Antizyme Restrains Centrosome Amplification by Regulating the Accumulation of Mps1 at Centrosomes

Running Title: Antizyme Targets Centrosomal Mps1

Christopher Kasbek, Ching-Hui Yang, and Harold A. Fisk∗
Department of Molecular Genetics
The Ohio State University
484 W. 12th Avenue
Columbus OH 43210-1292, USA.

∗To whom correspondence should be addressed (fisk.13@osu.edu)

Abbreviations:
APC/C, anaphase promoting complex/cyclosome; BrdU, 5-Bromo-deoxyUridine; DsRed, monomeric Discosoma sp. Red Fluorescent Protein; IIF, Indirect Immunofluorescence; miRDsRed, artificial microRNA embedded in the 3’UTR of the DsRed mRNA; miRGFP, artificial microRNA embedded in the 3’UTR of the GFP mRNA; MDS, Mps1 Degradation Signal;
Abstract:
Extra centrosomes are found in many tumors, and their appearance is an early event that can generate aberrant mitotic spindles and aneuploidy. Because the failure to appropriately degrade the Mps1 protein kinase correlates with centrosome overproduction in tumor-derived cells, defects in the factors that promote Mps1 degradation may contribute to extra centrosomes in tumors. However, while we have recently characterized an Mps1 degradation signal, the factors that regulate Mps1 centrosomal Mps1 are unknown. Antizyme (OAZ), a mediator of ubiquitin-independent degradation and a suspected tumor suppressor, was recently shown to localize to centrosomes and modulate centrosome overproduction, but the known OAZ substrates were not responsible for its effect on centrosomes. We have found that OAZ exerts its effect on centrosomes via Mps1. OAZ promotes the removal of Mps1 from centrosomes, and centrosome overproduction caused by reducing OAZ activity requires Mps1. OAZ binds to Mps1 via the Mps1 degradation signal and modulates the function of Mps1 in centrosome overproduction. Moreover, OAZ regulates the canonical centrosome duplication cycle, and reveals a function for Mps1 in procentriole assembly. Together, our data suggest that OAZ restrains the assembly of centrioles by controlling the levels of centrosomal Mps1 through the Cdk2-regulated Mps1 degradation signal.

Keywords:
Mps1/TTK, protein kinase, centrosome, ubiquitin-independent proteasome degradation, centrosome duplication, centriole assembly
Introduction:

Centrosomes are microtubule-organizing centers that coordinate mitotic spindle assembly, protecting genomic integrity by ensuring that each daughter cell inherits one copy of the duplicated genome. Extra centrosomes can lead to the formation of aberrant mitotic spindles that cause errors in chromosome segregation (Fisk et al., 2002; Azimzadeh and Bornens, 2007). Centrosome reduplication, or the formation of extra centrosomes within a single prolonged S-phase, represents one mechanism thought to produce extra centrosomes (Doxsey, 2002). By promoting inappropriate spindle-kinetochore attachments prior to being clustered into pseudobipolar spindles, extra centrosomes are sufficient to generate aneuploidy in vitro (Ganem et al., 2009). Moreover, extra centrosomes promote aberrant mitoses in situ (Lingle and Salisbury, 1999), and are apparent in breast (Lingle et al., 1998; Lingle et al., 2002) and prostate (Pihan et al., 2001; Pihan et al., 2003) tumors prior to aneuploidy, suggesting that centrosome reduplication might promote the genomic instability that is important in tumorigenesis (Lengauer et al., 1998; Ellsworth et al., 2004).

