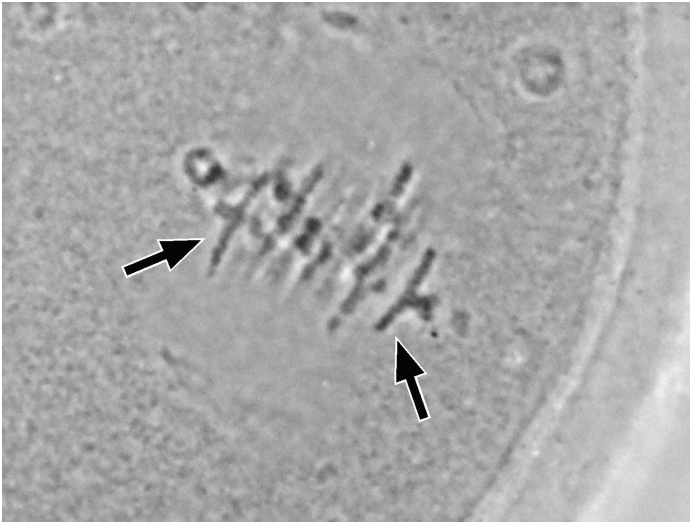


**SUPPLEMENTARY MATERIAL**

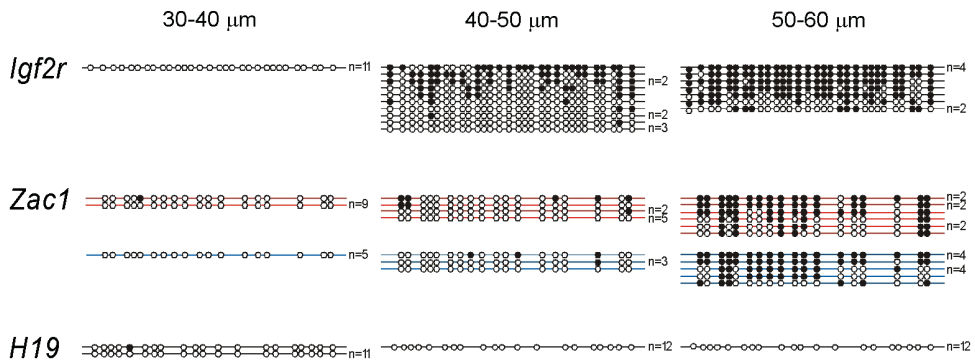
**corresponding to:**

**Large-scale production of growing oocytes *in vitro*  
from neonatal mouse ovaries**

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**Supplementary Fig. 1. Formation of tetrad chromosomes within MII ooplasm.** In vitro-growing oocytes were fused with MII ooplasm using inactivated Sendai virus (Obata and Kono, 2002) and their chromosomal integrity was observed. The donor chromosomes condensed within one hour and showed a typical tetrad composition with some undergoing chromosomal crossover (arrows).



**Supplementary Fig. 2. The methylation status of the differentially methylated regions of three imprinted genes, *Igf2r*, *Zac1*, and *H19*, in oocytes growing *in vivo* in juvenile mice.** Oocytes were collected from juvenile B6D2F1 (C57B/6 X DBA/2JF1) mice and analyzed. The methylation patterns of the two maternally imprinted genes, *Igf2r* and *Zac1*, suggest that genomic imprinting was established in an oocyte-size-dependent manner. The paternally imprinted gene, *H19*, remained unmethylated, irrespective of the size of the oocyte. Because there are DNA polymorphisms between the C57BL/6 and DBA/2 mouse strains in the methylation region analyzed in *Zac1*, the maternal allele sequences and paternal allele sequences are indicated by red and blue lines, respectively.