Initiation of Cyclin B Degradation by the 26S Proteasome upon Egg Activation

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Abstract. Immediately before the transition from metaphase to anaphase, the protein kinase activity of maturation or M-phase promoting factor (MPF) is inactivated by a mechanism that involves the degradation of its regulatory subunit, cyclin B. The availability of biologically active goldfish cyclin B produced in Escherichia coli and purified goldfish proteasomes (a nonlysosomal large protease) has allowed the role of proteasomes in the regulation of cyclin degradation to be examined for the first time. The 26S, but not the 20S proteasome, digested recombinant 49-kD cyclin B at lysine 57 (K57), producing a 42-kD truncated form. The 42-kD cyclin was also produced by the digestion of native cyclin B forming a complex with cdc2, a catalytic subunit of MPF, and a fragment transiently appeared during cyclin degradation when eggs were released from metaphase II arrest by egg activation. Mutant cyclin at K57 was resistant to both digestion by the 26S proteasome and degradation at metaphase/anaphase transition in Xenopus egg extracts. The results of this study indicate that the destruction of cyclin B is initiated by the ATP-dependent and ubiquitin-independent proteolytic activity of 26S proteasome through the first cutting in the NH₂-terminus of cyclin (at K57 in the case of goldfish cyclin B). We also surmise that this cut allows the cyclin to be ubiquitinated for further destruction by ubiquitin-dependent activity of the 26S proteasome that leads to MPF inactivation.
coli and of purified goldfish proteasomes allows the role of proteasome in the regulation of cyclin degradation to be examined for the first time. Here we propose that the 26S proteasome initiates cyclin degradation through the first cut in its NH₂ terminus.

Materials and Methods

Materials

Goldfish were purchased from a local supplier and maintained at 15°C until use. The 20S and 26S proteasomes were purified from immature goldfish oocytes by conventional column chromatography as described (Toku moto et al., 1995a,b). X. laevis were obtained from a dealer and maintained at 20°C. X. laevis CSF-arrested egg extracts were prepared by the method of Murray et al. (1989).

Electrophoresis and Immunoblot Analysis

Electrophoresis proceeded as described by Laemmli (1970), using 12.5% gels under denaturing conditions. Cyclin B degradation was assessed by immunoblotting against anti-goldfish cyclin B (B63 and B112) monoclonal antibodies (Yamashita et al., 1992b). Immunocomplexes were visualized using the ECL detection kit (Amersham Intl., Arlington Heights, IL).

Determination of the Digestion Site of Cyclin B

Electrobotted 42-kD fragments of cyclin B were prepared as described (LeGendre and Matsudaira, 1989). The NH₂-terminal amino acids were determined using a protein sequencer (470A; Applied Biosystems, Chiba, Japan).

Production of Recombinant Cyclin Bs

Full length (Δ0) and NH₂-terminal truncated (Δ41 and Δ68) goldfish cyclin Bs were produced as described (Hirai et al., 1992; Katsu et al., 1993; Yamashita et al., 1995). Mutant cyclin B in which lysine 57 was replaced by arginine (Δ0K57R) was produced as follows. A cDNA clone encoding full length goldfish cyclin B (Hirai et al., 1992) was mutated using a site-directed mutagenesis system (Mutan K; Takara, Tokyo, Japan), following a strategy based on the method of Kunkel (1985), according to manufacturer’s instructions. Double-strand, mutated cDNA was prepared by T3 polymerase using single-strand cDNA and the following oligonucleotide:

AAGAAGGAAGTG

This oligonucleotide was designed to produce a mutation site (bold type) and a restriction enzyme site (underlined) as described (Yamashita et al., 1995). Mutant clones were screened by digestion with the restriction enzyme and confirmed by sequencing.

Recombinant proteins were produced in E. coli BL21 (DE3) and purified by SDS-PAGE followed by electroelution from the gel, as described previously (Hirai et al., 1992). ³⁵S-labeled cyclins were produced using a TNT T7-coupled Reticulocyte Lysate System (Promega Biotech, Madison, WI) according to the manufacturer’s instructions.

