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# Cyclin B3 controls anaphase onset independent of spindle assembly checkpoint in meiotic oocytes

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**Key words:** Cyclin B3, oocyte, meiosis, SAC, anaphase initiation

Cyclin B3 is a relatively new member of the cyclin family whose functions are little known. We found that depletion of cyclin B3 inhibited metaphase-anaphase transition as indicated by a well-sustained MI spindle and cyclin B1 expression in meiotic oocytes after extended culture. This effect was independent of spindle assembly checkpoint activity, since both Bub3 and BubR1 signals were not observed at kinetochores in MI-arrested cells. The metaphase I arrest was not rescued by either Mad2 knockdown or cdc20 overexpression, but it was rescued by securin RNAi. We conclude that cyclin B3 controls the metaphase-anaphase transition by activating APC/C<sup>cdc20</sup> in meiotic oocytes, a process that does not rely on SAC activity.

## Introduction

The proliferation of all eukaryotic cells depends on the E3 ubiquitin ligase activity of the anaphase promoting complex/cyclosome (APC/C), as without it cells would not enter anaphase and sister chromatids would not separate. The main co-factors of APC/C are Cdh1 and Cdc20; they are the direct activators of APC/C, and they share the same sequence: C-box<sup>1</sup> and IR-tail<sup>2,3</sup>, both mediating their binding to APC/C, and WD40 domain helping to recognize the D-box<sup>4</sup> and KEN-box<sup>5</sup> in the substrates of APC/C.<sup>6</sup> APC/C is a very large complex of 1.5 MDa and it has over 10 subunits; the core subunits have cullin and Ring-finger domains, which are the markers of E3 ubiquitin ligase.<sup>7-9</sup> The Apc2's cullin domain associates with Apc11's Ring-finger domain<sup>3,10,11</sup> and interacts with E2 ligase.<sup>11,12</sup> Without considering the complexity of APC/C, Apc11 itself can perform E3 ligase activity in *vitro*, indicating that other subunits may not be essential for E3 function.<sup>2,10,12</sup>

In both mitosis and meiosis, there are several checkpoints that control cell cycle progression. One of them, the spindle assembly checkpoint (SAC) senses the existence of kinetochores not being attached to spindle microtubules or not being under tension; if the checkpoint is activated, cells are not able to continue mitosis or meiosis. Through this mechanism a cell ensures accurate chromosome segregation to maintain genomic stability, thus avoiding aneuploidy.<sup>13-17</sup> The key target of SAC is Cdc20, the activator of APC/C, and if Cdc20 is inhibited the APC/C will not be able to perform ubiquitin ligase activity thus cyclin B1 and securin

remain stable, thereby delaying the anaphase onset and mitotic exit.<sup>18,19</sup>

SAC proteins recruited onto kinetochores and other multiple components form a large complex between kinetochores and microtubules. The components of SAC are conserved from yeast to humans, indicating that this pathway has been consistent throughout evolution. Aurora B kinase and Mps1 are at the top of this pathway and they are thought to regulate each other,<sup>20-23</sup> the downstream components Bub1, BubR1, Bub3, Mad1 and Mad2 form 3 complexes: Bub1-Bub3, BubR1-Bub3 and Mad1-Mad2.<sup>24</sup> They are recruited to kinetochores in a KMN-dependent manner,<sup>25-27</sup> and when the kinetochore is unattached, the checkpoint is activated.

The function of SAC that inhibits APC/C is mainly related to the activity of the mitotic checkpoint complex (MCC), consisting of BubR1-Bub3, Cdc20 and Mad2.<sup>24</sup> When Cdc20 forms the MCC, Mad2 and BubR1 change Cdc20 binding to APC/C, and Cdc20 loses its function as an activator. Cdc20 is the activator of APC/C in anaphase when it is released from the complex. On the other hand, knockdown of Mad2 causes precocious chromosome separation and cells divide ahead of schedule.<sup>28</sup>

Cyclin B3 is a newly discovered cyclin, and its functions are little known. One study reports that its activity regulates leptotene and zygotene events such as recombination and synapsis in meiotic cells,<sup>29</sup> while another report shows that its accurate expression is required for mitotic anaphase onset in *C. elegans*.<sup>30</sup> Cyclin B3-depleted *C. elegans* embryos have defects in mitotic

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entry and their SAC proteins and dynein accumulate on chromosomes. But aside from these few findings our understanding of cyclin B3s function to date is very limited.

