A β-Tubulin Mutation Selectively Uncouples Nuclear Division and Cytokinesis in *Tetrahymena thermophila*

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Received 3 December 2003/Accepted 23 June 2004

The ciliated protozoan *Tetrahymena thermophila* contains two distinct nuclei within a single cell—the mitotic micronucleus and the amitotic macronucleus. Although microtubules are required for proper division of both nuclei, macronuclear chromosomes lack centromeres and the role of microtubules in macronuclear division has not been established. Here we describe nuclear division defects in cells expressing a mutant β-tubulin allele that confers hypersensitivity to the microtubule-stabilizing drug paclitaxel. Macronuclear division is profoundly affected by the *btu1-1* (K350M) mutation, producing cells with widely variable DNA contents, including cells that lack macronuclei entirely. Protein expressed by the *btu1-1* allele is dominant over wild-type protein expressed by the *BTU2* locus. Normal macronuclear division is restored when the *btu1-1* allele is inactivated by targeted disruption or expressed as a truncated protein. Immunofluorescence studies reveal elongated microtubular structures that surround macronuclei that fail to migrate to the cleavage furrows. In contrast, other cytoplasmic microtubule-dependent processes, such as cytokinesis, cortical patterning, and oral apparatus assembly, appear to be unaffected in the mutant. Macronuclear division is also perturbed in the K350M mutant, producing nuclei with elongated early-anaphase spindle configurations that persist well after the initiation of cytokinesis. The K350M mutation affects tubulin dynamics, as the macronuclear division defect is exacerbated by three treatments that promote microtubule polymerization: (i) elevated temperatures, (ii) sublethal concentrations of paclitaxel, and (iii) high concentrations of dimethyl sulfoxide. Inhibition of phosphatidylinositol 3-kinase (PI 3-kinase) with 3-methyladenine or wortmannin also induces amacronucleate cell formation in a *btu1-1*-dependent manner. Conversely, the myosin light chain kinase inhibitor ML-7 has no effect on nuclear division in the *btu1-1* mutant strain. These findings provide new insights into microtubule dynamics and link the evolutionarily conserved PI 3-kinase signaling pathway to nuclear migration and/or division in *Tetrahymena.*

Microtubules are multifunctional cellular structures that mediate intracellular transport, cell motility, chromosome segregation, and cell division (40, 51). These polymers consist of noncovalently linked heterodimers of α- and β-tubulin that are decorated with microtubule-associated proteins that specify different biochemical and cellular properties (reviewed in reference 35). A critical event in chromosome transmission is the proper division and segregation of daughter nuclei. Astral microtubules that radiate from the spindle pole toward the cell cortex direct the migration of the mitotic nuclei prior to karyokinesis (reviewed in reference 32). This process is regulated by the concerted action of microtubule-dependent motors (kinetins and dyneins) and microtubule growth and shrinkage at the plus end. Actomyosin filaments generate the contractile ring filament at the cleavage furrow, and microtubules and septins complete cytokinesis (reference 43 and references therein).

The ciliated protozoan *Tetrahymena thermophila* has been used extensively as a model for studying microtubule-mediated cellular processes and deciphering the role of posttranslational modification of tubulin subunits (45). This research has been facilitated by the presence of a single α-tubulin gene (*ATU1*) and two β-tubulin genes (*BTU1* and *BTU2*) that encode identical proteins (19, 31). DNA transformation methodologies allow for the functional analysis of these genes by targeted homologous recombination. Cytological and biochemical studies have thus far identified at least 17 distinct types of microtubules with different subcellular localization patterns and/or associated proteins (16).