Results

Restricted Digestion of Cyclin B by 26S Proteasome

Although the 26S proteasome is a ubiquitin-dependent protease in general (Armon et al., 1990; Driscol and Goldberg, 1990; Kanayama et al., 1992), it also catalyzes an ATP-dependent and ubiquitin-independent proteolysis (Tanka et al., 1983; Matthews et al., 1989; Murakami et al., 1992). Therefore, we initially investigated whether or not the 26S proteasome can degrade in vitro nonubiquitinated, full length goldfish cyclin B produced in E. coli (cyclin Δ0). Goldfish proteasomes (20S and 26S) were purified from immature goldfish oocytes by sequential chromatography (Tokumoto et al., 1995a,b). SDS-PAGE has demonstrated that the 26S proteasome consists of multiple subunits with a molecular mass ranging from 23.5 to 140 kD, whereas the 20S proteasome includes subunits ranging from 23.5 to 31.5 kD (Fig. 1 A). The 26S, but not the 20S proteasome, digested 49-kD cyclin Δ0 and produced a 42-kD cyclin (Fig. 1 B). Protease inhibitors of microbial origin (anti-pain, chymostatin, and leupeptin) blocked the cyclin digestion at high (200 μM) but not at low concentrations (50 μM). Protease inhibitors (MG115, MG132, and PSI) that inhibit the chymotrypsin-like activity and ubiquitin-dependent protein degradation (Figueiredo-Pereira et al., 1994; Rock et al., 1994; Jensen et al., 1995), blocked the digestion at 50 μM (Fig. 1 C). No digestion proceeded when

![Figure 1. Digestion of E. coli-produced goldfish cyclin B by 26S proteasome purified from immature goldfish oocytes. (A) Subunit composition of 20S and 26S proteasomes. Purified 20S (10 μg) and 26S (27 μg) proteasomes were resolved by electrophoresis and stained with Coomassie brilliant blue R-250. (B) Digestion of full length cyclin B by purified proteasomes. Cyclin Δ0 (5 μg/ml) was incubated at room temperature with purified 20S or 26S proteasomes (60 μg/ml) in reaction buffer (100 mM Tris-HCl, 5 mM MgCl₂, 0.04 mM ATP, pH 7.6). Samples were exposed to Laemmli’s SDS sample buffer at the indicated times during incubation. Cyclin B was detected by immunoblotting against an anti–goldfish cyclin B (B63) monoclonal antibody. The position to which the digested cyclin B migrated is indicated by an asterisk. (C) Effect of protease and proteasome inhibitors on cyclin B digestion by the 26S proteasome. Cyclin Δ0 was incubated for 60 min without (None) or with the 26S proteasome in the absence (Control) or presence of various inhibitors at 50 μM. Cyclin B was detected by the B63 antibody. The position of the digested cyclin B is indicated by an asterisk.](jcb.rupress.org)
the 26S proteasome was depleted with an anti-proteasome antibody (Fig. 2, A and B). The reaction was also prevented when ATP was depleted from the reaction mixture (Fig. 2 C). These results indicate that the digestion of cyclin B is not due to a contaminating protease in the 26S proteasome fraction, but is catalyzed by the 26S proteasome itself.

We determined whether the 26S proteasome digests the NH$_2$- or COOH-terminal region of the cyclin B. We used monoclonal anti-cyclin B63, which recognizes the COOH-terminal region of goldfish cyclin B. The site digested by the 26S proteasome (COOH terminus of K57) and truncated sites of deletion mutants (Δ41, Δ68) are indicated. The destruction box and lysine-rich stretch are also indicated. (B) Digestion of full length and truncated cyclin Bs by the 26S proteasome. Cyclins Δ0, Δ41, and Δ68 were incubated in the absence (−) or presence (+) of the 26S proteasome for 120 min at room temperature. Cyclin degradation was assessed by immunoblotting against two kinds of anti-cyclin B (B63 and B112) monoclonal antibodies. B112 recognizes the NH$_2$-terminal portion of goldfish cyclin B. The position of the digested cyclin B is indicated by an asterisk.
The NH₂-terminal region of cyclin B, including the destruction box and half of the lysine-rich stretch determined. The site of cyclin B cleavage by the 26S proteasome in vitro. We produced two NH₂-terminal truncated cyclins: cyclin A (60 μg/ml) for 120 min in the absence (Control) or presence of various concentrations of B2Nt or lysozyme. Cyclin B was detected with the B63 antibody. The migrating position of the digested cyclin B is indicated by an asterisk.