To reveal the functions of cyclin B3 in meiotic cells, we conducted a series of experiments in mouse oocytes. When cyclin B3 was knocked down, oocytes did not undergo anaphase onset during first meiosis. Unlike in *C. elegans*, cyclin B3-depleted mouse oocytes showed SAC inactivation, while cyclin B1 remained undegraded; so we conclude that cyclin B3 affects APC/C activity.

## Results

### Cyclin B3 depletion does not affect GVBD, but arrests oocytes at the MI stage in the presence of normal spindle and chromosome alignment

To clarify the expression of cyclin B3, we collected oocytes at the germinal vesicle (GV), germinal vesicle break down (GVBD), metaphase I (MI), anaphase I, and metaphase II stages.

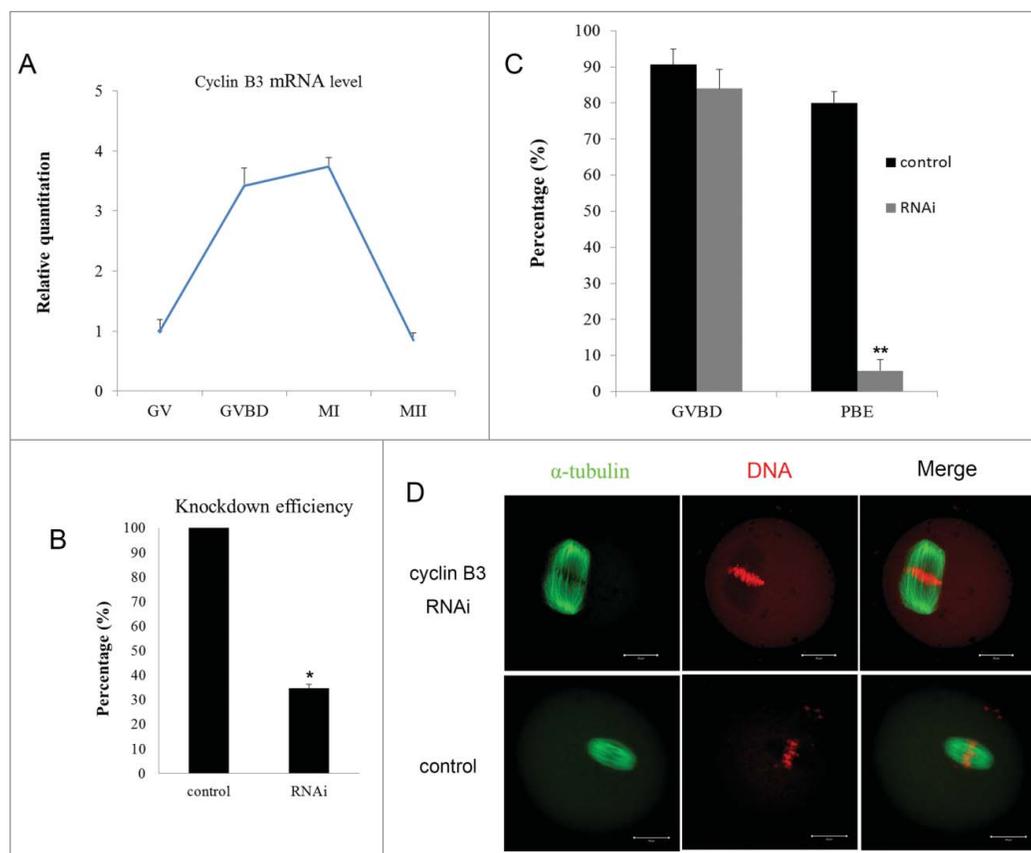
Because a working antibody is not available, we used real-time PCR (RT-PCR) to measure cyclin B3 mRNA expression. We found that in mouse oocytes cyclin B3 expression increased about 3 to fold4- in metaphase compared with the GV stage (Fig. 1A), indicating that cyclin B3 may have functions after GVBD.

SiRNAs (50  $\mu$ M) were injected into GV oocytes to investigate the effects of cyclin B3 depletion. To verify the knockdown efficiency, we mixed 3 siRNAs for injection, and  $65.3 \pm 1.7\%$  of cyclin B3 mRNA was depleted ( $p < 0.05$ ) (Fig. 1B). RNAi oocytes went through the GVBD stages, but they hardly could complete first meiosis, and the polar body extrusion (PBE) rate was only  $5.8 \pm 3.08\%$  (Fig. 1C). In contrast, control oocytes injected with ddH<sub>2</sub>O displayed high rates of GVBD and PBE, and the difference was significant ( $p < 0.05$ ). This phenotype shows that cyclin B3 is a critical regulator of oocyte maturation.

