Okadaic Acid Induces Interphase to Mitotic-like Microtubule Dynamic Instability by Inactivating Rescue

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Abstract. We used high-resolution video microscopy to visualize microtubule dynamic instability in extracts of interphase sea urchin eggs and to analyze the changes that occur upon addition of 0.8–2.5 μM okadaic acid, an inhibitor of phosphatase 1 and 2A (PP1, PP2a) (Bialojan, D., and A. Takai. 1988. Biochem. J. 256:283–290). Microtubule plus-ends in these extracts oscillated between the elongation and shortening phases of dynamic instability at frequencies typical for interphase cells. Switching from elongation to shortening (catastrophe) was frequent, but microtubules persisted and grew long because of frequent switching back to elongation (rescue). Addition of okadaic acid to the extract induced rapid (<5 min) conversion to short, dynamic microtubules typical of mitosis. The frequency of catastrophe doubled and the velocities of elongation and shortening increased slightly; however, the major change was an elimination of rescue. Thus, modulation of the rescue frequency by phosphorylation-dependent mechanisms may be a major regulatory pathway for selectively controlling microtubule dynamics without dramatically changing velocities of microtubule elongation and shortening.

The conversion from the long, persistent microtubules of interphase into the short, dynamic microtubules of mitosis is controlled by phosphorylation initiated by the activation of maturation promoting factor (MPF) (p34cdc2 kinase) (4, 22, 45) and a cascade of downstream kinases, including MAP kinase (15, 16). Addition of either p34cdc2 (22, 45) or MAP kinase (16) to interphase cytoplasm or cytoplasmic extracts of Xenopus embryos produces conversion to short dynamic microtubules. Conversely, addition of drugs like 6-dimethylaminopurine, an inhibitor of p34cdc2 kinase, induces long microtubules when added to mitotic cells (30).

In both interphase and mitotic animal cells, the plus ends of microtubules exhibit dynamic instability, abruptly switching between persistent, constant velocity phases of elongation and shortening (7, 17, 34, 35). Microtubule dynamics and lengths depend on the switching frequencies, catastrophe and rescue, as well as on the velocities of elongation and shortening. Catastrophe and rescue frequencies are difficult to measure accurately because of their stochastic nature, and data must be obtained from measurements of individual microtubules (37, 47). There is now some information on how the parameters of microtubule dynamic instability change between interphase and mitosis (7, 17), and upon activation of p34cdc2 kinase in interphase cytoplasm with cyclin B (4). Okadaic acid inhibits the activity of the serine, threonine protein phosphatases 1 and 2a (PP1 and PP2a) (5, 9, 18), and recent studies have shown that treatment with okadaic acid promotes premature and prolonged mitosis in sea urchin eggs (28) and shortens microtubules in interphase tissue culture cells (44). In the experiments reported here, we measured the changes in microtubule dynamics in interphase extracts of sea urchin eggs when phosphorylation was enhanced by inhibiting PP1 and PP2a with okadaic acid.

Materials and Methods

Preparation of Extracts

Interphase extracts from Lytechinus pictus (Marinus, Long Beach, CA) eggs were made as previously described (Gliksman, N. R., and E. D. Salmon. 1987. J. Cell Biol. 109:30a), from unfertilized eggs, or eggs 15 min after fertilization. Gametes were isolated by intracoelomic injection of 0.56 M KCl. Sperm were collected dry and stored on ice for up to 5 h. Eggs were collected into and washed 3x with artificial sea water (Instant Ocean, Mentor, OH). Eggs were then dejellied by washing 2x in isotonic 19:1 (28 mM NaCl, 0.53 mM KCl, 5 mM Tris base, pH 8.0). Egg fertilization was carried out in sea water containing 3-amino-1,2,4-triazole (1 mM) and passed through a 110-μm mesh (Tetko Inc., Elmsford, NY) to remove the fertilization envelopes (36). Unfertilized eggs or eggs 15 min after fertilization were collected in isotonic HEMG buffer (180 mM Heps, pH 7.35, 10 mM EGTA, 10 mM MgCl₂, 0.5 M glycine, 1 mM ATP, 1 mM GTP, 1 mM DTT, and protease inhibitors: 0.3 mM phenyl methyl sulfonyl fluoride, 10 μg/ml p-toluene-sulfonyl-L-arginine methyl ester HCl, 40 μg/ml soybean trypsin inhibitor, 1 μg/ml pepstatin, 1 μg/ml benzamidine HCI, 10 μg/ml aprotonin, and 1 μg/ml leupeptin) and packed with a minimum volume of buffer (20–30% dilution of the cytosol) using a hand centrifuge. The eggs were homogenized and centrifuged at 50,000 g for 45 min at 3°C. The clear middle layer was collected with a 25-gauge needle/syringe and used fresh or frozen in liquid nitrogen and stored at −80°C. L. pictus egg extracts contained 15-19 μM tubulin (Western blot.
Measuring the Dynamic Instability of Microtubules