The NH₂-terminal sequences of cyclins, including a consensus sequence called the destruction box, play a critical role in targeting cyclins for degradation (Glotzer et al., 1991; Lorca et al., 1991; Luca et al., 1991; Kobayashi et al., 1992). We therefore examined the role of NH₂-terminal sequences in cyclin digestion by the 26S proteasome in vitro. We produced two NH₂-terminal truncated cyclins: cyclin Δ41 lacking the destruction box and cyclin Δ68 lacking the destruction box and half of the lysine-rich stretch (Fig. 3A). Neither cyclin Δ68 nor cyclin Δ41 were digested by 26S proteasome (Fig. 3B). This finding suggests that the NH₂-terminal region of cyclin B, including the destruction box, supplies an interaction site between cyclin B and 26S proteasome that is necessary for the subsequent cutting of cyclin B at K57. Interaction between the NH₂-terminus of cyclin B and the 26S proteasome was also suggested by an experiment with a truncated protein containing the first 89 amino acids of Xenopus cyclin B2 (B2Nt, a gift from Dr. M.J. Lohka, University of Calgary, Calgary, Canada; Velden and Lohka, 1993). B2Nt inhibited the digestion of cyclin B by the 26S proteasome in a dose-dependent manner (Fig. 4). Control lysozyme, a basic and low molecular weight protein like B2Nt, did not inhibit the cyclin digestion (Fig. 4).

The 26S proteasome also digested native cyclin B that had been isolated as a complex with cdc2, yielding a truncated cyclin of ~42-kD (Fig. 5A). This was the same size as the fragment produced by digestion of recombinant, full length cyclin B with the 26S proteasome. The digestion of the cdc2–cyclin B complex with 26S proteasome, however, did not cause kinase inactivation of cdc2 (Fig. 5B).

**Detection of Intermediate Cyclin B during Egg Activation**

Mature goldfish oocytes are arrested at metaphase of meiosis II (metaphase II) and can be activated by the contact with water (Yamashita et al., 1992a,b). Cyclin B was degraded within a few minutes after egg activation (Nagahama et al., 1995). A 42-kD fragment of cyclin B transiently appeared during the initial phase of goldfish egg activation. In a partially purified and highly concentrated fraction from egg extracts, intermediate cyclin B was detected 3 min after egg activation (Fig. 5C). The monoclonal antibody B112, which recognizes the NH₂-terminal region of cyclin B, did not react with the intermediate (Fig. 6C). These results suggest that NH₂-terminal digestion of cyclin B by the 26S proteasome is not an artifact of in vitro proteolysis but an initial reaction of cyclin B degradation that proceeds upon egg activation.

**Destruction Analysis in Xenopus Egg Extracts**

Although the results so far suggested that the 26S proteasome interacts with the NH₂-terminal region of cyclin B and then cuts it at K57, digestion of cyclin B by the 26S proteasome in vitro is limited to cleavage of a single peptide bond and does not induce kinase inactivation of MPF (Fig. 5B). We believe that the incomplete digestion of cyclin B by the 26S proteasome in vitro is due to the absence of factors responsible for further degradation of cyclin B. The most likely candidate is ubiquitin and its ligase. In fact, goldfish oocytes contain high levels of free ubiquitins (Tokumoto et al., 1993b).