Immunofluorescence staining showed that the spindles of RNAi-injected oocytes were still arrested at metaphase I (Fig. 1D) at 14 hour of culture, which is the time when normal oocytes had extruded their first polar body. Both chromosomes and microtubules were the same as in MI oocytes, and this phenotype remained until 24 hour of culture. Overall, our results show that cyclin B3 depletion does not destroy spindle structure, but blocks oocytes in metaphase.

### Cyclin B3 knockdown causes metaphase-anaphase transition failure in a SAC-independent pathway

As cyclin B3 knockdown blocked oocytes in metaphase I, we wondered if cyclin B3 is essential for inactivation of SAC function. In mouse oocytes, SAC contains Bub1, BubR1, Bub3 and Mad2, and they assemble on kinetochores as dot signals when analyzed with immunofluorescence microscopy. To analyze SAC in clear detail we applied chromosome spread staining and Bub3 antibody staining. At 6 hour of culture, both cyclin B3 RNAi oocytes and control oocytes were observed in typical prometaphase and Bub3 was located on kinetochores (Fig. 2A). Unexpectedly, both groups lost Bub3 signal on kinetochores at 9 hour of culture, indicating that SAC was inactivated in cyclin B3



**Figure 1.** Cyclin B3 regulates meiotic anaphase onset. (A) Oocytes in 4 different maturation stages were collected and used for RT-PCR, each sample contained 100 oocytes. The polyline showed cyclin B3 mRNA changing tendency. (B) Oocytes were injected with mixed cyclin B3 siRNAs (50  $\mu$ M), incubated in M2 medium containing Milrinone for 24 h, then oocytes were washed with Milrinone-free M2 medium to resume meiosis. Knockdown efficiency was measured through RT-PCR. (C) In cyclin B3-depleted oocytes, polar body extrusion rate was observed after 14h of maturation; the RNAi group contained 236 oocytes and the control group contained 241 oocytes. (D) Cyclin B3 RNAi oocytes cultured for 14h were fixed and used for confocal imaging to show the spindles and chromosomes. Bar = 20  $\mu$ M.

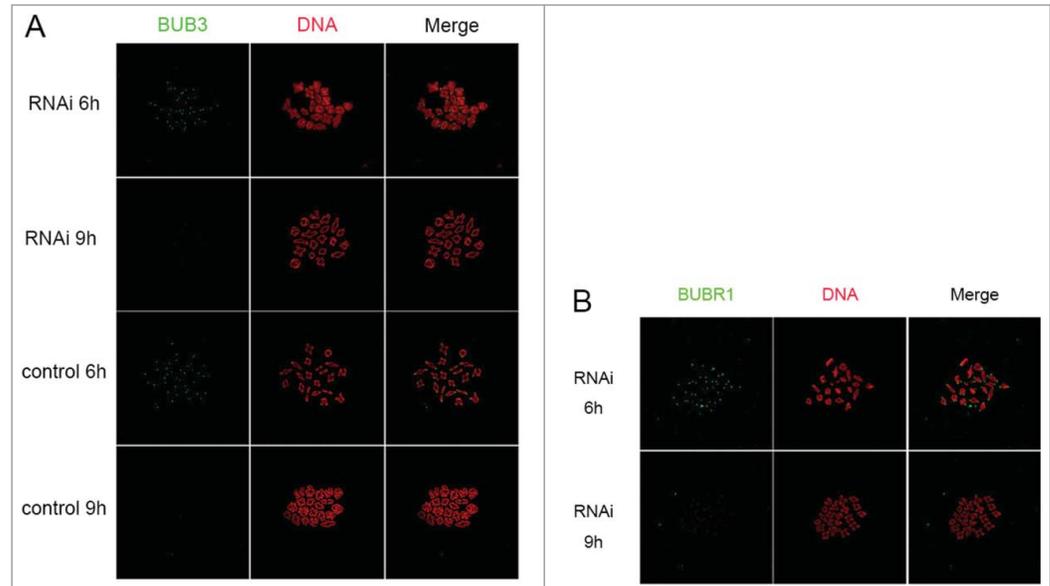
RNAi oocytes (Fig. 2A). To confirm the phenotype we incubated the oocytes with BubR1 antibody, and obtained the same result (Fig. 2B). Through these experiments we conclude that cyclin B3 regulates metaphase-anaphase transition through a SAC-independent pathway.