The role of microtubules in *Tetrahymena* nuclear division is likely to be complex, as *Tetrahymena* cells harbor two genetically related, but functionally distinct, nuclei—the germ line macronucleus and the somatic macronucleus (reviewed in reference 25). The transcriptionally silent diploid macronucleus serves as the reservoir of genetic information transmitted during conjugation and is thus dispensable for vegetative growth. The transcriptionally active polyploid macronucleus confers the cellular phenotypes during vegetative division and development. Microtubules play an essential role in mitotic and meiotic chromosome segregation in the diploid macronucleus. Furthermore, a meshlike microtubule network that forms at the conjugation junction facilitates the reciprocal exchange of pronuclei between mating cells (36).
The contribution of microtubules to macronuclear division is less clear. Although intramacronuclear microtubules appear prior to karyokinesis, they are disordered (23). Since macronuclear chromosomes lack centromeres, sister chromatids segregate randomly (9) and spindle pole bodies do not form. Instead, a nonconventional yet ordered array of cytoplasmic microtubules assembles perpendicular to the division plane of the replicated macronucleus. The temporal reorganization of cytoplasmic microtubules suggests their involvement in macronuclear division. In support of this idea, abnormal macronuclear division can be induced with microtubule-destabilizing drugs such as colchicine (23, 44). The blockage of cell division in Tetrahymena by disruption of the kinesin-II gene (3) or by mutations that eliminate β-tubulin polyglycation (45, 51) also implicates microtubules in cytokinesis.

Single amino acid substitutions in tubulin genes can dramatically alter microtubule stability. For example, the polymerization and stability of Chlamydomonas reinhardtii microtubules are enhanced by the β-tubulin K350M mutation, which imparts resistance to the microtubule-destabilizing drug vinblastine. This same mutation confers hypersensitivity to the microtubule-stabilizing drug paclitaxel (Taxol) (42). Similar drug sensitivities are observed in connection with the T. thermophila btu1-1 (K350M) allele (18).

The BTU1 locus has been exploited as a target site for introducing native or foreign genes that do not impart a selectable growth advantage on their own. Targeted disruption of the btu1-1 (K350M) allele in strain CU522 confers resistance to the microtubule-stabilizing drug paclitaxel (17), making positive selection for transformants possible. In the course of experiments using the btu1-1 locus for gene targeting, we uncovered cytogenetic evidence that microtubules are required for macronuclear division in vegetatively dividing Tetrahymena. We show that the btu1-1 (K350M) allele acts in a dominant-negative manner to disrupt microtubule dynamics and decrease the fidelity of macronuclear transmission to daughter cells. We also provide in vivo evidence for the involvement of the signaling protein phosphatidylinositol 3-kinase (PI 3-kinase) in microtubule-mediated macronuclear division.

MATERIALS AND METHODS

General methods. Molecular techniques were performed as described elsewhere (1). T. thermophila genomic DNA was isolated by detergent lysis procedures as previously described (53). Total RNA was isolated from vegetative and conjugating cells by using RNeasy total RNA kits (QIAGEN, Valencia, Calif.) as previously described (53). Total RNA was isolated from vegetative and conjugating cells by using RNeasy total RNA kits (QIAGEN, Valencia, Calif.).

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(b) (K350M) allele (18).

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Tetrahymena strains and growth conditions. T. thermophila strains CU522, CU725, and CU727 (Jacek Guertig, University of Georgia) contain the mutant β-tubulin allele btu1-1 (K350M), which confers hypersensitivity to paclitaxel. CU428 and SB1969 are wild-type strains, and TC219 is a spontaneous paclitaxel-resistant mutant CU522 revertant. All strains were grown in 2% PPYS in 50% ethanol; Sigma Chemicals) inhibited cell growth by 50% at the final concentrations used (1 μM for paclitaxel and 5 μM for vinblastine). When used alone to study amacronucleate cell formation, dimethyl sulfoxide (DMSO) was added to a final concentration of 2%. Concentrations of 1.0% or less had no effect on amacronucleate cell formation. The PI 3-kinase inhibitor wortmannin (250 μM stock in 100% DMSO; Calbiochem) was diluted to a final concentration of 250 nM (the working concentration range in mammalian cell studies is 10 to 250 nM). The PI 3-kinase inhibitor 3-methyladenine (3-MA; 1 μM stock in 100% DMSO; Sigma) was diluted to a final concentration of 10 nM (the working concentration range in typical mammalian cell studies is 10 to 250 nM). The PI 3-kinase inhibitor 3-methyladenine (3-MA; 1 μM stock in 100% DMSO; Sigma) was diluted to a final concentration of 10 nM (the working concentration range in typical mammalian cell studies is 10 to 250 nM).

