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Restricted development of mouse triploid fetuses with disorganized expression of imprinted genes

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Running Head: Imprinted genes in mouse triploid fetuses

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Abstract

Eukaryotic species commonly contain a diploid complement of chromosomes. The diploid state appears to be advantageous for mammals because it enables sexual reproduction and facilitates genetic recombination. Nonetheless, the effects of DNA ploidy on mammalian ontogeny have yet to be understood. The present study shows phenotypic features and expression patterns of imprinted genes in tripronucleate diandric and digynic triploid (DAT and DGT) mouse fetuses on embryonic day 10.5 (E10.5). Measurement of crown-rump length revealed that the length of DGT fetuses (1.87 ± 0.13 mm; mean $\pm$ SEM) was much smaller than that of diploid fetuses (4.81 ± 0.05 mm). However, no significant difference was observed in the crown-rump length between diploid and DAT fetuses (3.86 ± 0.43 mm). In DGT fetuses, the expression level of paternally expressed genes, *Igf2*, *Dlk1*, *Ndn*, and *Peg3*, remained significantly reduced and that of maternally expressed genes, *Igf2r* and *Grb10*, increased. Additionally, in DAT fetuses, the *Igf2* mRNA expression level was approximately twice that in diploid fetuses, as expected. These results provide the first demonstration that imprinted genes in mouse triploid fetuses show distinctive expression patterns independent of the number of parental-origin haploid sets. These data suggest that both DNA ploidy and asymmetrical functions of parental genomes separately influence mammalian ontogeny.

Key words: triploid, genomic imprinting, fetal growth, gene expression

Introduction

Polyploidy occurs relatively frequently among some animal groups, in particular, in fish and amphibians (McKinnell, 1964; Cuellar and Uyeno, 1972). In mammals, triploids and tetraploids show severe developmental anomalies, resulting in lethality at midgestation (Triploid: embryonic day 11.5 (E11.5), Tetraploid: E14.5) (Kaufman *et al.*, 1989a; Kaufman and Webb, 1990; Henery and Kaufman, 1992). As an exception, some human triploids could develop until the later stages of gestation, 46 to 308 days, and rarely survive more than one month after birth, exhibiting neonatal mortality (Arvidsson *et al.*, 1986; Fryns *et al.*, 1977; Hasegawa *et al.*, 1999; Iliopoulos *et al.*, 2005; Niemann-Seyde *et al.*, 1993; Sherard *et al.*, 1986). Triploidy has been estimated to occur in 1–3% of pregnancy cases in humans, which is one of primary causes of not only early miscarriage but also aberrations in a fetus or infant (Beatty, 1978; Carr, 1971a; Carr, 1971b; Niebuhr, 1974). The possible causes include dispermy or inhibited extrusion of the second polar body during fertilization (Jacobs *et al.*, 1978). Hence, triploid embryos could be classified into two types according to whether the extra haploid set of chromosomes (in triploidy) was derived from a sperm or oocyte, namely diandric triploid (DAT) and digynic triploid (DGT) embryos, in which the extra haploid set is either paternal or maternal, respectively. Nonetheless, it is unknown how triploidy in mammals leads to lethality, which is not the case in other animal groups such as fish and amphibians.

Because the triploid human conceptuses at pre- and postimplantation stages are unavailable for

experimental analysis, researchers have to use experimentally constructed mammalian triploid embryos that are not human. In mice, both DAT and DGT embryos show distinct phenotypes, particularly on E9.5 (Kaufman *et al.*, 1989a). The DGT fetuses are significantly smaller than diploid fetuses with many morphological aberrations such as cardiac defects, craniofacial anomalies, and neural tube defects, although the frequency of these problems has not been determined after E9.5 precisely (Kaufman *et al.*, 1989a; Kaufman and Speirs, 1987). In contrast, all of the DAT fetuses are relatively normal in terms of morphological characteristics. DAT fetuses had about 20-25 pairs of somites, of which the cephalic and cardiac regions were developmentally advanced as well as those of diploid fetuses at E9.5 (Henery and Kaufman, 1992; Kaufman *et al.*, 1989a). In both types of triploid embryos, the composition of sex chromosomes; i.e., XXX and XXY, appears not to cause significant phenotypical changes including fetal development, although DAT fetuses with the XXY set have not been observed (Kaufman *et al.*, 1989b; Speirs and Kaufman, 1989). The above-mentioned contrasting phenotypes of DAT and DGT embryos are evidence that the male and female genomes are not functionally equivalent, which is also the case for uniparental gynogenetic (parthenogenetic) and androgenetic diploids (McGrath and Solter, 1984; Surani *et al.*, 1984; Surani *et al.*, 1986). The functional and transcriptional asymmetry between parental genomes is explainable in terms of genomic imprinting. The latter is a mechanism for making one parental allele different from the other, and this process leads to parental origin-dependent transcription in mammals.