Microtubules were visualized by video-enhanced differential interference contrast (VE-DIC) microscopy at room temperature (23°C) as previously described (37, 47). Slide preparations were made by diluting the extract samples 4% with HEMG buffer (without protease inhibitors or DTT) containing 0.2 mM ATP, 0.1 mM GTP, 10 μM cytochalasin B, 0-2.5 μM okadaic acid, and salt extracted sea urchin axoneme seeds (47) or sperm centriole complexes (Sluder, G., and E. J. Miller, manuscript in preparation). A 4-μl aliquot of the sample was added to a slide cleaned with ethanol, covered with a biologically clean (23) coverslip (22 × 22 mm, thickness #1), and sealed with valap (a 1:1:1 mixture of Vaseline/lanolin/beeswax) to prevent flowing and drying of the chamber contents.

Perfusion chambers were made from 25 × 25-mm coverslips mounted over two parallel pieces of double-stick tape forming a 70-μm-high by 4-mm-wide channel (43). Perfusion was carried out by adding five chamber volumes of sample to one side of the chamber, and drawing the liquid through with filter paper on the other side. Control experiments showed that perfusion induced no changes in the parameters of dynamic instability of unfertilized interphase egg extract.

Microtubule assembly was videotaped and later analyzed using an IBM-compatible PC-based system for data measurement (47). The standard deviation measured for a stationary object was ± 0.1 μm for repetitive determinations.

Three strategies were used to measure microtubule lengths over time. Straight, nonmotile microtubules were measured by tracking the plus end in real-time or at fixed time intervals (between 0.033 and 2.0 s). Straight, motile microtubules were measured by tracking the two ends of the microtubule independently over the same time interval. The two data sets were combined, and the distance between the two ends was interpolated from the same starting time. Curved microtubules were measured at fixed time intervals (between 0.033 and 2.0 s) by adding up straight lines along the curved microtubule. With all three strategies, velocities of elongation and shortening were calculated from least squares regressions of distance versus time using a minimum of three data points. The catastrophe and rescue frequencies were calculated from the total number of catastrophe or rescue transitions observed, divided by the total length of time spent in either the assembly phase or the shortening phase, respectively (47).

Figure 1. Microtubule assembly in cytoplasmic extracts of *Lytechinus pictus* eggs. Plus-end microtubules assembled in untreated (A and B) or okadaic acid treated (C and D) interphase sea urchin egg extracts. Microtubules were nucleated from sperm tail axoneme seeds (A and C) or sperm centriole complexes (large arrows in B and D). Microtubule motors on the coverslip surface pulled the minus ends of microtubules (small arrows in B and D) from the centriole complexes and translocated microtubules across the coverslip surface away from the nucleating centers. Note that contrast and apparent width of the microtubules and axonemes depend on orientation in VE-DIC. Bars, 5 μm.
Phosphatase Regulation of Microtubule Dynamics