The canonical centrosome duplication pathway, or the duplication of the single centrosome inherited at mitosis, is initiated at the G1/S transition. A centriole assembly pathway consisting of SPD-2, the Zyg-1 protein kinase, and the Sas-4, -5, and -6 proteins was recently elucidated in C. elegans (O'Connell et al., 2001; Leidel and Gonczy, 2003; Delattre et al., 2004; Kemp et al., 2004; Pelletier et al., 2004; Leidel et al., 2005) and many of the components are conserved in vertebrates. In humans, the procentriole is templated by a cartwheel structure assembled adjacent to each of the two mother centrioles that contains the human Sas-6 orthologue, HsSas-6, and requires the presumptive Zyg-1 orthologue, Plk4 (H Abedanck et al., 2005; Strnad et al., 2007). While the human SPD-2 and Sas-4 orthologues Cep135 and CPAP also participate in the assembly of a procentriole that has the nine-fold symmetry characteristic of a centriole (Kleylein-Sohn et al., 2007; Tang et al., 2009), additional proteins not present in worms such as Centrin2 and CP110 then cooperate in extending the procentriole at a growing distal tip (Salisbury et al., 2002; Tsang et al., 2006; Kleylein-Sohn et al., 2007). In addition to its well known role in the spindle assembly checkpoint (Abrieu et al., 2001; Stucke et al., 2002; Fisk et al., 2003; Liu et al., 2003), and its more recently documented roles in DNA damage signaling (Wei et al., 2005) and the p53-dependent post-mitotic checkpoint (Huang et al., 2009), the Mps1 protein kinase is also required for centrosome duplication (Fisk and Winey, 2001; Fisk et al., 2003; Kasbek et al., 2007). However, the precise function of Mps1 in the centriole assembly pathway is unknown.

Centrosome-specific proteasome-mediated degradation is known to play a role in centrosome duplication and function in a variety of organisms. Proteasome subunits, along with several ubiquitin pathway enzymes, were identified in the recent proteomic analysis of the human centrosome (Andersen et al., 2003) and active proteasome complexes are present at centrosomes (Fabunmi et al., 2000). The anaphase promoting complex/cyclosome (APC/C) E3 ubiquitin ligase regulates mammalian centrosome separation through the degradation of Nek2 (Hames et al., 2005), and Skp1/Cdc53/F-box-dependent degradation is involved in centrosome duplication in flies (Wojcik et al., 2000) frogs (Freed et al., 1999) and mammals (Nakayama et al., 2000). We have shown that the function of Mps1 in mammalian centrosome duplication is regulated by proteasome-mediated degradation (Fisk and Winey, 2001; Fisk et al., 2003; Kasbek et al., 2007).
Cdk2 phosphorylates Mps1 at T468, attenuating the function of a degradation signal found in amino acids 420-507 (encoded by exons 12 and 13) and allowing the accumulation of a centrosomal pool of Mps1 that represents no more than 10% of total cellular Mps1 (Kasbek et al., 2007). This centrosomal Mps1 pool is critical for centrosome duplication; mutating T468 to alanine (T468A) prevents the accumulation of Mps1 at centrosomes, and a mutant Mps1 protein harboring T468A cannot substitute for the endogenous Mps1 in centrosome duplication (Kasbek et al., 2007). In contrast, replacing T468 with aspartic or glutamic acid to mimic phosphorylation at T468, or removing exons 12 and 13 that contain the Mps1 degradation signal prevents the removal of Mps1 from centrosomes even in the absence of Cdk2 activity, and the Mps1^T468A, Mps1^T468E and Mps1^Δ12/13 proteins cause centrosome reduplication (Kasbek et al., 2007; Kasbek et al., 2009). Cdk2 and the proteasome modulate centrosomal pools of Mps1 and GFP-Mps1 with little effect on whole cell levels of Mps1 or cytoplasmic pools of GFP-Mps1. In addition, GFP-Mps1^T468A that fails to accumulate at centrosomes readily fills the cytoplasm, and an exclusively centrosomal form of Mps1 (GFP-Mps1-PACT) is modulated by proteasome activity, suggesting that the Mps1 degradation signal specifically regulates the centrosomal pool of Mps1 (Kasbek et al., 2007).