Imprinted genes, where only one of the two parental chromosome copies is expressed, are the result of epigenetic modifications, such as DNA methylation (Li *et al.*, 1993). Acquisition of methylation of key controlling elements, i.e., imprinting of control regions (ICRs), occurs in parental germ lines during male or female gametogenesis and these acquired modifications (methylation) are usually retained thereafter (Obata and Kono, 2002). A number of imprinted genes are clustered with *cis*-acting ICRs carrying allele-specific methylation marks (Birger *et al.*, 1999; Kim *et al.*, 2012; Lin *et al.*, 2003; Shiura *et al.*, 2009; Thorvaldsen *et al.*, 1998; Verona *et al.*, 2008). Because the dosage of imprinted genes is crucial for normal development, growth, and behavior in mammals, the loss or overexpression of those transcripts can cause dysontogenesis in some cases, resulting in death of uniparental mouse embryos (Ogawa *et al.*, 2006; Sotomaru *et al.*, 2002). In contrast, nothing is currently known about transcriptional dynamics of imprinted genes in triploid mouse embryos.

Uniparental diploid embryos in mice are thought to be a good experimental model for elucidation of the roles of parental genomes during early embryogenesis (Surani *et al.*, 1986). Recent studies demonstrated that the parent-specific expression of imprinted genes is not always maintained in uniparental fetuses (Ogawa *et al.*, 2006; Sotomaru *et al.*, 2002). For instance, *H19* and *Igf2r*, which are usually expressed from maternal alleles are transcribed in androgenetic fetuses consisting of paternal genomes only (Sotomaru *et al.*, 2002). Furthermore, the expression of *U2af1*, which is a paternally expressed gene was confirmed in parthenogenetic fetuses (Ogawa *et al.*, 2006). These

results support the existence of yet unknown mechanism(s), such as those involving *trans*-acting parental alleles (act between parental genomes), that regulate the appropriate expression of imprinted genes (time and location). Because the imbalance of parental genomes in an embryo is one of the innate characteristics of triploid embryos *per se*, studies of expression of imprinted genes in both DAT and DGT embryos might be a promising way to elucidate why triploidy leads to mortality in mammals but not in other animal groups.

Here, we report the morphological features of murine DAT and DGT embryos on E10.5 produced using the pronuclear transplantation technique. We further examined expression patterns of eight primary imprinted genes in each type of triploid fetus.

Results

Developmental competence of triploid mouse embryos during pre- and postimplantation development

We first analyzed the developmental competence of triploid embryos until the blastocyst stage.

Triploid mouse embryos were produced using pronuclear transplantation of one male and female pronucleus from IVF diploid and parthenogenetic haploid embryo into another IVF embryo,

respectively: DAT and DGT embryos (Fig. 1). There were no significant differences in the rate of

blastocyst formation among diploid, DAT, and DGT embryos ($79.2\% \pm 7.1\%$, $86.0\% \pm 3.8\%$, and

$85.6\% \pm 3.4\%$, respectively; mean $\pm$ SEM; Table 1). Next, we assessed the developmental

characteristics of DAT and DGT embryos until E10.5. Compared to diploid conceptuses ($85.2\% \pm$

9.3%), the implantation rate of triploid conceptuses significantly decreased ($P < 0.05$; Table 2). In

addition, the proportions of a live fetus (both DAT and DGT, $5.1\% \pm 2.4\%$ and $5.2\% \pm 2.1\%$,

respectively) were much smaller than in diploids ($53.7\% \pm 5.0\%$; Table 2). The number of resorbed

fetuses was 15 in diploid, 41 in DAT, and 68 in DGT. These results demonstrated that triploidy of the

mouse embryos prevented them from developing beyond the early implantation stage until E10.5.

Morphological characterization of triploid mouse conceptuses

DAT and DGT conceptuses on E10.5 showed contrasting phenotypes in relation to the entire size

and vascularization on the surface of yolk sac (Fig. 2). The size of a DGT conceptus, particularly extraembryonic tissues, was smaller compared to DAT and diploid conceptuses, concomitant with impaired vasculogenesis on the surface of the yolk sac (Fig. 2C). At the same time, there appeared to be no differences in both entire size and vasculogenesis between diploid and DAT conceptuses. In order to assess morphological characteristics more precisely, we analyzed the crown-rump length and embryonic turning of Diploid, DAT, and DGT fetuses on E10.5 (Fig. 2D–G). As expected, the crown-rump length of DGT fetus (1.87 ± 0.13 mm) was significantly smaller than that of diploids (4.81 ± 0.05 mm; $P < 0.01$; Fig. 2H) and DAT (3.86 ± 0.43 mm; $P < 0.01$; Fig 2H).