We purified and characterized the 26S proteasome from immature Xenopus oocyte extracts (Tokumoto and Ishikawa, 1995). We then examined whether or not this proteasome digests goldfish cyclin B, like the goldfish proteasome can. Xenopus 26S, but not 20S proteasomes, digested the NH₂ terminus of goldfish cyclin B and produced the 42-kD intermediate (Fig. 6A and C). NH₂-terminal truncated cyclins were not digested by the Xenopus 26S proteasome (Fig. 6B). These results indicate that the Xenopus 26S proteasome can digest goldfish cyclin B, suggesting a similar role of goldfish and Xenopus proteasomes in the regulation of cyclin degradation. We then examined the involvement of 26S proteasome in cyclin B degradation using a Xenopus cell-free system widely used for cell cycle studies, which contains the complete system necessary for cyclin degradation (Murray et al., 1989).

Cyclin Δ0 was completely degraded within 30 min after adding Ca²⁺ to Xenopus egg extracts, although it was stable in the absence of Ca²⁺ (Fig. 7A). NH₂-terminal truncated cyclins Δ41 and Δ68 were not degraded in Xenopus egg extracts even after activation with Ca²⁺ (Fig. 7B). In contrast to cyclins Δ41 and Δ68, the 42-kD cyclin fragment (cyclin Δ57), which had been produced by the prior digestion of cyclin Δ0 with the purified 26S proteasome, was degraded in Xenopus extracts after adding Ca²⁺ (Fig. 8). To exclude the possibility that the 42-kD cyclin fragment remained in complex with the NH₂-terminal portion after digestion with the 26S proteasome, we performed gel chromatography under the buffer conditions that were used for cyclin digestion. When cyclin treated with 26S proteasome...
was chromatographed, 42-kD cyclin and NH₂-terminal fragment (∼9 kD) were clearly separated (Fig. 9). This result indicates that there is no significant interaction between 42-kD cyclin and the NH₂-terminal fragment after digestion.

**Digestion by the 26S Proteasome and Degradation of In Vitro Translated Cyclin B₅ in Xenopus Egg Extract**

Our experiment using *E. coli*-produced cyclins demonstrated that the 26S proteasome restrictively digests the NH₂ terminus of cyclin B and that only cyclins digestible or digested by 26S proteasome are degradable in *Xenopus* egg extracts activated with Ca²⁺. To confirm that these results are not artifacts derived from the incorrect form of cyclins produced in *E. coli*, we performed a similar experiment using cyclin B proteins translated in vitro in rabbit reticulocyte lysate. Cyclin B proteins produced in *E. coli* and translated in vitro gave the same results. Cyclin D₀ translated in vitro was digested by the 26S, but not the 20S proteasome (Fig. 10 A), and degraded in *Xenopus* egg extracts (Fig. 10, C and D). NH₂-terminal truncated cyclins D₄₁ and D₆₈ were resistant to 26S proteasome digestion (Fig. 10 B) and to degradation in *Xenopus* egg extracts (Fig. 10, C and D). Furthermore, the point mutant, cyclin D₀K₅₇R (in which the position of cleavage by the 26S proteasome, lysine 57, was converted to arginine), was neither digested by the 26S proteasome nor degraded in *Xenopus* extracts (Fig. 10, B–D).

**Discussion**

Immediately before the transition from metaphase to anaphase, the kinase activity of MPF is inactivated through the degradation of its cyclin B subunit. The mechanism of cyclin degradation, which must be a highly selective process since few other proteins are degraded only at this time, is poorly understood. Using recombinant goldfish cyclin B and purified 26S proteasomes, we investigated the role of proteasomes in the regulation of cyclin degradation during egg activation. We found that purified 26S proteasome digests not only recombinant cyclin B, but also native cyclin B at its NH₂-terminal portion, producing a 42-kD intermediate form. Since the 42-kD cyclin B appears transiently during the initial phase of normal egg activation, this digestion should not be an artifact but rather an initial step in cyclin B degradation upon egg activation.