Taken together, our previous data suggest that proper control over the centrosomal Mps1 pool prevents multiple rounds of centrosome duplication. In fact a variety of tumor-derived cell lines that undergo centrosome reduplication fail to properly degrade Mps1, and in one series of cell lines this correlates with both centrosome overproduction and tumorigenic potential (Kasbek et al., 2009). However, the factors that bind to the Mps1 degradation signal to regulate the centrosomal pool of Mps1 have not been identified. While the yeast Mps1 has been shown to be a substrate of the APC/C, this degradation is responsible for turning off the spindle checkpoint (Palframan et al., 2006), and its relevance to duplication of the spindle pole body (the yeast centrosome equivalent) is not clear. In addition, the centrosomal pool of vertebrate Mps1 is unlikely to be under the control of the APC/C; the Mps1 degradation signal lacks all known APC/C and SCF recognition motifs, and degradation of Mps1^AAA is not enhanced by roscovitine (Kasbek et al., 2007), a treatment that activates the APC/C and enhances the degradation of Cdc6^AAA, a non-phosphorylatable version of Cdc6 whose APC/C-dependent degradation is also attenuated by Cdk2 (Mailand and Diffley, 2005).

Recently, Mangold et al. identified a novel degradation pathway at centrosomes when they showed that ornithine decarboxylase Antizyme (OAZ) and its inhibitor (AZI) modulate centrosome number (Mangold et al., 2008). First described for its regulation of ornithine decarboxylase (ODC), an enzyme involved in polyamine biosynthesis (Coffino, 2001), OAZ binds to ODC and promotes its ubiquitin-independent degradation via the 26S proteasome (Murakami et al., 1992). Members of the antizyme family are negatively regulated by an inhibitor, called Antizyme Inhibitor (AZI), that is structurally similar to ODC but lacks enzymatic activity (Murakami et al., 1996). OAZ is a potential tumor suppressor whose overexpression reduces cell proliferation in vitro (Koike et al., 1999) and in vivo (Iwata et al., 1999; Tsuji et al., 2001; Fong et al., 2003). This phenotype is mimicked by reduced AZI expression (Choi et al., 2005; Kim et al., 2006), suggesting that OAZ and AZI cooperate to regulate ubiquitin-independent degradation of proteins involved in cell proliferation. At least three other proteins, cyclin D1, Smad1, and Aurora-A have also been shown to be targeted for destruction by OAZ (Lin et al., 2002;
Newman et al., 2004; Lim and Gopalan, 2007). Mangold et al. (2008) demonstrated that OAZ and AZI localize to centrosomes, that decreasing OAZ levels or activity lead to an increased number of cells in asynchronously growing cultures that have multiple centrosomes, and that increasing OAZ activity can suppress centrosome reduplication in tumor-derived cells. Because they also found that inhibition of ODC had no effect on centrosomes, they suggested that OAZ promotes the degradation of at least one additional protein whose continued presence promotes centriole amplification (Mangold et al., 2008). Based on our data suggesting that preventing the proteasome-mediated degradation of Mps1 causes aberrant centriole assembly (Kasbek et al., 2007), we set out to test the hypothesis that OAZ controls centrosome number by targeting Mps1. Our data not only demonstrates that OAZ regulates the centrosomal accumulation of Mps1, but reveals that OAZ also regulates the function of Mps1 in the canonical centrosome duplication process. The regulation of Mps1 degradation by a tumor suppressor suggests a connection between the failure to degrade Mps1 and centrosome defects in tumors.
Materials and Methods:

Plasmids:
Previously described plasmids used in this study include: pHF 7 (GFP), pHF 36 (GFP-Mps1), pHF 60 (GFP-Mps1<sup>Δ12/13</sup>), pHF 136 (GFP-Mps1<sup>T468A</sup>), pHF 140 (GFP-Mps1<sup>T468D</sup>), pHF 142 (GFP-Mps1-PACT), pHF 145 (GFP-Mps1<sup>Δ12/13-PACT</sup>), and pHF 148 (GFP-PACT). Plasmids created for this study include: pHF 253 (lacZ miRGFP), pHF 256 (lacZ miRDsRed), pHF 254 (OAZ miRGFP), pHF 257 (OAZ miR<sub>DsRed</sub>), pHF 258 (AZI miRGFP), pHF 259 (AZI miRDsRed), pHF 261 (DsRed-OAZ), pHF 260 (GFP-AZI), pHF 262 (DsRed-AZI), pHF 263 (GFP-OAZ-PACT), and pHF 264 (GFP-AZI-PACT). Vectors expressing MicroRNAs embedded in the 3’ UTR of the GFP mRNA (miRGFP) were created by inserting overlapping 64 base-pair oligos (see supplement for sequences) into pcDNA6.2-GW/EmGFP-miR (Invitrogen). A Dral-flanked PCR product containing DsRed (Clontech, Mountain View, CA) was used to replace GFP in pcDNA6.2-GW/EmGFP-miR to create analogous miRDsRed expression plasmids. OAZ and AZI were amplified by PCR from IMAGE clones 5533661 (Invitrogen) and 2823000 (Open Biosystems, Huntsville, AL), and ligated into pHF7 or pHF 252 to create GFP-OAZ, GFP-AZI, DsRed-OAZ, and DsRed-AZI expression plasmids. OAZ and AZI were amplified from pHF 259 and 260 with primers containing complementarity to pHF 148 and inserted into pHF 148 by In-Fusion™ cloning (Clontech) to create pHF 263 (GFP-OAZ-PACT) and pHF 264 (GFP-AZI-PACT). Thymidine at position 205 of the OAZ open reading frame was removed by site-directed mutagenesis of pHF 259, pHF 261, and pHF 263.

Cell Culture:
HeLa, U2OS, and 293 cells were cultured in DMEM supplemented with 10% FBS (HyClone, Logan, UT), 20 units/ml penicillin G, and 50µg/ml streptomycin. MCF7-GFPCentrin2 cells (the kind gift of Dr. Jefferey Salisbury, Mayo Clinic, Rochester MN) were cultured in MEM supplemented with 20 units/ml penicillin G and 50µg/ml streptomycin. Cells were cultured at 37°C in a humidified chamber in the presence of 5% CO2, and all media, supplements, and antibiotics were from Invitrogen unless otherwise indicated. S-phase arrest was achieved using treatment with 4 mM hydroxyurea (HU; Sigma, St. Louis, MO) for the times indicated. Proteasome inhibition was achieved using treatment with 5 µM MG115 (Sigma) for 4 hrs. HeLa G1 enrichment was achieved as previously described by the removal of FBS and antibiotics for 48 hrs. HeLa cells were then stimulated with DMEM containing 30% FBS and 40µM BrdU (Fisher, Pittsburgh, PA) for 8 hrs. MCF7-GFPCentrin2 cells were starved in phenol red-free MEM supplemented with 5% charcoal-dextran stripped FBS (HyClone) and 2mM L-glutamine. MCF7-GFPCentrin2 cells were released from starvation by the addition of 10nM estradiol (Calbiochem, San Diego, CA), 10ng/ml IGF (Sigma), and 10ng/ml EGF (Sigma). 293Mps1<sup>Δ12/13</sup>, HeLaGFPMps1, and HeLaGFPMps1<sup>Δ12/13</sup> cells were created as previously described (Kasbek et al., 2009). The expression of Mps1<sup>Δ12/13</sup>, GFPMps1, or GFPMps1<sup>Δ12/13</sup> was induced by the addition of Doxycycline at a final concentration of 1µg/ml.

DNA and siRNA transfections:
Plasmids were transfected using Effectene reagent (Qiagen, Valencia, CA). Stealth Mps1 (nucleotides 1360-1384) and control (scrambled nucleotides 1360-1384) siRNAs (Invitrogen) were transfected at a final concentration of 200 nM using Oligofectamine (Invitrogen). siRNA’s targeting AZI and OAZ were Smartpool AZI and Antizyme 1 (Dharmacon, Lafayette, Colorado), used at respective final concentrations of 50nM and 5nM, and were delivered into cells using Lipofectamine RNAiMAX (Invitrogen). Control siRNA for these transfections was SIGLO Lamin A/C (Dharmacon).