No significant difference was observed in the crown-rump length between diploid and DAT fetuses. Moreover, all DAT fetuses were morphologically normal, similar to the diploid one (Fig. 2D and 2E). This result is consistent with the observation that all fetuses in both diploid and DAT groups exhibited normal embryonic turning on E10.5 (Fig. 2I). In contrast, the number of fetuses (%) showing embryonic turning in the DGT group significantly decreased, although DGT fetuses with normal turning were found occasionally (Fig. 2F, 2G, and 2I). These results showed that the extra copy of a haploid genome had detrimental effects on fetal development during the early somite stage, ~E10.5. The extent of inhibitory effects on fetal growth depended on whether the extra copy of a haploid genome in a triploid fetus came from a male or female pronucleus.

Expression levels of imprinted genes in triploid mouse fetuses

We wanted to better understand the relationship of morphological characteristics in a triploid fetus on E10.5 with the expression pattern of imprinted genes. To this end, we performed quantitative expression analysis of eight imprinted genes (*H19*, *Gtl2*, *Igf2r*, and *Grb10* among maternally expressed genes and *Igf2*, *Dlk1*, *Ndn*, and *Peg3* among paternally expressed genes) in diploid, DAT and DGT groups by means of real-time RT-PCR (Fig. 3). As expected, the expression level of paternally expressed genes, *Igf2*, *Dlk1*, *Ndn*, and *Peg3*, remained significantly reduced, and that of maternally expressed genes, *Igf2r* and *Grb10*, increased in DGT fetuses ($P < 0.05$; Fig. 3). Additionally, in DAT fetuses, *Igf2* mRNA expression was approximately 2-fold higher than in diploid fetuses.

Unexpectedly, DAT fetuses showed appropriate expression levels equivalent to that of diploid fetuses in the paternally expressed genes *Dlk1*, *Ndn*, and *Peg3*. In addition, among maternally expressed genes, the *H19*, *Gtl2*, *Igf2r*, and *Grb10* mRNA expression levels in DAT fetuses expressed approximately normal levels seen in diploid fetuses. Even though DGT fetuses possessed the redundant maternal haploid set of chromosomes, the *H19* mRNA expression level in DGT fetuses significantly decreased ($P < 0.05$; Fig. 3). These results serve as clear demonstration that DAT fetuses had an almost normal expression pattern of imprinted genes except for *Igf2* mRNA expression, whereas in DGT fetuses, seven out of the eight imprinted genes tested exhibited aberrant

expression patterns, consistent with the results of morphological analysis that DAT fetuses appeared to be relatively normal in contrast to DGT fetuses.

Discussion

Polyploidy exists across all major taxonomical groups of animals including fish and amphibians (Cuellar and Uyeno, 1972; McKinnell, 1964). Mammals also show the types of localized polyploidy, endopolyploidy, where only some tissues are polyploid, for example platelets, bone marrow, hepatic precursor cells, or trophoblast (Brodsky and Uryvaeva, 1977; Carriere, 1969; Ilgren, 1980; Saito *et al.*, 1996). In addition, the diploid-triploid and diploid-tetraploid mouse chimeras could show full term development (Lu and Markert, 1980; Suwinska *et al.*, 2005). Nevertheless, in mammals, polyploidy including triploidy is definitely nonviable (the latter means that one additional haploid set of chromosomes of either maternal or paternal origin is present in the karyotype) (Kaufman *et al.*, 1989a; Kaufman and Webb, 1990). Likewise, it is well-known that uniparental diploidy (the genome consists of either two maternally or two paternally derived haploid sets of chromosomes) causes early embryonic mortality (E9.5), which can be explained by functional differences between parental genomes as a result of genomic imprinting (Surani *et al.*, 1986). Genomic imprinting leads to the parental origin-dependent pattern of gene expression, which means that a group of genes, called *imprinted genes*, is transcribed from one parental allele (Surani *et al.*, 1984). Therefore, the contribution of these complementary parental genetic programs is essential for proper development in mammals (McGrath and Solter, 1984). Nonetheless, there are no data in the literature on the relationship between developmental characteristics of triploid embryos and their expression patterns

of imprinted genes. In this work, we report morphological phenotypes of DAT and DGT embryos at E10.5, where the embryos were created using the pronuclear transfer technique. We examined the expression patterns of eight primary imprinted genes in each type of triploid fetus.