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**Figure 5.** Digestion of native cyclin B by 26S proteasome. The truncated cyclin B produced by the 26S proteasome digestion is indicated by an asterisk. (A) Digestion of cyclin B in MPF complex by the 26S proteasome. The MPF complex in mature carp oocytes was prepared using suc1 beads (Yamashita et al., 1992b). The beads were washed with buffer (50 mM Tris-HCl, 20% glycerol, 10 mM 2-mercaptoethanol, 0.1 mM ATP, pH 7.5) and shaken in the absence (−) or presence (+) of 60 μg/ml of the 26S proteasome at room temperature with agitation. Samples were treated with SDS sample buffer at the indicated times and immunoblotted against the B63 antibody. Two cyclin bands were detected, and only the upper band was digested by the 26S proteasome. It is unlikely that these two bands correspond to different phosphorylation states of cyclin B (Yamashita et al., 1992b). In C, only a single band of cyclin B was detected when oocytes were directly exposed to SDS sample buffer. Therefore, the lower band is probably produced by undesirable proteolysis during treatment with the suc1 beads. (B) Protein kinase activity of suc1 precipitates before and after the digestion with 26S proteasome. The kinase activity of suc1 precipitates incubated for 60 min in the absence (−) or presence (+) of 26S proteasome was measured with a synthetic peptide substrate for cdc2, as described (Yamashita et al., 1992a). Activities are indicated as a percentage of the activity at 0 min for each condition. (C) Detection of a truncated cyclin B during goldfish egg activation. Ovulated eggs (2 ml) were placed in 3 ml goldfish Ringer’s solution (Yamashita et al., 1992b) and immediately homogenized in 5 ml SDS sample buffer at the indicated times. Before detecting truncated cyclin B by immunoblotting with B63 or B112, proteins with a molecular mass of 40–50 kD were separated by SDS-PAGE (Prep Cell Model 491; Bio Rad, Richmond, CA) and concentrated.
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Figure 6. Digestion of goldfish cyclin B by the Xenopus 26S proteasome. Cyclin was visualized by immunoblotting with B63 (A–C) and B112 (C). The position of the digested cyclin B is indicated by asterisks. (A) Digestion of full length cyclin B. Cyclin Δ0 (5 μg/ml) was incubated at room temperature with purified 20S or 26S proteasomes (60 μg/ml) in the reaction buffer (100 mM Tris-HCl, 5 mM MgCl₂, 0.04 mM ATP, pH 7.6). Samples were exposed to Laemmli’s SDS sample buffer at the indicated times during incubation. (B) Digestion of truncated cyclin B. Cyclins Δ0, Δ41, and Δ68 were incubated in the absence (−) or presence (+) of 60 μg/ml of 26S proteasome for 120 min at room temperature. (C) Cyclin was digested by Xenopus 26S proteasomes at the NH₂ terminus. Goldfish cyclin Δ0 was digested by 26S proteasome for the indicated times and then stained with B63 or B112 antibody.

Figure 7. Degradation of goldfish cyclin B in Xenopus egg extracts. Cyclin B was detected with B63 antibody. (A) E. coli–produced goldfish cyclin Δ0 was added to Xenopus egg extract at a final concentration of 5 μg/ml. Incubations proceeded in the absence (−Ca²⁺) or presence (+Ca²⁺) of 0.4 mM CaCl₂ for the indicated times. (B) E. coli–produced cyclin Δ0, Δ41, and Δ68 were added to Xenopus extracts at the final concentration of 5 μg/ml. Cyclin degradation was induced by 0.4 mM Ca²⁺ and terminated by adding SDS sample buffer at the indicated times.

Using Xenopus egg extracts, we also showed that the initial digestion and further degradation of cyclin B are tightly linked. Cyclins Δ41 and Δ68 that are indigestible by the 26S proteasome are not degraded, whereas cyclins Δ0 and Δ57 digested by the 26S proteasome, are degraded. These findings strongly suggest that the initial cutting of the NH₂-terminal region of cyclin B by 26S proteasome is a prerequisite for the subsequent degradation that leads to the inactivation of MPF at the metaphase/anaphase transition.