### Knockdown of Mad2 does not rescue meiotic defects caused by RNAi cyclin B3 depletion

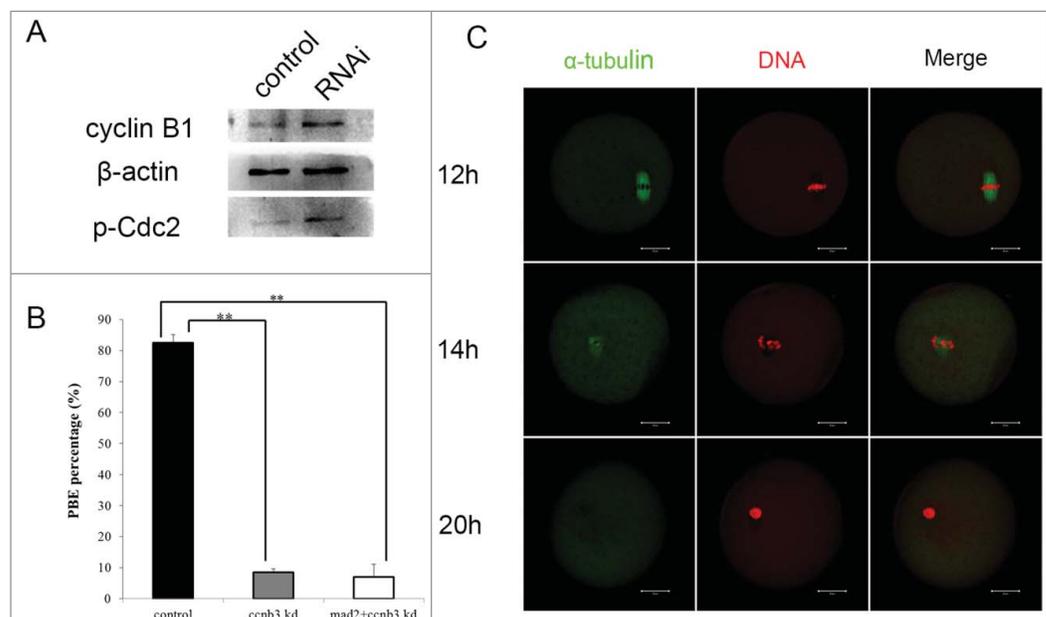
Because we found that SAC inactivation was not affected by cyclin B3 depletion, the mechanism(s) underlying metaphase arrest might be related to later events. Next to the SAC checkpoint, it is the APC/C that controls the metaphase-anaphase progression for which the prerequisites are degradation of cyclin B1 and securin, and inactivation of MPF. Western blotting showed that both cyclin B1 expression and p-CDK1 in control oocytes had already been degraded while in cyclin B3 RNAi oocytes they remained un-degraded (Fig. 3A) at 9.5 hour of culture, providing direct evidence for the correlation between APC/C activation and cyclin B3 function.

In general, APC/C is activated by the regulatory subunit Cdc20 and inhibited by Emi2. Cdc20 forms MCC with SAC components like Mad2 and inhibits APC/C in normal cells; we therefore wondered if cyclin B3 might promote MCC degradation. So we knocked down Mad2 and released Cdc20 from MCC, considering that it might activate APC/C. We mixed cyclin B3 and Mad2 siRNAs (50  $\mu$ M) and injected the mixture into GV oocytes. After more than