First, we determined the morphological characteristics of DAT and DGT fetuses at E10.5. From the standpoint of fetal growth, DGT fetuses were much smaller than the diploids; in contrast, DAT fetuses were similar to diploids in size. Additionally, the size of both types of triploids never exceeded that of diploids. These results point to two possible explanations as follows. One is that in triploids, the additional haploid set of chromosomes of either maternal or paternal origin has negative effects on postimplantation development (and there are mutual effects of reverse causation). The second explanation is that a triploid genome itself restricts fetal growth via some mechanism that detects the copy number of chromosomes. If we assume that the paternally and maternally derived genomes function for the benefit and detriment of fetal growth, respectively, then the morphological characteristics of triploids in this study are expected and logical results (Kaufman *et al.*, 1989a). Furthermore, the morphological phenotypes of triploid conceptuses show clearer differences between these functions of two parental genomes compared to experimental models that involve parthenogenetic and androgenetic conceptuses. These differences are well pronounced in both fetal and extraembryonic development because androgenetic conceptuses show growth retardation compared to parthenogenetic ones (Surani *et al.*, 1986). In addition, a growing body of

evidence points to the possibility that fetal growth is restricted by a triploid karyotype *per se*, regardless of which side (paternal or maternal) the triploidy is biased toward. This notion may be supported by the observation that DGT embryos show delayed cleavage during the preimplantation period, compared to developmentally matched controls (Takagi and Sasaki, 1976). Although this result was obtained with preimplantation embryos, there might be a delay in triploid embryos in the duration of the cell cycle during the postimplantation period.

In order to gain a further understanding of differences between DAT and DGT fetuses during postimplantation development, we analyzed mRNA expression patterns of eight imprinted genes including four paternally expressed genes and four maternally expressed genes in DAT and DGT fetuses. Among them, *Igf2*, *Dlk1*, *Grb10*, and *Igf2r* are directly responsible for growth and development of fetuses (Charalambous *et al.*, 2003; DeChiara *et al.*, 1990; Ludwig *et al.*, 1996; Moon *et al.*, 2002). The first two genes in the list, strongly promote the fetal size; in contrast, the last two genes, *Grb10* and *Igf2r*, act as a potent growth inhibitor and a mediator, respectively, of insulin-like growth factor signaling, which promotes growth (Charalambous *et al.*, 2003; DeChiara *et al.*, 1990; Ludwig *et al.*, 1996; Moon *et al.*, 2002). As expected, in DGT fetuses, the expression level of paternally expressed genes, *Igf2*, *Dlk1*, *Ndn*, and *Peg3*, remained significantly reduced, and that of maternally expressed genes, *Igf2r* and *Grb10*, increased (Fig. 3). These data are consistent with severe growth retardation of DGT fetuses (Fig. 2). We also found decreased expression of the

Ndn and *Peg3* genes that is responsible for the neurodevelopmental aberrations in DGT fetuses (Broad *et al.*, 2009; Li *et al.*, 1999; Pagliardini *et al.*, 2005; Yang *et al.*, 1998). One early study showed craniofacial anomalies such as axial neural tube defects in DGT fetuses at E9.5 (Kaufman *et al.*, 1989a), and in this study, we also observed severe morphological aberrations in DGT fetuses at E10.5 (Fig. 2). These problems might be due to the reduced expression of *Ndn* and *Peg3* in DGT fetuses.

Contrary to overexpression of *Igf2* in DAT fetuses that is particularly important for fetal growth during embryogenesis (DeChiara *et al.*, 1991), the body size of DAT fetus tended to be smaller than that of Diploid fetus. One possible reason is the polyploid states. Tetraploid fetuses possessing twice DNA contents also show smaller body size than developmentally matched controls (Kaufman and Webb, 1990). This fact indicates that polyploidy per se has negative effect on fetal size for any reason. Therefore, it was possible that fetal-size-promoting effect by *Igf2* mRNA overexpression in DAT fetus was compensated by polyploid state. Additionally, both DAT and DGT fetuses at E10.5 showed unregulated expression of the imprinted genes examined (Fig. 3). A likely explanation is that loss of methylation in these triploid fetuses happened during embryo manipulation or pregnancy, and then caused aberrant expressions of imprinted genes (Doherty *et al.*, 2000; Sato *et al.*, 2007).

When taken together, our results demonstrate that imprinted genes in mouse triploid fetuses show distinctive expression patterns independent of the number of parental-origin haploid sets.

These unregulated expression patterns of imprinted genes observed in triploids suggest that the appropriate number of parental alleles is essential for the maintenance of the parentally specific expression. Further research using triploid mouse embryos could provide a greater insight into the characteristics of mammalian ontogeny governed by both DNA ploidy and asymmetrical functions of parental genomes.

Materials and methods

Preparation of fertilized and parthenogenetic embryos

In order to create diploid fertilized embryos with an external haploid set from the other fertilized/parthenogenetic embryos by means of pronuclear transplantation, we carried out *in vitro* fertilization (IVF) and artificial oocyte activation. The oocyte donors were female B6D2F1 mice (C57BL/6N $\times$ DBA/2). They were superovulated with injections of 7.5 IU (international units) of equine chorionic gonadotropin (eCG; ASKA Pharmaceutical, Tokyo, Japan) and 7.5 IU of human chorionic gonadotropin (hCG; ASKA Pharmaceutical) given 48 h apart. Oocytes at the metaphase of the second meiosis (MII) were removed from the oviducts 16 h after hCG administration, and used for IVF and oocyte activation. All research and experimental protocols were approved by the Regulatory Committee for the Care and Use of Laboratory Animals, Hokkaido University.