Transformation of T. thermophila. Strain CU522 was grown at 30°C in 100-ml cultures (2% PPYS) to a density of 1 × 105 to 3 × 105 cells/ml and starved in an equal volume of 10 mM Tris-HCl (pH 7.5) for 14 to 21 h. Cells were gently pelleted and resuspended in 1 to 3 ml of 10 mM HEPEs (pH 7.5) to a density of approximately 106 cells/ml. The concentrated cells were transferred to a 100-mm-diameter petri dish that contained a sterile Whatman 114 filter paper pre-soaked with 2 ml of 10 mM HEPEs (pH 7.5). The petri dish was placed in the Bio-Rad gene gun sample chamber, and gold particles (1.0 μm in diameter) coated with approximately 1.0 μg of a neomycin resistance cassette targeted to the btu1-1 (K350M) locus (18) were introduced into cells biologically as per the instructions of the manufacturer (Bio-Rad). After transformation, cells were incubated in 50 ml of warmed 2% PPYS at 30°C for 2 h. Paclitaxel was then added to the culture to a final concentration of 20 μM. Cells were distributed into well plates (100-μl aliquots) and incubated in a dark humidity chamber at 30°C for 4 to 7 days. Colony cell lines were screened by Southern blot analysis for disruption of the btu1-1 (K350M) locus (29). Continuous selection in 20 to 40 μM paclitaxel was maintained until 100% phenotypic assortment relative to the disrupted allele was achieved, as confirmed by RT-PCR analysis of the targeted btu1-1 (K350M) allele.

Cell synchronization and flow cytometry. Cell cycle synchronization was achieved by slight modification of a stationary-phase synchronization protocol described previously (34). Brieﬂy, saturated cultures were starved for 8 h prior to dilution in growth media. Cells were stained with propidium iodide and monitored for synchronization by flow cytometry as previously described (7).

Feulgen staining and cytophotometry. Cells were grown to a density of 1 × 105 to 2 × 105 cells/ml in 3 ml of growth media. Cells were transferred to 20 to 40 μM paclitaxel or 5 μM vinblastine (Calbiochem) and incubated for 1 to 3 h. Cells were then processed as previously described (28). A single drop of VectraShield with 6-diamidino-2-phenylindole (DAPI; Vector Labs) was added to the cells prior to transfer to 20 μl of fixed cell suspension on microscope slides. To obtain vegetative dividing cells for DNA flow cytometry analysis, a magniﬁcation of ×320 by using a 40× oil immersion lens objective, a 1.6-optimizer setting, and a 5× ocular lens objective or with a Bio-Rad MRC1000/1024 laser confocal microscope (University of Minnesota Imaging Center). Micrographs were recorded and images were overlaid using a SPOT camera and imaging software.

Acridine orange and Hoechst 33342 staining. Approximately 0.1 ml of cultured cells was doubly stained with Acridine orange (0.001% acridine orange and 5 μg of Hoechst 33342/ml). Stained cells were immediately observed by fluorescence microscopy as described above.

Cell division and nuclear segregation analysis. Cells were fixed and stained with DAPI and visualized as previously described (28). Micrographs were recorded digitally using a SPOT camera and imaging software to create the negative images. Six micrographs of each strain were captured, and the percentage of cells lacking macronuclei was recorded. Each strain was measured in three independent experiments when grown at 25, 30, or 35°C.
RESULTS

Targeting of the btu1-1 (K350M) allele reveals a role for β-tubulin in macronuclear division. Drugs such as vinblastine and paclitaxel alter normal microtubule dynamics and are commonly used to block mitosis. A previous study of wild-type strains demonstrated that microtubule-inhibitory drugs interfere with amitotic macronuclear division in *T. thermophila* (20). In an effort to better understand the role of microtubules in both mitotic and amitotic nuclear division, we examined the fate of micro- and macronuclei in strains expressing either the wild-type (CU428) or paclitaxel-hypersensitive (CU522) β-tubulin allele *btu1-1* (K350M). In contrast to that in wild-type cells, macronuclear morphology in CU522 (*btu1-1*) cultures grown in the absence of paclitaxel showed significant variability, the most obvious differences being cells that had diminutive macronuclei or that lacked macronuclei entirely (Fig. 1A and B). Inactivation of the macronuclear *btu1-1* locus by targeted disruption with several unrelated transgenes restored wild-type macronuclear morphology and division (data not shown; see below), suggesting that the *btu1-1* mutation is responsible for the macronuclear division defect. Other *btu1-1* strains (CU725 and CU727) were similarly defective in macronuclear division.

