (Fig. 5B). This suggested that long cRNA probes was able to detect relatively small amount of mRNAs in the cytoplasm of cells during ISH, but may indeed be difficult to localize RNA species within mitochondria. It also suggested that the differentially distributed mtrRNAs detected by long cRNA probes were probably localized outside of the organelle. Since small

**Figure 5** Distribution of cytoplasmic 16S mtrRNA outside of mitochondria in mouse 2-cell embryos. (A) Quantitative RT-PCR of mitochondrial genes. Shown are average expression levels relative to that of 16S mtrRNA from three independent experiments. (B) In situ hybridization of pre-implantation mouse embryos. Collected embryos (from 1-cell to blastocyst) were hybridized with long cRNA probes for indicated genes. Signals for mitochondria encoded Cytb and Nd2 were not detectable in cytoplasm despite their higher levels of expression (A), whereas nuclear encoded Cyc1 could be seen at morula and blastocyst stages. Anti-sense and sense probes for mtrRNAs were used as controls. (C) Co-staining of 16S mtrRNA and mitochondria. Signals of 16S mtrRNA (detected by fluorescent oligonucleotide probes) were found to partially co-localize with mitochondria (detected by MitoTracker Red) from MII oocytes to blastocysts. Inner cells of morula and blastocysts contain higher levels of 16S mtrRNA than MitoTracker Red signals. (D) In situ hybridization of 16S mtrRNA at electron microscopic level (ISH-EM). 16S mtrRNA (arrow heads) was detected in the cytoplasm outside of mitochondria (arrows), as revealed by anti-DIG-colloidal gold secondary antibodies. Sense probe was used as negative control (right panel). Average numbers of gold particles per frame were calculated from ten frames of electron micrographs in each group (Inset, n=10). Significance of the difference was assessed using Mann–Whitney test. Error bars: s.d.
oligonucleotide probes could better penetrate mitochondria, FISH signals of mtrRNAs may represent both cytoplasmic and mitochondrial fractions of mtrRNAs. Using fluorescently tagged oligonucleotide FISH probes, together with MitoTracker Red, the subcellular localization of 16S mtrRNA in mouse embryos was further examined. In control experiments, fluorescent signals of 16S mtrRNA were found to completely overlap with that of MitoTracker Red in mouse embryonic fibroblast (MEF) and 293T cells (Supplementary Figure S2B). However, in mouse MII oocytes and pre-implantation embryos, signals of 16S mtrRNA were found to only partially overlap with MitoTracker Red staining, suggesting that 16S mtrRNA was also present in the cytoplasm outside of mitochondria in mouse female germ cells and embryos (Fig. 5C). Green fluorescent signals of 16S mtrRNA that did not overlap with MitoTracker Red were found in the cytoplasm of MII oocytes toward animal pole and more concentrated in inner cells in morula and blastocysts, suggesting that 16S mtrRNA at these regions was likely outside of mitochondria, consistent with ISH results using long cRNA probes (Fig. 1).

Mitochondrial rRNAs have been found outside of mitochondria in early embryos of Drosophila, Sea urchin and Xenopus (Kobayashi et al. 1993, 1998, Ogawa et al. 1999). In mouse, it has been suggested that MII oocytes also contain mtrRNAs in the cytoplasm (Ninomiya & Ichinose 2007). To further determine the sub-cellular localizations of 16S mtrRNA in relation to mitochondria, 2-cell embryos were examined at electron microscopic level following ISH. In control experiment, transmission electron microscopy of thin sections (100 nm) of 2-cell embryos showed that mitochondria appeared to be spherical in shape with clear cristae (Supplementary Figure S2C). Following ISH, mitochondria were less well maintained and the internal cristae of mitochondria were sometimes lost due to repeated sample treatments. However, the overall morphology of mitochondria remained identifiable (Supplementary Figure S2D). Using anti-DIG antibody conjugated with 12-nm colloidal gold particles, it was found that 16S mtrRNA was distributed in the cytoplasm of 2-cell blastomeres outside of mitochondria (Fig. 5D). The average number of gold particles on thin sections of 2-cell embryos was significantly higher when anti-sense probes of 16S mtrRNA were used (16.7 ± 7.8), comparing with that of sense control (5.3 ± 2.7; P = 0.002, n = 10, Mann–Whitney test) (Fig. 5D, inset). Collectively, these results indicate that mouse 2-cell blastomeres contain 16S mtrRNA in the cytoplasm outside of mitochondria.