IVF was performed using a previously described method (Kono *et al.*, 1993). In brief, spermatozoa were collected from the cauda epididymis of mature B6D2F1 male mice and preincubated in the HTF medium (Quinn *et al.*, 1985) in the humidified atmosphere containing 5% CO₂ at 37 °C for 1.5 h. MII oocytes that were collected as described above were transferred into 100- μ L droplets of the HTF medium. Likewise, the spermatozoa after preincubation were added to the same HTF droplets. Six hours after insemination, the single-cell embryos were washed with the M2 medium (Quinn *et al.*, 1982) and transferred into droplets of the M16 medium containing 0.1

mM ethylene diamine tetraacetic acid (EDTA; DOJINDO LABORATORIES, Kumamoto, Japan) (Whittingham, 1971). The IVF embryos were used as donors and recipients of pronuclear transplantation.

For creation of parthenogenetic embryos, the cumulus cells of collected MII oocytes were removed by means of a brief incubation in hyaluronidase (300 units/mL; Sigma-Aldrich, Tokyo, Japan) in the M2 medium. After the hyaluronidase treatment, stripped oocytes were washed with the M2 medium several times, and transferred into droplets consisting of 50 μ L of the Ca^{2+} -free M16 medium containing 10 mM SrCl_2 (KANTO CHEMICAL, Tokyo, Japan). After incubation for 4–6 h, activated haploid embryos containing both a second polar body and a female pronucleus were collected into M16 droplets at 37 °C in the humidified atmosphere containing 5% CO_2 until they were used for pronuclear transplant as a source of a maternal haploid set of chromosomes.

Pronuclear transplantation

Prior to micromanipulation, donor embryos; i.e., the IVF and activated embryos, were incubated for 1 h in the M16 medium and for 45 min in the same medium containing cytochalasin B (5 μ g/mL; Sigma-Aldrich) and colcemid (0.1 μ g/mL; Wako Pure Chemical Industries, Osaka, Japan). The male pronuclei were easily recognized within the IVF embryos because of the larger features compared to the female pronuclei. The male pronuclei were used as a source of an external haploid paternal

genome for the production of diandric triploid (DAT) embryos (Fig. 1).

For creation of digynic triploid (DGT) embryos, the female pronuclei in the activated embryos (that were produced without diploidization by means of a cytoskeleton inhibitor) were used as a source of external haploid maternal genome (Fig. 1). Either a male or female pronucleus was isolated with a small amount of the cytoplasm from donor embryos and in the presence of inactivated Sendai virus (hemagglutinating activity 2700 U/mL), was inserted into the perivitelline space of another IVF embryo. The IVF embryos that successfully fused with a karyoplast containing a male or female pronucleus were further cultured for 1 h.

***In vitro* culture and embryo transfer**

Both DAT and DGT embryos were cultured in the M16 medium supplemented with 0.1 mM EDTA in the humidified atmosphere containing 5% CO₂ at 37 °C. The number of cleaved embryos and blastocysts was assessed after one and three days of culture, respectively. The resulting blastocysts were transferred into the uterine horns of female mice after 2.5 days of pseudopregnancy. To assess development, the recipients were euthanized on E10.5, and conceptuses were excised from the uterus. Fetuses with noticeable heartbeat were used for the following tests, including measurement of crown-rump length, assessment of embryonic turning, and quantitative real-time RT-PCR analysis of gene expression. The diploid embryos were also cultured and implanted in uterine horns of

pseudopregnant females in the same manner, and the resulting fetuses served as developmentally matched controls.

Quantitative real-time PCR analysis

Total RNA from one fetus on E10.5 in each group, namely, the diandric triploid, digynic triploid, or control diploid group, was prepared using the ReliaPrep RNA Cell Miniprep System (Promega, Madison, WI) according to the manufacturer's instructions. RNA concentrations were assessed using NanoDrop (Thermo Fisher Scientific, Wilmington, DE), and then, quantity of the extracted RNA was normalized to the concentration of 1 µg among the number of samples necessary for one replicate. After normalization of RNA quantity, cDNA samples were synthesized using the ReverTra Ace qPCR RT Master Mix (Toyobo, Osaka, Japan) in a reaction mixture (10 µL) containing 1 µg of the total RNA extracted from each fetus. Finally, the RNA samples from three fetuses in each group were prepared, and we performed quantitative analysis of the gene expression using real-time PCR (LightCycler; Roche Diagnostics, Basel, Switzerland) after preparing the reaction mixture using Thunderbird SYBR qPCR Mix (Toyobo) in triplicate as described previously (n = 3) (Nagatomo *et al.*, 2013). The primers are listed in Supplemental Table S1. The transcript levels were calculated relative to the transcription of the housekeeping gene *Gapdh* in every sample.