Results and Discussion

We viewed the dynamic instability of individual microtubules in cytoplasmic extracts using high-resolution, VE-DIC light microscopy (37, 47). Concentrated interphase cytoplasmic extracts were made from unfertilized *Lytechinus pictus* eggs or eggs 15 min after fertilization. At pH 7.3, the natural pH of fertilized eggs, little spontaneous self-assembly occurred in the extracts. However, sperm tail axonemal fragments (Fig. 1A) and sperm centriole complexes (Fig. 1B) nucleated plus end (but not minus end) microtubule assembly as previously observed (12, 37, 38, 39). We used low concentrations of nucleation sites (1 or 2 per 35-μm video field) so that microtubule assembly did not significantly deplete the concentration of unpolymerized tubulin in our preparations.

Long, persistent microtubules typical of interphase cytoplasm (7, 34, 35) grew in the egg extracts at 23°C (Figs. 1, a and b, and 2, a and c). The rate of plus-end microtubule elongation was 9.1 ± 2.9 μm/min, while the rate of shortening was 12.3 ± 4.4 μm/min (Table I). Elongating ends grew for a mean duration of 53 s and a mean length of 8.0 μm before catastrophe and switching to shortening. Shortening occurred for only 28 s and 5.7 μm on average before rescue back to elongation. Nearly all shortening events were rescued, so that the interphase microtubules did not achieve steady-state lengths. Instead, microtubules persisted and continued to grow longer at a net rate of ~1.7 μm/min.

\[
\text{net growth rate} = \frac{[\text{elongation rate} \times \text{elongation duration}] - [\text{shortening rate} \times \text{shortening duration}]}{(\text{elongation duration + shortening duration})}
\]

Pauses in microtubule assembly, noted previously both in vivo (34) and in vitro (37, 47), represented only a minute fraction (6-7%) of the observed microtubule lifetimes.

Addition of 0.8-2.5 μM okadaic acid (the predicted concentration of PP1 and PP2a in the sea urchin egg) (9, 28) to the interphase extracts reduced microtubule lengths and lifetimes to the short steady-state values typical of mitosis (1, 33, 45) (Fig. 1C). This conversion in microtubule length and dynamics was not produced by major changes in either the velocities of elongation or shortening; there was little change in the velocity of elongation, and the velocity of shortening increased by <30% (Table I; Fig. 3). Microtubules were much shorter and more labile due to a twofold increase in catastrophe frequency, an elimination of pauses, and a complete inhibition of rescue (Table I; Fig. 3). Rescue was the parameter of dynamic instability most sensitive to okadaic acid. Rescue was undetectable within 5 min of okadaic acid addition and after a catastrophe, microtubules shortened all the way back to the nucleation center. Methyl okadaate and 1-nor okadaone are analogs of okadaic acid which are not phosphatase inhibitors (27). Neither analog inhibited microtubule rescue in our interphase extracts.

Perfusion chambers were also used to determine how quickly rescue was eliminated after okadaic acid treatment for microtubules initially grown to long lengths in untreated extract. We first grew long microtubules in interphase ex-

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Measuring the Microtubule Length Distribution

Microtubule length distributions were measured from the same video recordings used to measure microtubule dynamics using the programs described above. Microtubule lengths were measured at each new field of view for microtubules initially grown to long lengths in untreated extract. We first grew long microtubules in interphase extracts with okadaic acid-pretreated extract rapidly eliminated and pauses. (C) Perfusion of long microtubules grown in interphase elongation, rapid shortening, frequent catastrophes, but no rescues and pauses. Note the scale changes between A, B, and C.

**Figure 2.** Length histories of microtubules assembled in clarified sea urchin extracts. Different symbols represent different microtubules. (A) Microtubule assembly in the interphase extracts was dynamic with rapid elongation, rapid shortening, frequent catastrophes, frequent rescues, and infrequent pauses (P). (B) Microtubules assembled in okadaic acid–treated extracts displayed rapid elongation, rapid shortening, frequent catastrophes, but no rescues and pauses. (C) Perfusion of long microtubules grown in interphase extracts with okadaic acid–pretreated extract rapidly eliminated rescues and pauses. Note the scale changes between A, B, and C.