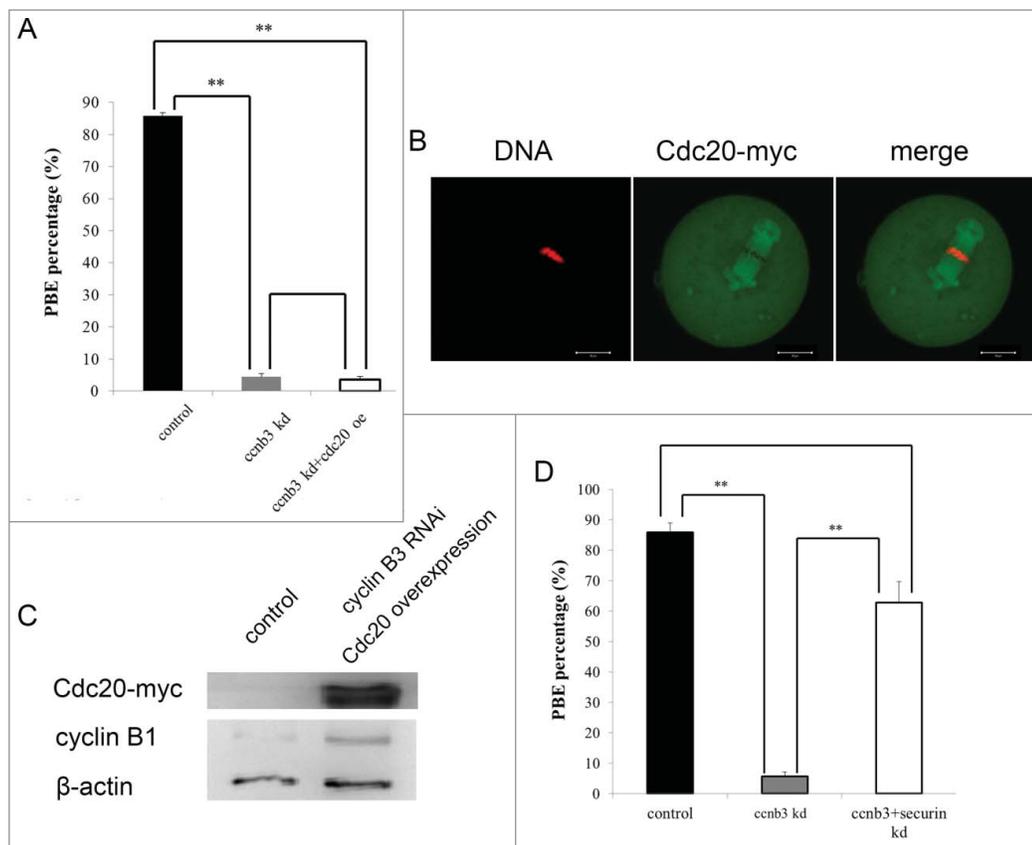
14 hours of culture we analyzed the maturation rates. The oocyte maturation rate in the co-RNAi group ( $6.9 \pm 4.2\%$ ) showed no



**Figure 2.** SAC activity in cyclin B3 RNAi oocytes. (A) Cyclin B3 RNAi oocytes and control oocytes were used for chromosome spread staining at 6h and 9h of culture. Bub3 (green fluorescence) was used as marker of SAC; the red crosses are metaphase chromosomes stained by PI. (B) BubR1 (green fluorescence) staining was used to confirm the phenotype.



**Figure 3.** (A) Cyclin B3 RNAi oocytes and normal oocytes at 9h of culture were collected and used for WB. Cyclin B1 and p-CDK1 first antibodies were incubated to show the MPF activity and  $\beta$ -actin was the reference protein. (B) Cyclin B3 and Mad2 siRNAs were mixed and injected into GV oocytes at a concentration of 50  $\mu$ M, and oocytes were incubated in M2 medium containing Milrinone for 24 h, then they were washed in Milrinone-free M2 medium to resume meiosis. After 14 h the PBE was analyzed. The RNAi group contained 217 oocytes and the control group contained 194 oocytes. (C) Cyclin B3 RNAi oocytes were collected at different times of culture, 12 h, 14 h and 20 h, then used for confocal imaging;  $\alpha$ -tubulin was incubated with antibody displaying green fluorescence and DNA was detected with PtdIns staining. Bar = 20  $\mu$ M



**Figure 4.** The rescue of cyclin B3 RNAi phenotype. (A) GV oocytes were injected with cyclin B3 siRNA (50  $\mu$ M) either alone or with *cdc20* mRNA (5 mg/ml), incubated in M2 medium containing Milrinone for 24 h, then oocytes were washed in Milrinone-free M2 medium to resume meiosis. PBE rates were analyzed at 14 h of culture. The cyclin B3 group contained 224 oocytes, the mixed group contained 189 oocytes and the control group contained 207 oocytes. (B) Oocytes injected with *cdc20* mRNA were incubated with FITC-myc antibody; IF was used to show *Cdc20* expression and location. (C) Cyclin B3 siRNA plus *cdc20* mRNA injected oocytes and control oocytes were collected at 9 h of culture and protein gel blotting was performed. Cyclin B1 showed the MPF activity, myc antibody showed expression of injected *cdc20* mRNA and  $\beta$ -actin was stained as reference protein. (D) Cyclin B3 siRNA and securin siRNA were mixed to a concentration of 50  $\mu$ M and injected into GV oocytes; cyclin B3 alone injected oocytes and normal oocytes were used as control. Oocytes were incubated in M2 medium containing Milrinone for 24 h, then they were washed in Milrinone-free M2 medium to resume meiosis. PBE rates were analyzed at 14 h of culture. The cyclin B3 group contained 211 oocytes, the mixed group contained 208 oocytes and the control group contained 227 oocytes. Bar = 20  $\mu$ M.

difference to that of the cyclin B3 knockdown group ( $8.5 \pm 1.0\%$ ), but was significantly lower than that of the control group ( $82.5 \pm 2.7\%$ ) (Fig. 3B). Though oocytes did not recover to progress through maturation, knockdown of Mad2 caused apoptosis as indicated by chromatin condensation (Fig. 3C). These results further confirm that cyclin B3 activates APC/C downstream of the SAC checkpoint system.