RT-PCR:
Total cellular RNA was isolated from HeLa using RNAqueous-4PCR (Ambion, Austin, TX). Superscript III First-Strand Synthesis System (Invitrogen) was used to produce cDNAs, which were further amplified using conventional PCR. Primers used were Antizyme 5’-atggtgaatcctcctgcagcg-3’ (sense) and 5’-ctactcctctctctccgaagac-3’ (antisense) to amplify the full-length Antizyme message of 688 base pairs. Control primers used were GAPDH 5’-aggtcggtgtgaacggatttg-3’ (sense) and 5’-tgtagaccatgtagttgaggtca-3’ (antisense) to amplify a portion of GAPDH message of 123 base pairs.

Cytology:
Antibodies and working dilutions for indirect immunofluorescence (IIF) were as follows: 1:200 GTU-88 mouse anti-γ-tubulin (Sigma), 1:200 T5192 rabbit anti-γ-tubulin (Sigma), 1:5,000 affinity purified rabbit anti-Cetn2 (as described in Yang et al., 2010), 1:500 20H5 mouse anti-Cetn2 (a kind gift of Dr. Jeffrey Salisbury), 1:200 4G9 mouse anti-Mps1 (H00007272-M02, Novus Biologicals, Littleton, CO), 1:500 rabbit anti-Antizyme 1 (Biomol International, Plymouth Meeting, PA), 1:200 mouse anti-Antizyme Inhibitor 1 (ab57169, Abcam, Cambridge, MA), 1:100 mouse anti-Sas6 (Santa Cruz Biotechnology, Santa Cruz, CA), 1:500 rabbit anti-CP110 (a kind gift from Dr. Brian Dynlacht, NYU school of Medicine), and 1:500 rat anti-BrdU (Accurate Chemicals, Westbury, NY). Secondary antibodies for IIF were donkey anti-rabbit, donkey anti-mouse, or donkey anti-rat conjugated to Alexa 350 (1:500), Alexa 488 (1:1,000), Alexa 594 (1:1,000), or Alexa750 (1:200) (all from Invitrogen), and DNA was stained with Hoechst 33342 (Sigma). Centrosome reduplication assays were performed as described previously (Kasbek et al., 2009). Briefly, centrosome number was determined by IIF with antibodies against γ-Tubulin and centrin in 3 independent experiments where at least 100 cells were counted per replicate. Measuring Mps1 centrosomal levels was performed with the Slidebook software package (Intelligent Imaging Innovations, Denver CO) using a local background correction method as previously described (Kasbek et al., 2007). Briefly, HeLa cells transfected with a variety of constructs were arrested in S-phase with a 24 hr HU treatment, then analyzed by IIF with antibodies against Mps1 and γ-Tubulin. After No Neighbors deconvolution and projection of Z-series along the Z-axis, background corrected Fluorescence Intensity of the Mps1 signal at centrosomes (F_M) was calculated from the integrated fluorescence intensities of the Mps1 signal in a small 15 x 15 pixel box (F_S) surrounding each centrosome, that of a large 20 x 20 pixel box (F_L) surrounding the first box, and the area of each box (A_S and A_L), using the formula described by Howell et al. (Howell et al., 2000); F_M = F_S - ((F_L - F_S) x (A_S + (A_L - A_S)). Twenty five cell pairs were analyzed for each condition. For Mps1-PACT experiments,
the percentage of cells in which a GFP signal (which was always exclusively centrosomal) could be detected was determined in 250-300 cells for each sample. Co-localization of GFP-Mps1 and mutants with centrosomes (using $\gamma$-tubulin) was assessed by visual inspection in 30-50 cells for each construct.