16S mtrRNA in 2-cell embryos regulates cell allocations in blastocyst

First found in Drosophila, 16S mtrRNA was shown to drive the germ cell formation during early embryogenesis (Iida & Kobayashi 1998). To find out whether the differentially expressed 16S mtrRNA in mouse 2-cell embryos could have functional roles in cell-lineage allocations during pre-implantation embryogenesis, levels of 16S mtrRNA were altered in one sister blastomere of 2-cell embryos using microinjection of either sense or antisense RNAs. In vitro transcribed Egfp mRNA was used as control or co-injected into the same blastomere in order to trace the fate of its descendant into blastocysts (Fig. 6A). Injected embryos were developed till blastocyst stage in vitro and scanned at 2-μm interval using confocal microscopy. Confocal sections of each individual blastocyst were compiled into single 3-D images, from which the number and locations of cells were analyzed (Supplementary Movies 1 and 2, see section on supplementary data given at the end of this article). In the three dimensional space, cells that are located on the outer-most layer on either axis of the three dimensions (X, Y and Z) were considered as the TE cells, whereas cells that are surrounded by others on all three axes were counted as the ICM cells (Fig. 6B). Cell positions were determined by the nuclear DAPI staining and green fluorescence was used to identify the EGFP-labeled (EGFP⁺) cells. For the ease of analyses of 3D cell allocations, embryos containing 40–45 cells (46–48 h in vitro culture following microinjection), corresponding to early blastocysts were used in most cases. At this stage, ICM and TE cells are segregated, whereas epiblast and primitive endoderm cells of the ICM are yet to be differentiated. To confirm the cell positioning method, embryos were co-stained with cell–cell junctional E-Cadherin or plasma membrane marker TRITC-Phalloidin in some cases so that cell boundaries were clearly defined. It was found that cells in both ICM and TE could be similarly positioned with or without cell membrane labeling (Supplementary Figure S3). Thus for most of the cell allocation analyses, embryos without cell membrane labeling were used. Total numbers of EGFP⁺ cells that were allocated into either ICM or TE were then counted in order to assess the effects of changing 16S mtrRNA levels on cell lineage allocations.

Quantitative RT-PCR showed that anti-sense mtrRNA decreased 16S mtrRNA level in injected blastomeres, while sense RNA increased it by about 60% (Supplementary Figure S4A, see section on supplementary data given at the end of this article). The decrease of 16S mtrRNA following anti-sense microinjection was also supported by the decreased fluorescent signals in injected blastomeres when compared to the non-injected ones using FISH (Supplementary Figure S4B). Examination of Egfp mRNA-injected embryos first showed that EGFP⁺ cells derived from microinjected blastomere occupied blastocyst in both ICM and TE without affecting the overall development of embryos in vitro. All groups of blastocysts contained similar numbers of cells when cultured for the same length of time in vitro (Supplementary Table S6). There were about 34% of EGFP⁺ cells in ICM and 66% of EGFP⁺ cells in TE in Egfp-injected embryos (Fig. 6F). Comparing
with the Egfp-injected control, increasing 16S mtrRNA (sense injection) increased the number of EGFP⁺ descendent in ICM (to about 49%) \( (P=0.005, n=18) \), while reduced 16S mtrRNA level (anti-sense injection) caused an increase in the number of EGFP⁺ cells in TE (to about 78%) \( (P=0.0001, n=19, \text{Mann–Whitney test}) \) (Fig. 6F and Supplementary Table S6). These changes of cell allocations of EGFP⁺ cells following microinjection appeared statistically significant when they were compared together across different conditions \( (P < 0.001, \text{one-way ANOVA}, \alpha = 0.05) \) (Supplementary Table S7).

