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Reporting Summary

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For all statistical analysis, confirm that the following items are present in the figure legand, table legand, main text, or Methods section

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n/a	Со	nfirmed
	×	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
×		A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
	×	The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.
x		A description of all covariates tested
x		A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	×	A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
	×	For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i>) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted <i>Give P values as exact values whenever suitable.</i>
x		For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
x		For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
	x	Estimates of effect sizes (e.g. Cohen's d, Pearson's r), indicating how they were calculated
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Software and code

Policy information about availability of computer code

Data collection

The sequencing data was derived from Pacific BioSciences (PacBio) sequencing platform.

Data analysis

Ccs (version 3.4.1) was used to process subreads generated by PacBio SMRTbell sequencing. Minimap2 (version 2.15-r905) was used to align ccs reads to reference genome. TAPIS (version 1.2.1) was used to detect polyadenylation sites. ClusterProfiler (version 3.10.1) was used for gene set enrichment analysis. R (version 3.5.0) was used to plot violin plot, density plot, bar plot and line plot of PacBio data. Statistical analyses were conducted with R. Key scripts used to process and analyze PAIso-seq data is available on GitHub (https:// github.com/niehu2018/GV oocyte PAlsoSeqAnalysis).

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors/reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

The PAIso-seq CCS reads have been deposited into NCBI Sequence Read Archive under the accession number PRJNA529588. The SRA accession number of GV rep.1 and GV rep.2 are SRR8798075 and SRR9130368, respectively. The SRA accessions of 15 single GV oocyte PAlso-seq data are SRR9130400, SRR9130401, SRR9130402, SRR9130403, SRR9130404, SRR9130405, SRR9130406, SRR9130407, SRR9130408, SRR9130409, SRR9130410, SRR9130411, SRR9130412, SRR9130413, SRR9130414. All other data are available upon request. The source data underlying Figs 1b and 4d are provided as a Source Data file.

Field-spe	ocific r	enorting			
		t is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.			
Life sciences		Behavioural & social sciences			
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Life scier	nces s	tudy design			
All studies must dis	sclose on the	se points even when the disclosure is negative.			
Sample size	We choosed	sample size based on literatures in the field and experimental knowledge.			
Data exclusions	No data wer	ere excluded from the analyses.			
Replication	Almost all ex	all experiments shown in this study were performed independently at least three times and no inconsistent results were observed.			
Randomization	Samples we	ere randomly distributed into groups.			
Blinding	Analysis was	objective and did not require blinding.			
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Reportin	g tor s	specific materials, systems and methods			
		rs about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.			
Materials & ex	perimenta	systems Methods			
n/a Involved in the study		n/a Involved in the study			
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Human research participants					
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Animals and	other o	rganisms			
Policy information about studies involving animals; ARRIVE guidelines recommended for reporting animal research					
Laboratory animals CD1 (ICR) mice were female mice between		CD1 (ICR) mice were purchased from Beijing Vital River Laboratory Animal Technology Co., Ltd and bred in our facility. Male and female mice between the age group of 7-8 weeks were used in this study.			
Wild animals The study did not invo		The study did not involve wild animals.			

Field-collected samples Field samples were not collected for the study.

Mice were maintained in compliance with the guidelines of the Animal Care and Use Committee of the Institute of Genetics and Ethics oversight Development Biology, Chinese Academy of Sciences (CAS).

Note that full information on the approval of the study protocol must also be provided in the manuscript.