#### Cdc20 overexpression cannot rescue cyclin B3 knockdown phenotype but securin RNAi can rescue the phenotype

Our experiments revealed the function of cyclin B3 in the meiotic cell cycle and we tried to explain the mechanism, but we could only tell that cyclin B3 did not interact with SAC to control the activation of APC/C. If we could activate APC/C directly and rescue the phenotype of metaphase I arrest, we would be able

to narrow down and detail the scope of the investigation, so we applied *Cdc20* overexpression in cyclin B3 RNAi oocytes. Because cyclin B3 siRNA requires an incubation time for about 24 hours, but mRNA usually only requires 2–4 hours to take effect, we tested if the *cdc20*-myc mRNA injection time would make a difference. We showed that a reaction for 24 hours showed the same effect as a reaction for 4 hours, so we mixed cyclin B3 siRNA and *cdc20*-myc mRNA for injection and blocked oocytes at prophase for 24 hour, and we then released the inhibition. Unexpectedly, *Cdc20* overexpression could not increase the oocyte maturation rate, and there was no difference between cyclin B3 knockdown oocytes ( $4.4 \pm 1.2\%$ ) and cyclin B3 knockdown plus *Cdc20* overexpression oocytes ( $3.6 \pm 2.7\%$ ) (Fig. 4A). IF staining was used to show the expression and location of *Cdc20*-myc protein (Fig. 4B). We found that *Cdc20*-myc (green fluorescence) was expressed significantly, and it was mainly located on the spindle. To test APC/C activity we performed another western blotting experiment and analyzed the cyclin B1 level at 9 hour of culture. We could observe an obvious signal of *Cdc20*-myc in *Cdc20* overexpression plus cyclin B3 RNAi oocytes and cyclin B1 was not degraded, while it was degraded in the control group (Fig. 4C). The results indicate that in cyclin B3 knockdown oocytes, APC/C could not be activated only by *Cdc20* overexpression.

To further clarify the mechanisms for the metaphase I arrest in cyclin B3 depleted oocytes, we set out to rescue the phenotype by artificially depleting the substrates of APC/C. As the substrates of APC/C were mainly cyclin B1 and securin, and cyclin B1 was essential for GVBD, we only knocked down securin. SiRNAs were mixed to a concentration of 50  $\mu$ M, and after injection the oocytes were blocked for 24 hours at the GV stage. After release and culture for 12 hours we calculated the PBE rate of each group, and obtained an interesting result:  $62.8 \pm 6.8\%$  of the

oocytes in which both cyclin B3 and securin were knocked down completed maturation (Fig. 4D), thus the phenotype caused by cyclin B3 depletion was rescued. These results confirmed that in cyclin B3 knockdown oocytes, the metaphase I arrest was caused by APC/C inactivation, and that cyclin B3 was an activator of APC/C, but the detailed mechanisms of its function still remain to be clarified.

## Discussion

It is generally accepted that the anaphase onset is strictly controlled by SAC both in mitosis and meiosis. When chromosomes are not properly attached to kinetochores the checkpoint system is activated, which inhibits APC/C from degrading cyclin B1 and securin, thus delaying metaphase-to-anaphase transition. Once SAC is satisfied Cdc20 is released from MCC and APC/C is activated to ubiquitinate cyclin B1 and securin for degradation, and thus separase is activated to cleave cohesin, followed by anaphase initiation. Here, we found a new molecular mechanism controlling the metaphase-to-anaphase transition in meiosis: cyclin B3 is also a strict controlling component of the anaphase onset, a process that is independent of SAC. Depletion of cyclin B3 results in oocyte arrest at metaphase I and this arrest is not controlled by SAC but by direct APC/C inactivation.