The NH₂-terminal sequences of cyclin B, including a consensus sequence that is called the destruction box, play

Figure 8. Degradation of intermediate cyclin B (cyclin Δ57) in Xenopus egg extracts. Cyclin Δ57 was obtained by digesting cyclin Δ0 with 26S proteasome. 1/20 vol of the digestion mixture containing cyclins Δ0 and Δ57 was added to Xenopus extracts, and cyclin degradation was examined in the absence (−Ca²⁺) or presence (+Ca²⁺) of 0.4 mM Ca²⁺. Samples were exposed to SDS sample buffer at the indicated times. Cyclin degradation was assessed by immunoblotting with B63 antibody. The position of cyclin Δ57 is indicated by an asterisk.
a critical role in targeting cyclins for degradation, since truncated sea urchin (Murray et al., 1989), human (Lorca et al., 1991), and clam (Luca et al., 1991) B-type cyclins missing the first 90, 72, or 97 amino acids, respectively, and clam (Luca et al., 1991) and Xenopus (Kobayashi et al., 1992) A-type cyclins missing the NH\textsubscript{2}-terminal 60 or 62 amino acids are resistant to degradation. Each of these truncated cyclins continuously activates cdc2, which prevents cells or cellular extracts from leaving mitosis. A truncated protein containing only the first 89 amino acids of Xenopus cyclin B2 (B2Nt), including sequences essential for cyclin degradation in other species, also inhibited cyclin degradation (Velden and Lohka, 1993). These results indicate interaction of the NH\textsubscript{2}-terminal portion of cyclin with the destruction machinery.

In this study, we proposed that the initial reaction of cyclin B destruction is the restricted cleavage of its NH\textsubscript{2}-terminal portion (K57 in the case of goldfish cyclin B) by the 26S proteasome. As well as cyclin D\textsubscript{68} that lacks the cutting site K57, cyclin D\textsubscript{41} containing K57 was not cleaved, indicating that the NH\textsubscript{2}-terminal region affords not only the cutting site but also the interaction site necessary for digestion by the 26S proteasome. This notion was confirmed by inhibiting the cyclin digestion with B2Nt that consists of the first 89 amino acid of Xenopus cyclin B2. Cyclins D\textsubscript{41} and D\textsubscript{68} were neither digested by the 26S proteasome nor degraded in Xenopus egg extracts activated by Ca\textsuperscript{2+}, whereas cyclin D0 was digested at K57 by 26S proteasome and degraded in the extracts. In addition, cyclin D57 produced by digesting cyclin D0 with the 26S proteasome, was degraded in the extracts. These results suggest that only cyclins that have undergone 26S proteasome digestion at K57 can be degraded upon egg activation.

The mechanism of cyclin B degradation after initial cleavage by the 26S proteasome remains to be determined. The most likely candidate will be ubiquitin-dependent proteolysis. Proteins to be degraded by the ubiquitin pathway are ligated to ubiquitin through their lysine amino acid groups and then degraded by the 26S proteolytic complex (Hershko and Ciechanover, 1982). The first evidence that cyclin B degradation is mediated by ubiquitin-dependent proteolysis was provided by Glotzer et al. (1991). Other support for the involvement of a ubiquitin-dependent pathway in the cyclin degradation arises from the observation that methylated ubiquitin, which prevents the polyubiquitination of proteins destined for degradation, delays cyclin degradation in an extract from clam embryos (Hershko et al., 1991). A complex containing cyclin-selective ubiquitin ligase activity has been identified in clam oocytes (Sudakin et al., 1995). These findings suggest that the cell cycle-specific cyclin degradation is mediated by a ubiquitin-dependent proteolytic system.

The cyclin B subunit of MPF must be ubiquitinated immediately before the onset of its destruction at the metaphase/anaphase transition. The findings of the present study indicate that the restricted cleavage of cyclin B triggers its ubiquitination. It is likely that the digestion of the