Exacerbation of the *btu1-1* (K350M) amacronucleate phenotype at elevated temperatures. The *btu1-1* allele confers hypersensitivity to the microtubule-stabilizing drug paclitaxel. Since microtubules are also stabilized by elevated temperatures, we tested the effect of temperature on amacronucleate cell formation. Under standard culture conditions (30°C), ~6% of CU522 cells lacked macronuclei (Fig. 1B). The frequency of amacronucleate cell formation increased to ~12% when growing cultures were maintained at 35°C. In contrast, almost no amacronucleate cells were detected at 25°C. Irrespective of growth temperature, the amacronucleate phenotype was not detected in wild-type strain CU428, indicating that the temperature-sensitive effect is linked to the *btu1-1* allele.

Cytophotometric analysis revealed that the macronuclear DNA contents of cells expressing the *btu1-1* allele varied considerably compared to those of wild-type cells (Fig. 1C). When normalized to the postmeiotic 4C micronucleus, the average macronuclear DNA content of CU522 cells cultured at 35°C was 117C (standard error, ±66C; range, 31C to 546C). In contrast, the macronuclear DNA contents of wild-type CU428 cells grown under the same conditions were much more moderate (mean, 93C; standard error, ±13C; range, 64C to 132C), with values similar to those reported previously for wild-type *T. thermophila* strains (11). The wide range in macronuclear DNA contents in CU522 populations indicates that asymmetric macronuclear division occurs frequently. These events not only generate amacronucleate cells but also produce the reciprocal product of asymmetric macronuclear division—cells with an overabundance of macronuclear DNA.

The microtubule-stabilizing drugs paclitaxel and DMSO promote amacronucleate cell formation. The dominant *btu1-1* allele causes hypersensitivity to the microtubule-stabilizing agent paclitaxel. Since the nuclear division defect in the mutant is exacerbated at elevated temperatures, it seemed likely that microtubule dynamics were being affected at the higher temperatures. To test this hypothesis, we treated cells with increasing concentrations of drugs that stabilize (paclitaxel) or destabilize (vinblastine) microtubules. Titration curves were first generated for paclitaxel and vinblastine to identify drug concentrations that slowed down, but did not completely inhibit, cell growth (Fig. 2A and B). Drug concentrations that reduced the growth rate by ~50% were then tested for their effects on macronuclear division. Paclitaxel (1 μM) caused a sixfold increase in the frequency of amacronucleate CU522 cells (20% of the treated population), but 5 μM vinblastine had no effect (Fig. 2C).

DMSO has been shown to inhibit the polymerization of astral microtubules in *Xenopus laevis* egg extracts (41, 47). Two-percent DMSO produced a dramatic effect on amacro-
nucleate cell formation: more than one-third of the CU522 cell population lacked a macronucleus (Fig. 2C). The titration curve for DMSO was steep, as 1% DMSO had virtually no effect on amacronucleate cell formation and 4% DMSO was toxic (data not shown). Remarkably, cells with two macronuclei were not detected in all populations treated with regimens that enhanced amacronucleate cell formation (exposure to elevated temperatures, paclitaxel, and DMSO). Similar to elevated temperatures, microtubule-stabilizing drugs had no effect on macronuclear division in cells expressing just the full-length wild-type H9252-tubulin gene (Fig. 2D shows data for TC219, a CU522 revertant; see below). The allele-specific effect of DMSO and paclitaxel suggests that both drugs target Tetrahymena microtubules in vivo.

Paclitaxel resistance correlates with the loss of the amacronucleate phenotype. As noted above, disruption of the btu1-1 (K350M) allele by insertion of a foreign gene confers paclitaxel resistance. During the selection for targeted btu1-1 transformants, nontransformed clonal lines resistant to paclitaxel were also isolated. RT-PCR analysis of mRNA from one of these spontaneous mutant strains (TC219) revealed wild-type levels of btu1-1 mRNA expression (Fig. 3A). TC219 and an independently isolated btu1-1::Neo transformant failed to produce the amacronucleate phenotype at all tested growth temperatures (Fig. 3B). Sequence analysis of btu1-1 cDNA from TC219 revealed a 4-bp deletion at the H9252-tubulin codon corresponding to S339, resulting in missense translation of btu1-1 until a nonsense codon was encountered 76 codons beyond the S339 codon (Fig. 3C). Thus, the loss of the amacronucleate phenotype when the btu1-1 allele is either disrupted or mutated demonstrates that expression of the H9252-tubulin K350M mutation is responsible for the macronuclear division defect. Since the CU522 strain also expresses wild-type H9252-tubulin from the BTU2 locus, the mutant btu1-1(K350M) allele must be dominant over the wild type.