The cell allocation effects of altered 16S mtrRNA levels were further examined by cell lineage specific gene expression. Using antibodies against cell-lineage markers NANOG (pluripotency marker) and CDX2 (TE marker), in vitro cultured blastocysts were immunostained following microinjection. It was found that patterns of marker gene expression in either ICM or TE cell lineages were not changed with respective allocations in blastocyst following either sense or anti-sense injection, comparing to the non-injected control group (Fig. 6C, D and E). NANOG was primarily found in ICM cells with occasional staining in TE cells, whereas CDX2 was exclusively localized in TE cells at the outer layer in all groups. Since NANOG is a pluripotency marker but not a strict ICM specific marker at the early blastocyst stage and CDX2 is an exclusive TE lineage marker, changes of CDX2 expressing (CDX2⁺) cells in microinjected embryos were further analyzed. It was found that, while overall development of the embryos remained unaltered (embryos contained similar number of cells during similar period of culturing time in all groups), EGFP⁺ cells contained higher fraction of CDX2⁺ cells \( (16.7 ± 5.0, 69.5 ± 13.3\% \text{ of total EGFP⁺ cells}) \) when 16S anti-sense was injected, comparing with sense injected embryos \( (9.4 ± 4.2, 42.6 ± 16\% \text{ of total EGFP⁺ cells}) \), \( P=0.0002, n=18–19, \text{Mann–Whitney test} \) (Fig. 6G, Supplementary Tables S8 and S9). These results suggested that changes of 16S mtrRNA did not change the marker gene expression prior to cell lineage allocations and levels of 16S mtrRNA in 2-cell blastomereres could deviate the segregation of ICM and TE cell lineages in blastocyst. Neither anti-sense nor sense 16S mtrRNA brought changes to mitochondrial gene transcription.

**Figure 6** Influences of 16S mtrRNA in 2-cell blastomereres on cell allocations in mouse blastocyst. (A) Schematic drawing of 2-cell embryo microinjection. In vitro transcribed Egfp mRNA was injected into one blastomere at random, either singly or together with 16S mRNA. Embryos were cultured in vitro for 46–48 h till blastocyst stage and EGFP⁺ descendent cells were traced for their locations in either inner cell mass (ICM) or trophoblast stem (TE) within blastocyst. (B) Representative confocal images showing locations of EGFP⁺ ICM and TE cells. Blastocysts cultured in vitro following microinjection were scanned under a confocal microscope at 2-µm interval. EGFP⁺ descendent cells were traced for their locations as marked by DAPI staining using 3-D compositional images. Cells that are surrounded by other cells along three axes \( (X, Y \text{ and } Z) \) are denoted as ICM cells (left panel) and cells that are located on the outside along either one of the three axes are denoted as TE cells (right panel). Shown are center sections of a scanned blastocyst with indicated ICM and TE cells on three dimensions. (C, D and E) Immunostaining of NANOG and CDX2 in mouse blastocysts with or without microinjection. Embryos were cultured in vitro from 2-cell stage, when one blastomere was microinjected with indicated RNA species. Similar to the control (C), NANOG was found to primarily express in ICM cells, with occasional staining in TE cells (D), whereas CDX2 was found to express exclusively in TE cells in all groups (E). Cell nuclei were stained with DAPI. Representative confocal images of immunostaining of NANOG or CDX2 are shown on the left and merged images with either DAPI or EGFP signals are shown on the right in each panel. (F) 16S mtrRNA influences cell allocations in blastocyst. Comparing to Egfp mRNA injected controls, 16S mtrRNA sense or anti-sense increased EGFP⁺ cells’ occupation in either ICM or TE, respectively \( (n=18–19; \text{see also Supplementary Tables S6 and S7}) \). (G) While total number of cells were not changed in blastocysts with or without microinjection, 16S mtrRNA anti-sense increased CDX2⁺ cells in EGFP⁺ fraction \( (n=18–20; \text{see also Supplementary Tables S8}) \). Pair-wise comparison was assessed for significance using Mann–Whitney test, while significance of differences among three groups were assessed using one-way ANOVA and post-hoc Turkey analyses (see also Supplementary Table S9).
activity between 2-cell blastomeres (Supplementary Figure S4C and D), suggesting that the extra-mitochondrial fraction of the mtrRNA may be accountable for the cell lineage allocation effects.