Statistical analysis

We compared the means using Student's t test. Data on the number of fetuses with embryonic turning were analyzed using the χ^2 -test. All calculations were performed using the software Statview (Abacus Concepts, Inc., Berkeley, CA). A P value either <0.05 or <0.01 was assumed to denote statistical significance.

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Figure legends

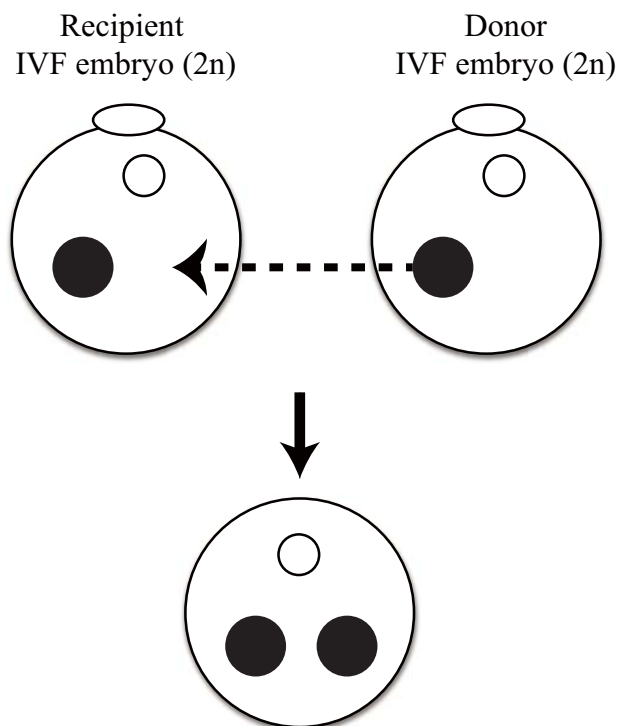
Figure 1. Creation of diandric and digynic triploid mouse embryos using the pronuclear transfer technique. To create a diandric triploid (DAT) embryo ($3n$), we fused a recipient IVF embryo ($2n$) with a male pronucleus (n) derived from another IVF embryo. Digynic triploid (DGT) embryos ($3n$) were created by transferring a female pronucleus derived from an activated haploid embryo (n) into a recipient IVF embryo ($2n$).

Figure 2. Morphological characterization of triploid mouse conceptuses at E10.5. The whole diploid (A), DAT (B), and DGT (C) conceptuses are shown. Note that the DAT conceptus is equivalent in size to the diploid. In contrast, a DGT conceptus is much smaller than the other types, concomitantly with impaired vasculogenesis on the surface of the yolk sac (see the inset). Representative fetuses from diploid (D), DAT (E), and DGT (F and G) embryos are shown. (H) Assessment of the crown-rump length revealed that the length of DGT fetuses (1.87 ± 0.13 mm; $n = 5$) was much smaller than that of diploids (4.81 ± 0.05 mm; $n = 3$; $P < 0.01$) and DAT (3.86 ± 0.43 mm; $n = 4$; $P < 0.01$). Note that there was no obvious difference in the crown-rump length between diploid and DAT fetuses. (I) Evaluation of the number of fetuses with embryonic turning revealed that this percentage was significantly decreased in DGT fetuses ($P < 0.01$), although only two of nine DGT live fetuses with normal turning showed embryonic turning as shown in (G). The scale bar

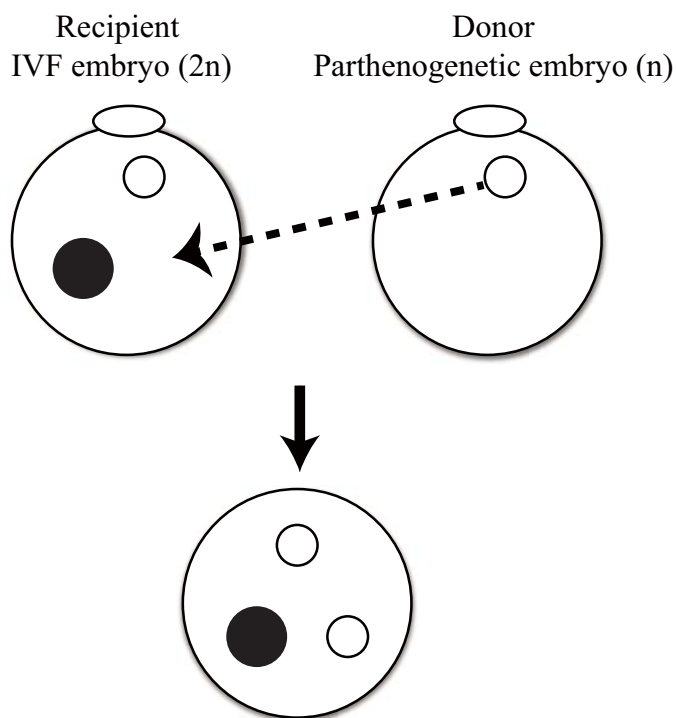
is 1 mm. The data are presented as mean $\pm$ SEM. The asterisks, * and **, indicate $P < 0.05$ and $P < 0.01$, respectively.