btu1-1 (K350M) nuclear phenotypes. CU522 and TC219 strains were examined by conventional and confocal microscopy to assess the organization of microtubules at various stages of cell division. A panel of confocal images clearly illustrates that the amacronucleate phenotype in CU522 results from a delay or failure in macronuclear elongation and division (Fig. 4A). Although the micronuclei in dividing CU522 cells often exhibit the normal round and condensed shape seen in wild-type cells during cytokinesis, a large percentage of cells possessed micronuclei with an elongated spindle shape characteristic of micronuclear anaphase (Fig. 4B). This apparent delay in micronuclear division was detected only in cells expressing the btu1-1 (K350M) allele.

Nuclear microtubules were visualized directly with a monoclonal antibody that cross-reacts with Tetrahymena α-tubulin. Confocal immunofluorescence images of dividing CU522 and TC219 cells are shown in Fig. 5. Three observations were immediately apparent. First, abnormally long microtubule bundles were detected early in cell division as spiraling whorls that encircled macronuclei in CU522 cells but not in TC219 cells (Fig. 5B, C, and E). These structures were reminiscent of those described by Fujiu and Numata (15) for wild-type Tetrahymena cells whose macronuclear microtubules had been stabilized artificially with high concentrations of paclitaxel. Second, immediately preceding cytokinesis, unusually robust
bundles of microtubules were associated with the macronuclei. The abnormally elongated macronuclei often failed to form macronuclear constrictions (compare Fig. 5D and F). The microtubule bundles occasionally wrapped around inside one of the two daughters in cells that failed to undergo macronuclear division (Fig. 5H). A third, less conspicuous phenotype was the persistence of micronuclei in an elongate, early-anaphase spindle configuration (Fig. 5). These structures were present long after micronuclear division was completed in strain TC219 (data not shown). The micronuclei often segregated wholly with one or the other future daughter (Fig. 5H) and as a result would be expected to generate both multimicronucleate and amicronucleate cells.

For the three paclitaxel-sensitive strains tested (CU522, CU725, and CU727), careful cytological examination failed to uncover any defects in cytokinesis, such as mislocalization of the cleavage furrow (data not shown). To more completely assess whether the btu1-1 (K350M) mutation affects non-nuclear microtubule-mediated processes, we examined the structure and propagation of the oral apparatus and cortical patterning in paclitaxel-resistant (TC219) and -sensitive (CU522) strains (Fig. 6). Representative images of TC219 were indistinguishable from those of CU522. Immunocytochemical staining with polyclonal antisera specific for Tetrahymena basal bodies (5) revealed normal oral and cortical development during cell division (Fig. 6F to K). Fission furrows and oral primordia were evident in cells that contained small macronuclei or no macronuclei at all. Cortical development was similarly unaffected. Consistent with the observed defect in macronuclear transmission (Fig. 1A and Fig. 4), the macronuclei were unequally positioned across the cleavage furrows during cytokinesis in many CU522 cells (Fig. 6F to J). In contrast, the elongated macronuclei were evenly distributed across the cleavage furrows in TC219 revertants (Fig. 6A to D). Thus, asymmetric positioning of the replicated macronuclei...
may possibly account for the wide variation in macronuclear DNA contents in CU522 cell populations (Fig. 1C).