Figure 9. Separation of intermediate cyclin B and NH\textsubscript{2}-terminal fragment by gel chromatography. (A) Sephadex G-50 column chromatography. The \textsuperscript{35}S-labeled cyclin D0 was produced in vitro in rabbit reticulocyte lysate. Digestion of \textsuperscript{35}S-labeled cyclin D0 was performed for 60 min at room temperature with (●) or without (○) 26S proteasome. Samples were then separated on Sephadex G-50 column (1.0 × 19.0 cm) in 100 mM Tris-HCl, 5 mM MgCl\textsubscript{2}, pH 7.6. Fractions of 0.5 ml were collected. Arrows indicate the eluted positions of molecular weight standards as follows: 1, bovine serum albumin; 2, myoglobin; 3, ubiquitin; 4, total column volume (Vt). (B) SDS-PAGE analysis of gel chromatography fractions. Sephadex G-50 column chromatography fractions from 26S proteasome-treated cyclin D0 and untreated (Control) were separated by SDS-PAGE (15% gel) followed by autoradiography on Imaging plates (Fuji Film). The positions of the digested cyclin B is indicated by an asterisk, and the positions of NH\textsubscript{2}-terminal portion of cyclin B is indicated by an arrowhead.
NH₂-terminal restricted portion by the 26S proteasome (K57 in goldfish cyclin B) changes the cyclin structure available for further chemical modifications including ubiquitination, which leads to the complete destruction of the cyclin at metaphase/anaphase transition. Since cyclin D₅₇ was destroyed, whereas cyclin D₆₈ was not, the lysine residues between amino acids 58 to 68 constitute the most likely ubiquitination site, and cutting by the 26S proteasome at K57 might be necessary to expose them to ubiquitinating enzymes. This notion should be verified by investigating the difference in the three-dimensional structure of cyclins D₀, D₀K₅₇R, D₄₁, and D₆₈.

Cyclin B is absent in immature (prophase I-arrested) goldfish oocytes. Cyclin B is de novo synthesized during oocyte maturation and forms an MPF complex with extant cdc2, which drives the prophase I-arrested oocytes to metaphase II (mature oocytes; Hirai et al., 1992; Katsu et al., 1993; Yamashita et al., 1995). As shown in this study however, the 26S proteasome purified from immature goldfish oocytes can digest cyclin B. If the initial cleave of cyclin B by the 26S proteasome triggers cyclin destruction as proposed in this study, the question remains why cyclin B is stable during oocyte maturation and in mature oocytes but destroyed upon egg activation. Based on the results obtained from clam oocytes, Sudakin et al. (1995) have suggested that the initiation of cyclin degradation is triggered by ubiquitination caused by the activation of cyclin-selective ubiquitin ligase near the end of M-phase, which targets cyclin B for destruction by the 26S proteasome that is constitutively active during the cell cycle. Contrary to this, we found that the 26S proteasome purified from mature goldfish oocytes cannot digest cyclin B.

Figure 10. Digestion and degradation of in vitro translated cyclin B. The ³⁵S-labeled cyclins D₀, D₀K₅₇R, D₄₁, and D₆₈ were produced in vitro in rabbit reticulocyte lysate. After the translation of each cyclin, the lysate was incubated in the presence of 100 µg/ml of cycloheximide at room temperature under the indicated conditions. The ³⁵S-labeled proteins were resolved by SDS-PAGE followed by autoradiography on Imaging plates (Fuji Film). The position of the digested cyclin B is indicated by an asterisk. (A) Digestion of full length cyclin B by purified 20S and 26S proteasomes. The reticulocyte lysate containing cyclin Δ₀ was incubated with 60 µg/ml of proteasomes. (B) Digestion of full length, point mutated, and NH₂-terminal truncated cyclin Bs by purified 26S proteasome. The reticulocyte lysate containing cyclin D₀, D₀K₅₇R, D₄₁, or D₆₈ was incubated in the absence (−) or presence (+) of 60 µg/ml of the 26S proteasome for 60 min. (C) Degradation of cyclin B in Xenopus egg extracts. One ninetieth of the lysate containing cyclin D₀, D₀K₅₇R, D₄₁, or D₆₈ was added to the Xenopus egg extracts, and its degradation was induced by 0.4 mM Ca²⁺. At the indicated times, the reaction was terminated by adding SDS sample buffer. (D) The same sample as in C. Cyclin contents were quantified using an image analyzer (BAS2000; Fuji Film).
and that at least two subunits in 26S proteasomes from immature and mature oocytes differ (Tokumoto, T., Horiguchi, R., Nagahama, Y., unpublished results). These findings suggest that some inhibitory mechanisms preventing cyclin B degradation proceed on the proteasome itself at least during metaphase II arrest. The amount of the proteasome in egg cytosol also changes dramatically during oocyte maturation and egg activation; the lowest level is in mature metaphase II-arrested oocytes, and there is a transient increase between the first and second meiotic cell cycles and upon egg activation in goldfish (Tokumoto et al., 1993a). Further studies should reveal how the subunit composition of proteasomes and their contents during oocyte maturation and egg activation are involved in controlling the cell cycle by regulating cyclin stability.