Studies of cyclin B3 have been scarce and sometimes results are inconsistent and confusing. Through our study we show that cyclin B3 has multiple functions because it is the activator of APC/C and APC/C plays roles in both prophase and anaphase. The expression pattern of cyclin B3 in oocytes exhibited a typical feature of a protein being activated in metaphase or anaphase, and cyclin B3 RNAi caused a phenotype of metaphase I arrest. Metaphase arrest is typically caused by unattached chromosomes, which activates SAC to delay the anaphase onset; this at first prompted us to hypothesize that cyclin B3 might be the inhibitor of SAC, either related to its transport or to its disassembly (inactivation). However, further experiments showed that this was not the case because neither SAC transport nor disassembly had been affected by cyclin B3 depletion. SAC disappeared from kinetochores at the correct time after reaching metaphase I, and Mad2 RNAi could not rescue the phenotype of metaphase I arrest. So we doubt the idea that cyclin B3 interacts with SAC protein and dynein, as reported in a previous study.<sup>30</sup>

Here we propose that cyclin B3 activates APC/C directly after metaphase, then Cdc20 can combine with APC/C, forming the functional APC/C<sup>Cdc20</sup> to promote cell cycle progression. APC/C is composed of 12 subunits in humans, and its structure is so complicated that we know little about it except for its 3D image. There are many phosphorylation sites on these subunits that either positively or negatively control the complex. We think that cyclin B3 may take part in some of the phosphorylation reactions. Apc2 and Apc 11 together are the catalytic domain<sup>3,31</sup> and they can perform functions like E3 ligase *in vitro*, but *in vivo* their activity is obviously controlled by multiple mechanisms, the classical Cdh1/Cdc20 activation mode and our proposed cyclin B3 control mode. Like other cyclins, cyclin B3 must

have its partner. Molecules that interact with cyclins are mainly CDKs; the typical Cdk1 and Cdk2. Cdk1 is known to combine with cyclin B1 to form MPF, while Cdk2 interacts with cyclin A. Cyclin B3 has both cyclin A and B domains so it is possible for both CDKs to be its partner. Because MPF is not inactivated in our research, it indicates that cyclin B3 acts earlier than Cdk1 releases from MPF. When we knocked down securin the oocytes completed maturation. There are reports in budding yeast that Cdk1 phosphorylates securin and protects it from being degraded by APC/C.<sup>32-34</sup> However, in human cells Cdk1 is less important in the metaphase-anaphase transition, and the main mechanism is that securin and cyclin B1 inhibit separase activity to protect cohesion so that chromosomes are not able to separate.<sup>34,35</sup> Considering others' findings and our results we conclude that securin is the main molecule that protects cohesion.

In summary, our study provides evidence showing that cyclin B3 is a critical regulator of APC/C activity and anaphase onset in oocyte meiosis, and its effect is independent of SAC activity.

## Materials and methods

### Ethics statement

Mouse handling was conducted following policies promulgated by the Ethics Committee of the Institute of Zoology, Chinese Academy of Sciences. ICR mice were housed in the animal core facility which holds a license from the experimental animal committee of the city of Beijing.

### Mouse oocyte collection and *in vitro* maturation

We isolated GV stage oocytes from minced ovaries from 8 to 10 week-old ICR female mice. Oocytes were cultured in M2 medium for at least 12 hours. The culture was conducted in an incubator under environmental conditions of 5% CO<sub>2</sub>, 37°C, and saturated humidity.

### RNA interference

The GV oocytes were microinjected with 5-10 pl of siRNA in M2 medium containing 2.5 uM Milrinone, cyclin B3-1 siRNA GCAGCAGGCUAUUACUAAAtt, cyclin B3-2 siRNA GGCCUCUAUAUGAGGAAUtt, cyclin B3-3 siRNA GAUCAGUGUUUGAAGAUGUtt, Mad21 siRNA GGACUCACCUUGCUUACAAtt, securin siRNA GAUGAUGCCUACCCAGAAAtt, control siRNA UUCUCCGAACGUGUCACGUtt, the concentration of siRNAs was 50 μM. Injected oocytes were incubated in M2 medium containing Milrinone for 24h, then oocytes were washed in Milrinone-free M2 medium and allowed to resume meiosis and further maturation.

### Real-time PCR

Total RNA was extracted from 100 oocytes using RNeasy micro purification kit (Qiagen). Single-strand cDNA generated with the cDNA synthesis kit (Takara), using polyT primers. The cDNAs were used as templates to amplify cyclin B3 and Ppia using the following primers cyclin B3: 5'-

GAAGCAACCATACAAAGAAGCC-3' (forward) and 5'-TTGTCTGGCAGTACAGATGGC-3' (reverse). Real-time PCR was performed using SYBR Premix (Kangwei) in Roche Light Cycler 480. Analysis of relative gene expression was measured by real-time quantitative PCR and the 2(-Delta Delta C (T)) method.