Figure 3. Graphical representation of the transcription levels of eight imprinted genes in the triploid mouse fetuses at E10.5. The expression of eight imprinted genes (*H19*, *Gtl2*, *Igf2r*, and *Grb10* among maternally expressed genes and *Igf2*, *Dlk1*, *Ndn*, and *Peg3* among paternally expressed genes) in diploid, DAT, and DGT fetuses (n = 3 each group). The values represent the levels of expression relative to that of an internal control gene (*Gapdh*). Diploid (n = 3, gray), DAT (n = 3, blue), and DGT (n = 3, pink). The data are presented as mean $\pm$ SEM. An asterisk indicates $P < 0.05$.

DAT embryo (3n)

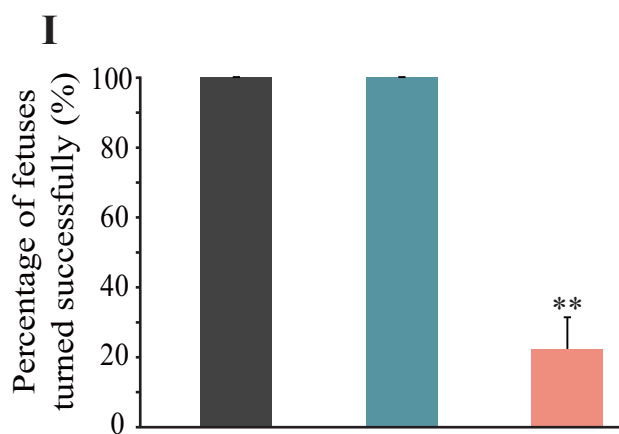
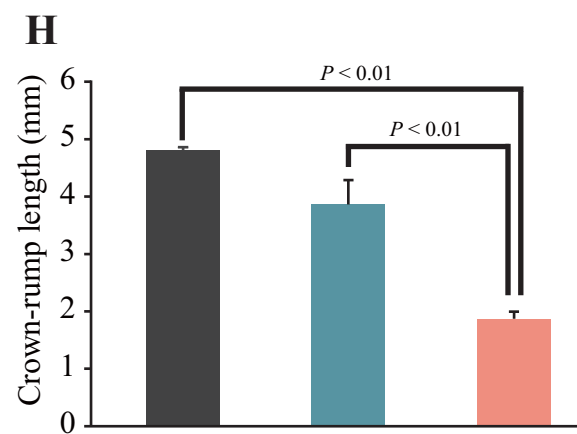
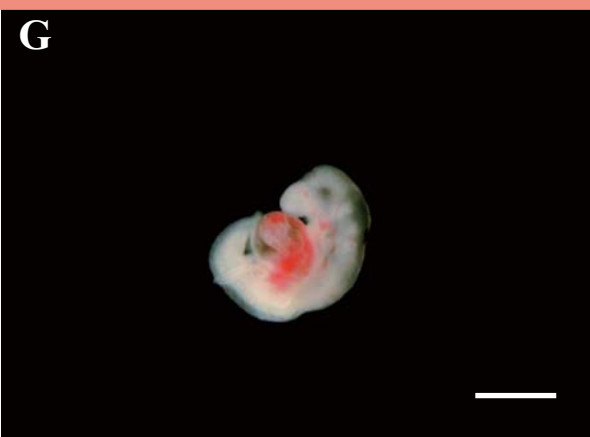
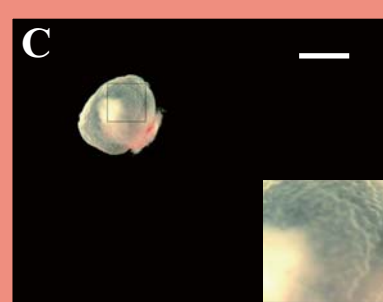
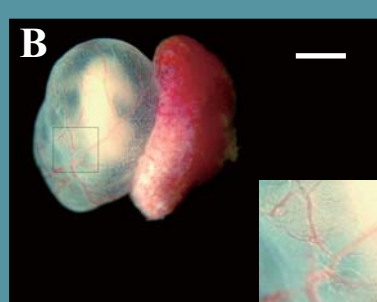
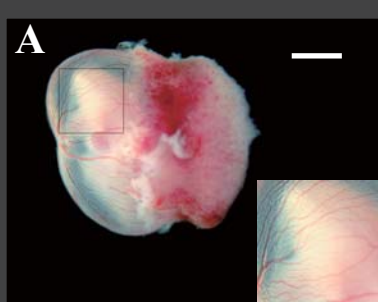