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The Simulation Program

Microtubule dynamic instability was simulated using a Monte Carlo computer program (see ref. 2, and Glikman, N. R., E. D. Salmon, and R. A. Walker. 1987. *J. Cell Biol.* 111:298a). The program was written in Turbo Pascal V. 5 (Borland International, Scotts Valley, CA) and run on IBM PC compatible computers. The program simulated a population of plus-end microtubules undergoing dynamic instability. The dynamic instability parameters were applied to each plus end in a Monte Carlo fashion at one-second time intervals for a period equivalent to 5 min. All histograms represent the average of 500 ends. All variables were made independent of tubulin concentration, and pauses were ignored for these simulations.
tricts from axonemes stuck to the inner coverslip surface of a perfusion chamber (Fig. 2 C). After 20 min, the microtubules tethered to the axonemes were perfused with extract which had been preincubated with 2.5 μM okadaic acid for 10 min at room temperature. Within 5 min of perfusion, all microtubule plus ends had switched to shortening. They shortened all the way back to the nucleation center without pause or rescue (Fig. 2 C).

Although microtubules in untreated interphase extracts had a net growth of 1.7 μm/min, after 5 min in okadaic acid there was no net growth of microtubules because of the absence of rescue. Instead, an average steady-state length was achieved (Fig. 4). As shown in Fig. 4, the length distributions of microtubules measured in the untreated and okadaic acid–treated extracts were similar to the microtubule length distributions predicted by a Monte Carlo computer simulation of microtubule assembly dynamics, using the mean values for the dynamic instability parameters listed in Table I. In the absence of rescue, the microtubule steady-state length distribution was determined only by the growth velocity and catastrophe frequency. We have neglected the role of microtubule severing activity in this analysis since the severing reported in mitotic Xenopus extracts (43) was not seen in our extracts before or after okadaic acid treatment.

Nucleation in the cytoplasmic extracts depended both on the type of nucleation center and the treatment history. When axonemes and sperm centriole complexes were added directly to okadaic acid–treated extracts, nucleation from axonemes was infrequent in comparison with the frequency in untreated extracts, but nucleation from the sperm centriole complexes remained robust (Fig. 1, B and D). This was not surprising since previous studies have shown that axonemal but not centrosomal nucleation is inhibited in meiotic or mitotic cytoplasm in vivo (20). A surprise in the perfusion experiments with okadaic acid–treated extract was the finding that microtubule renucleation occurred almost immediately and was not lost after perfusion (Fig. 2 B). Since microtubules were pregrown from the plus ends of axonemes in our perfusion studies, this result suggests that nucleation material in the cytoplasm is recruited to assembled microtubules and transported along the microtubules to the nucleation center (46).

Okadaic acid treatment increased the phosphorylation of a number of extract proteins (seen via SDS-PAGE and autoradiography), and doubled the TCA-precipitable 32P counts, but no phosphorylation of tubulin was detected (data not shown). Comparing the changes in microtubule dynamic instability induced by okadaic acid (Table I and Fig. 3) to those reported to occur upon cyclin B activation of p34<sup>cdc2</sup> kinase in interphase Xenopus extracts (4), we note that neither study found a decrease in elongation or shortening velocities in treated cytoplasm, a result consistent with available in vivo data (7, 17). Our interpretation of the okadaic acid results, along with the microtubule assembly data from Xenopus egg extracts treated with cyclin B activated p34<sup>cdc2</sup> (4) or MAP kinase (14), is that there is a catastrophe factor which is acti-