Since bi-laterally symmetric mouse blastocysts can be divided into embryonic half (containing ICM and polar TE) and abembryonic half (containing mural TE) along the A–V axis, cell allocation effects of 16S mtrRNA along the A–V axis following microinjection was further analyzed. Based on the distribution of EGFP + descendant in either embryonic or abembryonic allocations in blastocysts, embryos were separated into three different categories (Fig. 7A and B). Among 101 Egfp-injected embryos, 31.7% of embryos contained EGFP + progenies only at embryonic side (pattern 1, embryonic) and 25.7% of embryos contained EGFP + cells only at abembryonic side (pattern 3, abembryonic). The remaining 42.6% of embryos contained EGFP + cells in both cell allocations (pattern 2, mixed). However, when single 2-cell sister blastomeres were co-injected with anti-sense 16S mtrRNA and Egfp mRNA, blastocysts containing EGFP + progenies in embryonic half reduced from 31.7% to 15.3%, whereas blastocysts containing EGFP + cells in the mixed cell allocations increased to 68.1% (n = 144). Including the remaining 16.7% embryos with EGFP + cells located only at abembryonic half, a total of 84.8% embryos contained progenies from injected blastomeres at abembryonic allocations (comparing to the total of 68.3% in the control group). The dramatic increase of embryos with mixed cell allocations is probably caused by the randomness at choosing the 2-cell sister blastomeres for microinjection. In contrast, when in vitro transcribed 16S mtrRNA was co-injected with Egfp mRNA into single 2-cell blastomeres, blastocysts containing EGFP + descendant allocated to embryonic cell-lineage increased to 46.5%, whereas blastocysts with EGFP + abembryonic cells decreased to 12.6% (n = 127) (Fig. 7C). Statistical analysis suggested that these changes of embryos in different patterns were significantly relevant to the microinjection of either sense or anti-sense 16S mtrRNAs among experimental groups (χ² = 39.18, P < 0.001, χ² test, Supplementary Table S10). Taken together, these results suggested that the increase of 16S mtrRNA in 2-cell blastomeres facilitates their progeny cells to allocate into pluripotent embryonic cell lineage, whereas reduction of 16S mtrRNA deviates the descendant into abembryonic allocation.

Discussion

It has been reported that the total number of mitochondria and the mitochondrial DNA remain largely unchanged in pre-implantation mouse embryos (Piko & Taylor 1987). Since embryonic mitochondria are mostly inherited from the maternal source (Cree et al. 2008, Mishra & Chan 2014), the number of mitochondria in embryonic cells gradually decreases with cell division until post-implantation stage when cell proliferation accelerates. However, mtrRNAs are actively transcribed during pre-implantation development (Piko & Chase 1973, Piko & Taylor 1987). Consistent with this, quantitative RT-PCR showed that the level of 16S
mtrRNA is maintained constant from MII oocytes to 2-cell embryos and increased dramatically after 4-cell stage. This suggests that mitochondrial genome is probably expressed continuously in the absence of DNA replication, separating its gene expression regulation away from that of nuclear genome (Lee et al. 2014).

In unfertilized eggs, mtrRNAs appeared to localize more towards the animal pole where the first polar body emits. Uneven distribution of mtrRNAs was also observed throughout early embryos at different stages from 2-cell and 4-cell (higher in half of the blastomeres) to morula and blastocyst (higher in inner cells). It is not clear at this stage whether the maternal polarity is related to the molecular asymmetry observed in early embryos. Several studies have shown that LEPTIN and STAT3 are asymmetrically localized in unfertilized eggs and early embryos in mouse, accumulating at the animal pole toward the first polar body (Antczak & Van Blerkom 1997, Schulz & Roberts 2011). This is similar to that of mtrRNAs in unfertilized eggs (Fig. 1A and B). A large multi-protein complex (subcortical maternal complex (SCMC)) was also found to localize in the subcortical regions of mouse eggs and early embryos. Genetic depletion of the SCMC components in mouse caused developmental retardation at 2-cell stage (Li et al. 2008). During compaction, these polarized factors accumulate differentially in outer cells vs inner cells, generating polarity gradient within the embryo that may be linked to the cell-lineage determination and axis formation in blastocyst. However, none of them was found to differentiate in-between sister blastomeres following fertilization when embryogenesis starts. Recent research suggested that differentially localized epigenetic factors appearing at very early stage (4-cell stage) might regulate cell lineage determination in blastocysts (Torres-Padilla et al. 2007, Plachta et al. 2011, Burton et al. 2013). However, how these early molecular asymmetries are generated and whether there are even earlier cell fate determinants in mouse pre-implantation embryos remain elusive. Results presented here indicate that mouse 2-cell sister blastomeres contain quantitatively different mtrRNAs, suggesting that the molecular asymmetry may occur at the very beginning of mouse embryogenesis. Whether the differential mtrRNAs in 2-cell blastomeres are functionally related to the molecular asymmetries that occur during the development of early embryos requires further exploration.