PI 3-kinase inhibitors exacerbate the macronuclear division defect in the btu1-1 mutant. PI 3-kinase activates intracellular signaling cascades that are important for the regulation of many metabolic events in the cell cycle (4). The p55 and p85 regulatory subunits of mammalian PI 3-kinase bind to β-tubulin in vitro and exhibit perinuclear staining in vivo (22, 24). Although enzymatic evidence for PI 3-kinase has not yet been obtained for Tetrahymena, biochemical studies have identified the substrate for PI 3-kinase, PIP2 (phosphatidylinositol 4,5 diphosphate) (26, 27, 52). In addition, the PI 3-kinase inhibitors wortmannin and 3-MA inhibit caspase-induced autophagy (52), similar to their well-documented effects on mammalian cells and the slime mold Dictyostelium discoideum (38). BLAST analysis of the unannotated T. thermophila genome database (http://www.tigr.org/tdb/e2k1/ttg/) revealed several strong candidate orthologues of the PI 3-kinase catalytic subunit. The most conserved homology was found in a 357-kb Tetrahymena genome scaffold (no. 8254435) within six predicted exons that span ~60% of the human catalytic subunit coding region. When the combined exons were used to query the National Center for Biotechnology Information nonredundant sequence database, strong homology was observed with the PI 3-kinase catalytic subunits from a wide variety of eukaryotes, including the slime mold D. discoideum (expected E value, 2^{-76}) and vertebrates (Homo sapiens; E value, 3^{-63}) (Fig. 7A). Similar searches identified a strong Tetrahymena candidate for the phosphatase and tensin homologue (PTEN), the enzyme that reverses the reaction catalyzed by PI 3-kinase (Fig. 7B). These data indicate that Tetrahymena encodes the key components of the classical PI 3-kinase pathway. Since mammalian PI 3-kinase associates with β-tubulin, we asked whether Tetrahymena PI 3-kinase was involved in microtubule-mediated macronuclear division. Tetrahymena strains CU428 (BTU1) and CU522 (btu1-1) and the spontaneous revertant TC219 were treated with the PI 3-kinase inhibitors wortmannin and 3-MA at concentrations similar to those used

![FIG. 5. Immunocytological analysis of microtubules associated with nuclei in strains CU522 and TC219. Confocal fluorescent images of paclitaxel-resistant TC219 cells (A, D, and G) and paclitaxel-hypersensitive CU522 cells (B, C, E, F, H, and I) are shown. Vegetatively dividing cells were maintained at 35°C prior to fixation and labeling with an antibody specific for α-tubulin (CP06; Oncogene). (A to C) Cells in the first row are representative of those in the early stages of cell division, just prior to macronuclear division stage 1 (15) and at oral development stages 2 to 4 (14). (D to F) Cells in the second row represent those that have initiated cytokinesis and macronuclear division stages 2 to 3. (G to I) The third row represents cells near the completion of cytokinesis and macronuclear development stages 5 to 6 (15). Large arrows indicate the developing cleavage furrows, whereas small arrows point to micronuclear spindles. Bar, 20 μm.](http://www.tigr.org/tdb/e2k1/ttg/)

![FIG. 6. Immunocytological analysis of cell membrane-associated microtubule-containing compartments in vegetatively dividing strains CU522 and TC219. Shown are fluorescent images of cells grown as described for Fig. 5 prior to fixation and labeling with FV4F9, a polyclonal antibody specific for basal body antigens (fluorescein isothiocyanate), and counterstained with DAPI (blue). Images in panels A to D are of TC219 cells, which do not express full-length K350M β-tubulin. Images in panels E to K are of CU522 cells, which express K350M β-tubulin and are hypersensitive to paclitaxel. The fusion furrows formed during the initiation of cytokinesis are marked with arrows. Note the pattern of membrane-associated microtubule structures: basal bodies are along ciliary rows, apical (parental) oral apparatus, and medial (daughter) oral primordia. The indistinguishable pattern in these two strains suggests that the K350M mutation has no effect on basal body- or oral apparatus-associated microtubules (see the text for discussion). Bar, 20 μm.](http://www.tigr.org/tdb/e2k1/ttg/)
When these drugs were added to synchronized cells at various times following release from G0/G1 arrest, a significant increase in the frequency of amacronucleate cell formation was observed (Fig. 8A). This effect was limited to the \( T.t. -tubulin \) K350M mutant alone and was most pronounced when inhibitors were added to S/G2-phase cells. Thus, the effect of PI 3-kinase inhibitors on macronuclear division was dependent on expression of the \( btu1 \)-1 (K350M) allele.

Since high concentrations of wortmannin inhibit mammalian MLCK (46) and an unusual myosin-like protein is required for macronuclear elongation in \( T.thermophila \) (50), we asked whether the highly specific MLCK inhibitor ML-7 could also induce amacronucleate cell formation. ML-7 at concentrations of 1 to 10 \( \mu \)M had no effect on macronuclear division (data not shown). Higher concentrations of ML-7 (50 to 100 \( \mu \)M) were toxic and therefore uninformative.