#### Plasmid construction and Cdc20-myc overexpression

We used oocyte total RNA to conduct reverse transcription and generate cDNA as above, cdc20 DNA was amplified using nest PCR. The outer primers were 5'-TGTGTTCGGAGAGCTGAGTACG-3' (forward) and 5'-GGAAGATACGAACCTTATTAGA-3' (reverse), the primers with restriction enzyme site were 5'-CGAATTCGCGCTTGGTTCGCCTTTCGC-3' (forward with EcoRI site) and 5'-CCGGCGCCGCGGAAGATACGAACCTTATTAGA-3' (reverse with AscI site). The 1592bp fragment was then recombined into EcoRI and AscI cutted pCS2+-myc plasmid, transfected into Trans10 competent cells (TransGene). Plasmids were extracted from bacteria using TIANprep Mini Plasmid Kit (TIANGEN) and linearized by Sall then purified by Gel and PCR Clean-Up System (Promega). We used Sp6 MmessageMACHINE kit (Qiagen) to produce capped mRNA and we used RNeasy cleanup kit (Qiagen) to purify it. mRNA was dissolved in nuclease-free water to a concentration of 5.0 mg/ml before injection into oocytes. Injected oocytes were incubated in M2 medium containing Milrinone for 2h (changes in different conditions), then oocytes were washed in Milrinone-free M2 medium and allowed to resume meiosis and maturation.

#### Immunofluorescence (IF) labeling

Oocytes were first fixed in 4% paraformaldehyde at room temperature for 30 min, and then permeabilized in PBS containing 0.5% Triton X-100 for 20 min. Next, oocytes were blocked in PBS with 1% BSA for 1h, and then incubated with first antibody at 4°C overnight. After washing 3 times, they were incubated with second antibody at room temperature for 1 h. DNA was stained with PI. The samples were then mounted on slides. The antibodies used included FITC- $\alpha$ -tubulin (F2618, Sigma), FITC-myc (R953-25, Invitrogen) and Bub 3(sc-28258, Santa Cruz). The oocytes were observed under a laser-scanning confocal microscope (Zeiss LSM 710, Germany). At least 40 oocytes were examined in each treatment, and each treatment was repeated 3 times.

#### Western blotting analysis

First we gathered at least 150 oocytes and boiled for 5 min with loading buffer; then the proteins were separated by 10% SDS-PAGE and electrically transferred to polyvinylidene fluoride membranes. Following this the membranes were blocked in 5% BSA dissolved in TBST for 2 h at room temperature, then incubated with first antibodies overnight at 4°C. After washing 3 times in TBST, 10 min each time, the membranes were

incubated for 1 h at 37°C with secondary antibodies. Finally the membranes were imaged using the enhanced chemiluminescence detection system.

#### Chromosome spread staining

The chromosome spread was performed as described previously (Hodges and Hunt, 2002). Briefly, the oocytes were exposed to acid Tyrode's solution (Sigma) to remove the zona pellucida; the whole process was monitored under the microscope to avoid over-digestion. After a brief recovery in M2 medium, the oocytes were transferred onto glass slides and fixed in a solution of 1% paraformaldehyde in distilled H<sub>2</sub>O (pH 9.2) containing 0.15% Triton X-100 and 3 mM dithiothreitol. The slides were allowed to dry slowly in a humidified chamber for several hours, and then blocked with 1% BSA in PBS for 1h at room temperature or overnight at 4°C and incubated with primary antibodies of anti-Bub3 or anti-BubR1 (1:50), overnight at 4°C. After brief washes with washing buffer, the slides were then incubated with corresponding secondary antibodies for 2h at room temperature. DNA on the slides was stained with PtdIns for 10 min and slides were mounted for immunofluorescence microscopy observation.

#### Data analysis

All experiments were repeated at least 3 times. Statistical analysis was performed using Student's T test and shown as mean  $\pm$  SEM.

#### Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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#### Note

While we were preparing this manuscript, a paper was published to show that cyclin B3 is a mitotic cyclin that promotes the metaphase-anaphase transition (Yuan K, O'Farrell PH. Cyclin B3 is a mitotic cyclin that promotes the metaphase-anaphase transition. *Curr Biol* 2015; 25(6):811-6). Our conclusion in meiosis is similar to that in mitosis.

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