Although the origin of differentially distributed mtrRNAs remains to be determined, their differences between sister blastomeres appear to be more evident at late 2-cell stage. One possibility is that the partitioned maternal mtrRNAs are differentially retained or degraded with the development of embryos. The transcription activity of mitochondrial genome could also differ between sister blastomeres. Alternatively, since the differential distribution of mtrRNAs were detected mainly outside of the organelle, mitochondria in sister blastomeres could export differential amount of mtrRNAs into the cytoplasm. It will be interesting to find out whether the post-transcriptional modifications of RNAs and mitochondrial activities contribute to the differential distribution of mitochondrial small and large ribosomal RNAs between 2-cell sister blastomeres. Although it is known that embryonic mitochondria are mainly inherited from the maternal source, it remains possible that sperm entry or the paternal cellular components could contribute to the regulation of embryogenesis beyond fertilization and activation of zygote. Previous sperm labeling and tracing experiment in mouse showed that sperm tail enters the egg during fertilization and remains in one of the 2-cell blastomeres (Jefferson & Williams 2012). The experiment showed that sperm tail remained largely intact at 2-cell stage. Given that sperm mid-piece contains mitochondrial sheath with rather compacted configuration, this is different from what’s seen with unevenly distributed mtrRNAs, which have diffused localization patterns. Research in Drosophila showed that sperm mitochondria are destroyed through autophagic and endocytic effects following fertilization (Politi et al. 2014). Whether paternal mitochondria are also eliminated in the same manner in mouse remains to be determined. Previous studies suggested that the mode of the first cell divisions is relevant to the progenies’ cell lineage allocations in blastocysts (Piotrowska et al. 2001, Piotrowska-Nitsche & Zernicka-Goetz 2005). Sequential meridional–equatorial divisions of 2-cell blastomeres showed more predictable fate of the descendent in the blastocyst (the blastomere that divide meridionally first tend to allocate into the embryonic hemisphere) than 2-cell blastomeres that are divided in the same directions (meridional–meridional or equatorial–equatorial divisions). Perhaps the non-uniform dividing orientations could help to generate molecular gradients among progenies which are necessary preludes for cells’ further differentiation. In this regard, it will be interesting to find out whether the levels of mtrRNAs are relevant to the regulation of the first cell divisions and cell lineage allocations in early embryos.