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**Table I. Parameters of Microtubule Dynamic Instability**

| Extract conc. | OA Elongation rate ± SD (μm/min) (N) | Shortening rate ± SD (μm/min) (N) | Catastrophe rate* (Total time) (S<sup>-1</sup>) | Rescue rate* (Total time) (S<sup>-1</sup>) |
|---------------|-------------------------------------|----------------------------------|-----------------------------------------------|------------------------------------------|
|               | 1                                   | 1.25                             | 10.1 ± 3.1 (15)                               | 14.6 ± 2.9 (14)                         | 0.051 (215 s)                           | 0.0124 (244 s)                         |
|               | 2                                   | 0                                | 10.8 ± 2.6 (41)                               | 13.8 ± 3.5 (40)                         | 0.017 (1152 s)                         | 0.023 (986 s)                          |
|               | 3                                   | 2.5                              | 11.3 ± 3.3 (22)                               | 18.4 ± 4.3 (23)                         | 0.026 (352 s)                          | 0.00 (284 s)                           |
|               | 4                                   | 0                                | 7.8 ± 3.0 (49)                                | 12.7 ± 4.2 (27)                         | 0.016 (1698 s)                        | 0.034 (790 s)                          |
| Mean†         | 6                                   | 2.5                              | 9.1 ± 2.9 (142)                               | 12.3 ± 4.4 (99)                         | 0.019 (3937 s)                        | 0.036 (2148 s)                         |
| Mean‡         | 1.0-2.5                             | 2.5                              | 9.2 ± 3.0 (84)                                | 15.3 ± 4.9 (93)                         | 0.038 (1600 s)                        | <0.0024 (1670 s)                      |

The parameters of microtubule dynamic instability in sea urchin extracts. Extracts and length measurements were made as described in Figs. 1 and 2, and numbers were recorded from the first 20–25 min after warming the sample to 23°C. No differences in microtubule dynamics were noted between microtubules assembling on the glass surface (attached via microtubule motors) or assembling above the glass surface in solution. No differences in microtubule dynamics were noted between microtubules assembling off of axoneme seeds, or sperm centriole complexes, or between microtubules in simple slide coverslip preparations and in the perfusion chambers. Total time the duration of either elongation or shortening measured for each extract; N is the number of periods of either elongation or shortening measured for each extract.

* The catastrophe and rescue rates were calculated from the number of transitions observed divided by the total time observed in elongation or shortening.

† Combined data from all four extracts.

‡ Of the few rescue events seen, all occurred within the first 2 minutes of observation.

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**Figure 3. Okadaic acid–induced changes in the parameters of microtubule dynamic instability.** The catastrophe frequency doubled, the shortening rate increased slightly, and the rescue frequency decreased substantially when okadaic acid was added to the sea urchin egg extracts. The data was taken from the mean values in Table I.

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...vated by phosphorylation and a rescue factor which is inacti-
phorylate the factors (13, 14, 19, 25, 30, 32, 42). There are
and rescue factors in two ways: by directly blocking dephos-
are not yet known.

bles of the okadaic acid-treated extracts remained at a steady-state
length (30--40 #m); nevertheless, computer simulations predicted
mitotic-like microtubules after okadaic acid treatment. Both sets of
microtubules typical of untreated extracts, and the short, dynamic,
microtubule length distributions after 5 min of as-
puter simulation from mean values of the microtubule dynamic instabil-
Figure 4. The microtubule length distributions after 5 min of as-
the extracts as well as microtubule lengths predicted by computer
mulation parameters in Table I. The average microtubule lengths mea-
for the untreated and okadaic acid--treated samples, respect-
respectively. The computer simulations demonstrated that the
model the assembly of long interphase microtu-
the untreated and okadaic acid--treated extracts, and the short, dynamic,
microtubules after okadaic acid treatment. Both sets of
microtubules represent 5 min of assembly from 0 length. The
parameters for the untreated and okadaic acid--treated extract
predicted average microtubule lengths of 18.9 and 4.1 #m after 5
min of simulated assembly. By 15 min, microtubule lengths could
not be measured in the interphase extracts because of the video field
length (30--40 #m); nevertheless, computer simulations predicted
an average microtubule length of 28 #m. After 15 min the microtu-
bules of the okadaic acid--treated extracts remained at a steady-state
average length of 4.1 #m.

vated by phosphorylation and a rescue factor which is inactiv-
phorylation. There is some evidence that these are
different molecular complexes (37), but their identities
are not yet known.