Flow cytometry was used to examine the effect of 3-MA on cell cycle progression in \( C.U.522 \) cells. A partial block in the cell cycle was observed when the drug was added to G0-arrested and refed cells (Fig. 8B; compare panels for 3 h after refeeding). There was significant accumulation of cells with lower than normal DNA content (sub-G0 peak), most likely due to the asymmetric macronuclear division documented for this strain. The sub-G0 peak was more pronounced when 3-MA was added to cultures that were predominantly in the S/G2 stage of the cell cycle (Fig. 8B, lower panels). This observation is consistent with the increased frequency of amacronucleate cell formation documented micrographs.
scopically (Fig. 8A). Since the PI 3-kinase inhibitors produced the same effect on macronuclear segregation as three known treatments that stabilize microtubules, we conclude that the PI 3-kinase pathway is required for proper macronuclear division. The dependence of PI 3-kinase inhibitors on the \textit{btu1-1} (K350M) mutation suggests that this kinase regulates microtubule-mediated processes that are dedicated to nuclear division.

**DISCUSSION**

Previous cytological studies provide a framework for the events associated with macronuclear division (23, 44). Upon completion of micronuclear mitosis, the macronucleus aligns at the cleavage furrow and divides more or less equally prior to cytokinesis. Cytoplasmic microtubules associate with the macronucleus in large numbers at this time, often forming bundles at the beginning of the elongation phase. These bundles ultimately localize to the constricted center of the bilobed dividing macronucleus. Macronuclear division and segregation typically precede the formation of a pronounced cytoplasmic cleavage furrow. Colchicine and other microtubule polymerization inhibitors block macronuclear constriction and disrupt the temporal relationship between macronuclear division and cytokinesis, often resulting in two daughter cells that receive grossly unequal quantities of nuclear material. It has been suggested that macronuclear division under these circumstances is mediated by pressure exerted by the cleavage furrow (23, 49). Since the macronuclear division defect observed in the \textit{btu1-1} mutant is strikingly similar to that described for colchicine-treated wild-type cells, this process must be regulated by dynamic changes in microtubule polymerization and depolymerization. This prediction is borne out by our observation that three regimens that stabilize microtubule polymerization (elevated temperatures, high DMSO concentrations, and sublethal concentrations of paclitaxel) increase amacronucleate cell formation in the \textit{btu1-1} mutant.

Since amitotic macronuclear division is inherently imprecise, variation in macronuclear DNA contents occurs naturally in wild-type \textit{Tetrahymena} populations. Unconventional mechanisms have evolved to maintain genic balance. For example, excess macronuclear DNA is eliminated from the macronucleus in the form of chromosome extrusion bodies (2). When DNA content drops below a threshold, the macronucleus undergoes an endoreplication cycle prior to cell division (10). The broadened macronuclear DNA content profile of CU522 cells is best explained by a failure in macronuclear division rather
than replication, since cells with higher and lower than normal DNA contents were observed. Thus, the two extremes in the population (cells with abnormally high DNA contents and those with no macronuclei at all) represent failures in macronuclear migration and/or division.

Despite the fundamental differences in micro- and macronuclear division (mitotic versus amitotic), expression of btu1-1 (K350M) affects both processes. Since cytokinesis, cortical patterning, and oral apparatus assembly appear to be unaffected, the btu1-1 allele behaves like a separation-of-function mutation. Microtubule stability is enhanced by this mutation, resulting in hypersensitivity to paclitaxel and distinct aberrations in nuclear-associated microtubules. During the early stages of cell division, CU522 cells grown at 35°C exhibit pronounced macronuclear microtubules in a spiral array (Fig. 5) that closely resemble those detected in wild-type cells treated with paclitaxel (15). Coincident with the first stages of cytokinesis, macronuclear microtubules accumulate in strange and persistent bipolar spindles. The appearance of these unusually robust microtubule bundles correlates with the failure in macronuclear division. In contrast, persistent microtubule spindles are associated with the mitotic microtubule chromosomes during anaphase. These spindles are normally absent well before macronuclear elongation and the initiation of cytokinesis (Fig. 5). Taken together, it is likely that the enhanced stability of microtubules that have incorporated β-tubulin K350M alters normal microtubule dynamics, leading to the nuclear phenotypes described in this study.