Stewart et al. (1994) have shown that binding with cdc2 is necessary for the degradation of Xenopus cyclins A and B2 but not for that of cyclin B1. This implies that the mechanisms of cyclin degradation vary according to the types of cyclins. Since goldfish cyclin B exhibits higher homology to Xenopus cyclin B1 (66%) than to cyclin B2 (50%), the mechanism of cyclin degradation by the 26S proteasome proposed in this study may be specific to cyclin B1.

Our present data suggest that the NH2-terminal restricted cleavage of cyclin B by 26S proteasome allows cyclin to be ubiquitinated. Compared with the idea that the initiation of cyclin destruction is dependent on the activity of ubiquitin ligase, irrespective of 26S proteasome activity (Sudakin et al., 1995), we propose that cyclin destruction is primarily controlled by the activity of 26S proteasome.

Ubiquitination of cyclins has been well studied genetically and biochemically (for review see Murray, 1995). A cyclin-specific ubiquitin ligase complex, the cyclosome, or APC complex, has been characterized in clam and Xenopus, respectively (King et al., 1995; Sudakin et al., 1995). These ubiquitin ligases (E3) catalyze ubiquitination using a specialized ubiquitin carrier protein (E2). Among the multiple species of E2s, UBC9 is required for cell cycle progression in late G2 or early M-phase (Seufert et al., 1995). UBC4 protein can ubiquitinate cyclins in Xenopus egg extracts (King et al., 1995). Recently, a novel cyclin-selective UBC family member, E2-C, was reported which can ubiquitinate cyclin B(13-91)/protein A fusion protein in cyclosome-dependent manner (Aristarkhov et al., 1996). These reports have shown that destruction box mutants cannot be ubiquitinated or degraded after extract activation, suggesting that the destruction box is a recognition sequence for the ubiquitinating system. However, we have shown that mutants that lack the proteasome cleavage site cannot be degraded after extract activation. The relative importance of these two processes is unclear because of the discrepancy between in vivo and in vitro results. In vitro, proteasome cleavage and ubiquitination of cyclin seem independent of each other. Destruction box-dependent ubiquitination of cyclin by purified proteins does not depend on the previous proteasome cleavage of cyclin B, and cyclin cleavage by purified proteasome does not depend on previous ubiquitination of cyclin B. But, since mutations that block in vitro ubiquitination and mutations that block proteasome cleavage also block destruction in extracts, the simple conclusion is that the ubiquitination

and proteasome cleavage are both necessary for cyclin destruction in extracts. There are, however, no experimental data available at present that provide information on the relative order of these two steps in vivo. Therefore, there are three possible in vivo scenarios. (a) Proteasome cleavage precedes ubiquitination to expose an NH2-terminal lysine that is a good substrate for ubiquitination. An extreme view would be that destruction box-dependent ubiquitination is an artifact that plays no role in vivo and that the purpose of the destruction box is solely to induce the initial ubiquitin-independent cleavage of cyclin. (b) Destruction box-dependent ubiquitination precedes proteasome cleavage to recruit cyclin to the proteasome by virtue of the proteasome’s polyubiquitin binding subunit. An extreme view would be that proteasome cleavage is not necessary in vitro, and mutants like K57R are having a direct effect on destruction box-dependent ubiquitination. (c) There is no required order of proteasome cleavage and destruction box-dependent ubiquitination, although both events would be necessary for efficient cyclin destruction, they could occur in either order. Further studies are necessary to understand the molecular mechanism of cyclin degradation, especially identification of the lysine residue that is destined to be ubiquitinated.

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