Okadaic acid, by blocking PPI and PP2a activity, can en-
hance the phosphorylation of these presumptive catastrophe
and rescue factors in two ways: by directly blocking dephos-
phorylation of the factors in the presence of a basal level of
kinase activity and by activating the kinases which phos-
phorylate the factors (13, 14, 19, 25, 30, 32, 42). There are
a number of kinases which may be responsible for phosphor-
ylation of the catastrophe and rescue factors. Staurosporine
(100 #M), an inhibitor of protein kinase C, cAMP kinase,
cGMP kinase, pp60--Src, and myosin light chain kinase (24,
40), did not prevent the okadaic acid--induced loss of rescue
(data not shown). Although p34<sub>cdc2</sub> kinase is activated in
some cell types by okadaic acid (13, 30, 32), in the sea urchin
egg extracts p34<sub>cdc2</sub> kinase was stimulated only slightly (data
not shown). This result is not surprising, since unfertilized
eggs and eggs shortly after fertilization contain little of the
cyclin A or B needed to activate p34<sub>cdc2</sub> kinase (11). Okadaic
acid has been shown to activate MAP kinase in some cell
types (14, 19, 25, 42). In addition, Gotoh et al. (15, 16) have
reported that MAP kinase is activated downstream of p34<sub>cdc2</sub>
kine, producing short mitotic-like microtubules without
activating the p34<sub>cdc2</sub> kinase. Based on this information, we
tested whether okadaic acid would induce MAP kinase ac-
tivation in our sea urchin extracts by measuring the phosphor-
ylation of exogenous MAP2 and MBP, both known substrates
for MAP kinase (8, 14, 15, 19, 25, 31, 42). We found that
a MAP2 and MBP kinase activity increased two- to threefold
in the presence of okadaic acid (data not shown), similar to
the increase in this activity at mitosis reported for sea ur-
chins embryos (29). Our data support a role of MAP kinase
in the regulation of catastrophe and rescue frequency, but do
not preclude the contributions of other unknown kinases.

Cyclin B activated p34<sub>cdc2</sub> kinase (4) and okadaic acid in-
hibition of PPI and PP2a activity both activate MAP kinase
and produce similar redistributions of microtubule lengths.
For example, compare the microtubule length distributions
of Fig. 4 with Fig. 2 of reference 45, Fig. 5 of reference 21,
Fig. 3 of reference 16, Fig. 7 a of reference 1 and Fig. 6 of
reference 6. As a result, we expected them to change the
catastrophe and rescue frequencies in the same way. Belmont
et al. (4) reported that the short dynamic microtubules in cy-
clin B--activated extracts were produced by a six- to eightfold
increase in catastrophe frequency, with little decrease in res-
cue frequency. In contrast, we found that okadaic acid treat-
ment produces only a twofold activation of catastrophe, but
a total elimination of rescue. Belmont et al. (4) were uncer-
tain of their rescue measurements in the activated extracts
because of the high density of short microtubules near the
centrosomes. It is possible that the rescues they scored in
their activated extracts were not true rescues of a single end,
but rather elongating ends passing by adjacent shortening
ends. Similar technical considerations can not be used to ex-
plain the different changes in catastrophe frequency pro-
duced by cyclin B activation of p34<sub>cdc2</sub> activity and okadaic
acid; these differences remain an interesting and important
puzzle.

It now seems apparent that catastrophe and rescue fre-
quencies are key phosphorylation-dependent regulators of
cytoplasmic microtubule lengths and lifetimes, not the ve-
locity of elongation or shortening of individual microtu-
bules. The results of Belmont et al. (4) demonstrated that in-
creases in catastrophe frequency can cause conversion of
long, persistent, interphase microtubules into short, labile,
mitotic-like microtubules. Our experiments have shown that
elimination of rescue produces the same result. Changes in
phosphorylation make substantial changes in the frequencies
of catastrophe and rescue without major changes in the ve-
locities of elongation and shortening. These findings do not
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