In Saccharomyces cerevisiae and fungi, astral microtubules that emanate from the spindle pole body scan the mother cell cortex and direct the migration of the mitotic nucleus to the cleavage furrow. When nuclear migration is blocked, the nucleus still divides, thereby generating a binucleate and an anucleate daughter (reference 1a and references therein). Remarkably, cells with two (or more) macronuclei were not observed, suggesting that macronuclear division does not occur in missegregation mutants. Since the Tetrahymena macronucleus lacks a spindle pole body, the mechanism for macronuclear segregation must be inherently unique. Whether macronuclear migration is required for division is not known. Current data cannot rule out the possibility that the macronuclear division defect in the btu1-1 mutant is due solely to a defect in nuclear migration.

Recent studies have demonstrated a role for structural (actin and tubulin) and motor (myosin and kinesin) proteins in macronuclear division and cytokinesis in Tetrahymena. In contrast to the btu1-1 (K350M) allele described here, mutations documented for these proteins affect nuclear and cytoskeletal processes. For example, deletion of the MYO1 gene, which encodes an unusual myosin-like protein, causes defects in endocytosis and macronuclear division (50). Similarly, expression of green fluorescent protein-tagged actin blocks macronuclear division and cytokinesis (20). Thus, both actomyosin filaments (20, 50) and cytoplasmic microtubules (this work) are required for proper macronuclear division. Whether the β-tubulin K350M mutation disrupts an interaction between cytoplasmic microtubules and actomyosin filaments remains to be determined.

It is also possible that the increase in microtubule stability and the associated macronuclear division defect may be mediated by the net loss of a positive charge in β-tubulin K350M. Alternatively, the K350M substitution may block posttranslational modification of K350 or nearby residues. Since the K350M mutation is dominant over the wild type, this mutation may affect the integrity of mixed microtubule polymers. Consistent with this prediction, the deduced β-tubulin product in the TC219 revertant strain lacks the domain(s) at the β-tubulin C terminus required for microtubule assembly (12). Consequently, the truncated protein would not be expected to contribute to microtubule-related functions. The simultaneous loss of the amacronucleate phenotype and reacquisition of paclitaxel resistance support this contention.

The pharmacological studies presented here suggest a novel role for the PI signaling pathway in macronuclear division. Specific inhibitors of PI 3-kinase induce asymmetric macronuclear division and amacronucleate cell formation. These agents effectively amplify the nuclear division defect of the β-tubulin K350M mutant in a manner analogous to those of previously established treatments that enhance microtubule stability (elevated temperatures, high DMSO concentrations, and sublethal concentrations of paclitaxel). Furthermore, the PI 3-kinase inhibitors showed no evidence of general cytotoxicity at the concentrations used, and other cytotoxic drugs that target microtubules or actomyosin filaments showed no effect on macronuclear division. Additional studies are needed to determine the exact relationship of the PI 3-kinase signaling pathway to macronuclear division. In vitro mammalian studies have demonstrated interactions between the regulatory sub-units of PI 3-kinase and α-, β-, and γ-tubulins (24), and PI 3-kinase inhibitors increase the efficacy of paclitaxel in ovarian cancer models (21). Since several distinct point mutations in the β-tubulin gene (not the K350M mutation) can lead to clinical drug resistance (reviewed in reference 37), a better understanding of how PI 3-kinase regulates microtubule dynamics in the genetically tractable eukaryote Tetrahymena may provide new insights. Recent studies in plants have identified another phospholipid signaling enzyme, phospholipase D, that interacts with microtubules in vivo and mediates stress-induced microtubule reorganization (8). Our demonstration of a functional in vivo link between β-tubulin and PI 3-kinase in Tetrahymena suggests a concerted role for these proteins in the regulation of nuclear division.

ACKNOWLEDGMENTS

This work was supported by NSF grant MCB 0220085 (D.P.R.), NIH grant GM53572 (G.M.K.), and NSF RUI grant MCB 0131175 (E.S.C.).

We thank Dorothy Shippen for careful reading of the manuscript and valuable comments.

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