How are mitochondria encoded mtrRNAs localized in the cytoplasm outside of the organelle? Although mtrRNAs outside of mitochondria have been found in various species, including Drosophila, Sea urchin, Xenopus and mouse (Kobayashi et al. 1993, 1998, Ogawa et al. 1999, Ninomiya & Ichinose 2007), it is not clear what molecular mechanisms that govern their cytoplasmic distribution. It was suggested that various RNA species encoded by mitochondrial genome could be exported into cytoplasm of human cells via unknown mechanisms (Attardi & Attardi 1968, Maniataki & Mourelatos 2005). One intriguing phenomenon is that neither the number nor the membrane potential of mitochondria has apparent differences between 2-cell
sister blastomeres. It is known that mouse pre-implantation embryos utilize pyruvate and lactate instead of glucose to meet their energy and metabolic requirements and maintain relatively dormant mitochondrial oxidative phosphorylation activity until blastocyst stage (Krisher & Prather 2012). Interestingly, inner cells of morula and blastocyst appear to contain higher levels of mtrRNAs when long cRNA probes were used for hybridization, whereas outer cells showed higher contents of mitochondria as indicated by TMRM staining (Fig. 1Aa and B, Supplementary Figure S2A). The possible higher levels of cytoplasmic mtrRNAs in inner cells of late pre-implantation embryos are in parallel with the cell lineage allocation effects of 16S mtrRNA. It is possible that the expression and distribution of mtrRNAs is related to unknown mitochondrial activities in early embryos. MtrRNAs are important components of mitochondrial ribosomes for mitochondrial protein synthesis. Despite the dormant oxidative phosphorylation activity of mitochondria, mitochondrial protein synthesis remains active in early embryos (Piko & Chase 1973, Cascio & Wassarman 1981). One possible functional consequence of the cytoplasmic localization of mtrRNAs can be the participation of mitochondrial ribosomes, which are more close to prokaryotic ribosomes than to its eukaryotic cytoplasmic counterparts (Amunts et al. 2014), in the synthesis of specific subsets of mRNAs that may play crucial roles during pre-implantation embryogenesis (Amikura et al. 2001). Intriguingly, latest studies showed that mitochondrial genome contains short ORFs (sORFs) that reside within mtrRNAs genetic loci, which encode short polypeptides that are translated following the export of mtrRNAs (Hashimoto et al. 2001, Lee et al. 2015). These polypeptides (the 24-aa long Humanin from 16S mtrRNA and the 16-aa long MOTS-c from 125 mtrRNA) are biologically functional and elicit critical roles in regulating insulin sensitivity and maintaining metabolic homeostasis in neuronal and skeletal muscle cells (Hashimoto et al. 2001, Muzumdar et al. 2009, Lee et al. 2015). It will be interesting to fully elucidate whether mtrRNAs regulate cell lineage allocations during early embryogenesis via RNA-mediated or alternative mechanisms and how mitochondrial genome participates in the process.

The mechanisms of cell fate determination regulated by early embryonic events during embryogenesis have been under intensive investigation (Rossant & Tam 2009). It is known that blastomeres in early embryos possess developmental plasticity during experimental manipulations. Although molecular alterations within isolated embryonic cells have not been examined, the adaptive nature of embryos suggests that mammalian pre-implantation embryogenesis is a subject of more complicated regulation than previously anticipated (Zernicka-Goetz et al. 2009). During microinjection experiments, altered 16S mtrRNA levels influenced cell lineage allocations of descendant in the blastocyst. Although it is not clear whether effects brought by alterations of mtrRNAs will elicit in a stoichiometric or dose-dependent manner, statistical analyses showed that altered 16S mtrRNA levels brought significant changes to cell lineage allocations of descendant. Specific cell lineage markers (such as NANOG and CDX2), however, maintained constant within respective positions of embryonic cells, even when progeny cells of injected blastomeres were deviated to either ICM or TE. This suggests that 16S mtrRNA may participate in the regulation of cell allocations prior to cell lineage specific gene expression (Nance 2014, Rayon et al. 2014). Recent genome-wide gene expression analyses on early embryonic cells in both mouse and C. elegans indicated a bi-facet model of cell lineage determination, in which an earlier stochastic heterogeneous gene-expression stage when cells appear non-distinguishable is followed by a late reinforced homogeneous gene-expression stage when cells gain respective molecular signatures (Ohnishi et al. 2014, Du et al. 2015). These suggested that perhaps earlier events preluding more definitive cell-lineage determination pathways are regulated by post-transcriptional mechanisms, involving various RNA species and RNA binding proteins. The correlation between mtrRNAs and cell lineage allocations in the blastocyst indicates that differentially expressed molecules, such as mtrRNAs, are functionally involved in the regulation of cell fate determination at the very beginning of animal development. Future research is required to elucidate the molecular mechanisms that govern the partitioning of mtrRNAs and how they regulate the cell lineage allocation during mouse pre-implantation embryogenesis.

Supplementary data
This is linked to the online version of the paper at http://dx.doi.org/10.1530/REP-15-0301.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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