DGT embryo (3n)

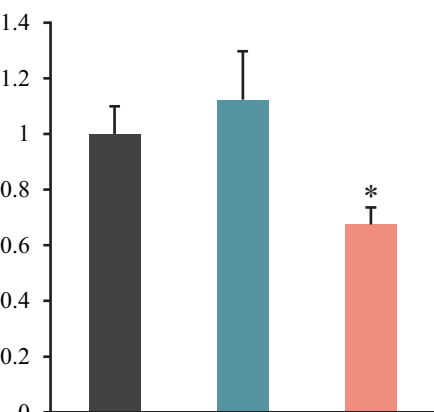
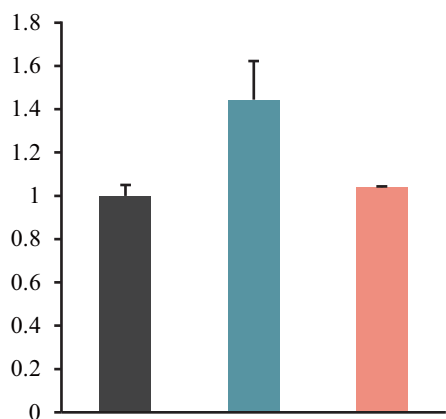
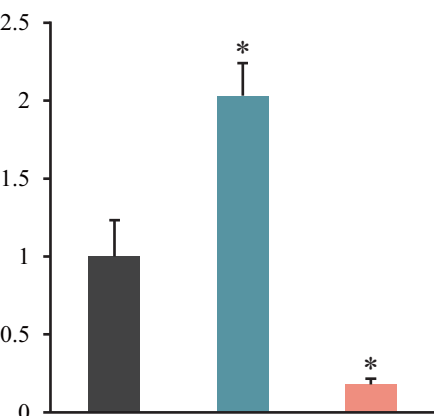
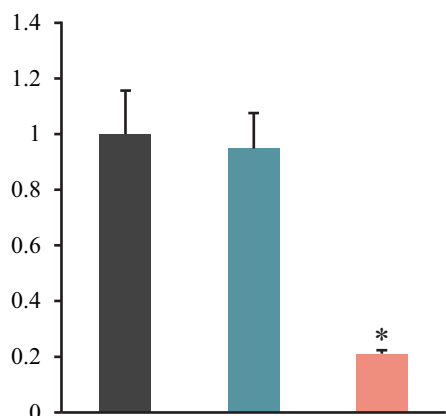
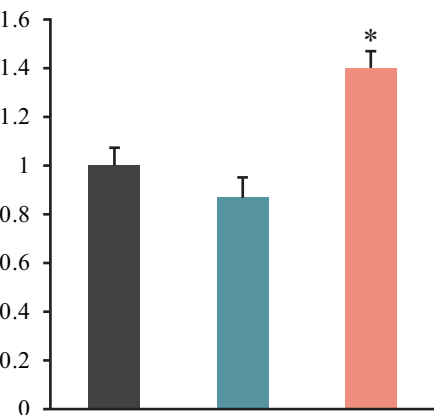
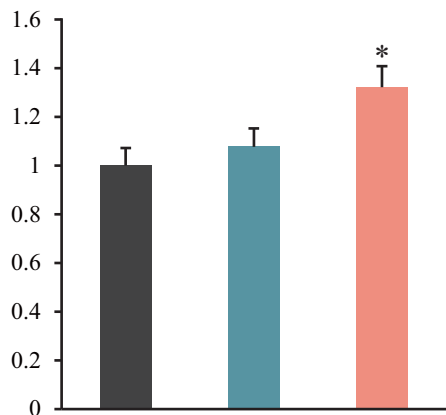
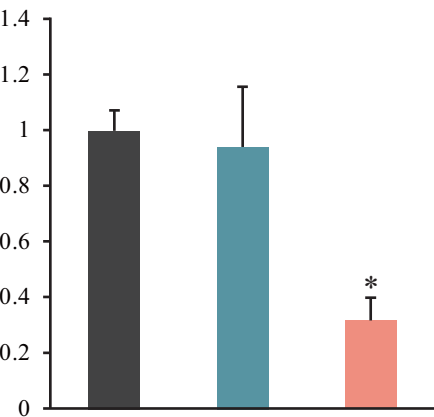


○ Female pronuclear ● Male pronuclear

○ 2nd polar body ← - - - Pronuclear transplantation



Diploid (control)
 DAT
 DGT

H19*Gtl2**Igf2**Dlk1**Igf2r**Grb10**Ndn**Peg3*