**Immunoprecipitation and Immunoblotting:**
HeLa and 293 cells were lysed in buffer composed of 50mM Tris-HCl pH8.0, 150mM NaCl, and 1% NP-40. For immunoprecipitation experiments, Mps1 complexes were immunoprecipitated by coupling N1 mouse anti-Mps1 (Invitrogen) or MDS rabbit anti-Mps1 ((Kasbek *et al.*, 2009), to Dynabeads® Protein G (Invitrogen) or anti-GFP (A11120, Invitrogen) to Dynabeads® Protein A (Invitrogen). Mps1/GFP complexes were run on SDS-PAGE and transferred to nitrocellulose for immunoblotting. Antibodies for immunoblot analysis were 1:1,000 C-19 rabbit anti-Mps1 (Santa Cruz Biotechnology), 1:2,000 rabbit anti-GFP (Sigma), 1:1,000 MDS rabbit anti-Mps1, 1:1,000 N1 mouse anti-Mps1, and 1:1,000 4G9 mouse anti-Mps1 (M02). For non-immunoprecipitation experiments, antibodies for immunoblot were as follows: 1:10,000 DM1A mouse anti $\alpha$-Tubulin (Sigma), 1:1,000 mouse anti-Antizyme Inhibitor 1, 1:2,000 rabbit anti-GFP (Sigma), 1:1,000 rabbit anti-DsRed (Clonetech), 1:2,000 mouse anti-cyclin D1 (BD Pharmingen, San Diego, CA), and 1:1,000 rabbit anti-cyclin A2 (Sigma). Secondary antibodies were Alexa680-conjugated donkey anti-mouse/rabbit (Invitrogen) and IRDye800-conjugated donkey anti-mouse/rabbit (Rockland, Gilbertsville, PA), both used at 1:10,000 for all primaries except anti-Antizyme Inhibitor 1, which required visualization by ECL™ sheep HRP-linked anti-mouse IgG (1:20,000, GE Healthcare, UK), as detected with SuperSignal® West Femto Maximum Sensitivity Substrate (Thermo Scientific, Rockford IL). All other immunoblot analysis was performed by dual-color quantitative immunoblot analysis on the Odyssey imaging system (Li-Cor, Lincoln NE) as previously described (Kasbek *et al.*, 2007). AZI levels were quantified using ImageJ (http://rsbweb.nih.gov/ij/).
Results:

Antizyme activity modulates centrosomal Mps1 levels

We chose to test our hypothesis that OAZ regulates the centrosomal pool of Mps1 in HeLa cells, because unlike U2OS cells HeLa cells do not normally reduplicate centrosomes. However, blocking the function of the Mps1 degradation signal is sufficient to cause reduplication in HeLa cells (Kasbek et al., 2007). As Mangold et al. found in other cell types (Mangold et al., 2008), both OAZ and AZI localize to centrosomes in HeLa cells (Supplemental Fig. 1), although while Mangold et al. found that OAZ was only present at centrosomes during interphase, in HeLa cells OAZ was present at centrosomes in both interphase and mitosis. Moreover, OAZ co-localized with Mps1 at centrosomes during interphase (see e.g. Fig 2B below). If our hypothesis that OAZ targets Mps1 for degradation at centrosomes is correct, increasing OAZ activity in HeLa cells should lead to the proteasome-dependent removal of Mps1 from centrosomes, while reduction of OAZ activity should increase centrosomal Mps1 levels. We tested this hypothesis using the OAZ- and AZI-specific siRNAs described by Mangold et al (2008), which in HeLa cells reduced AZI protein levels (Fig. 1A) by five-fold and rendered the OAZ mRNA undetectable (Fig. 1B; as did Mangold et al., we failed to detect OAZ on immunoblots). In addition, while both the OAZ and AZI antibodies generate apparently aspecific indirect immunofluorescence (IIF) signals that remain after siRNA treatment, the centrosomal signals generated by these antibodies are lost in virtually all cells after siRNA treatment (Fig. 1 C,D), suggesting that both siRNAs effectively deplete the centrosomal pools of the respective target proteins. Consistent with our hypothesis that OAZ regulates the centrosomal Mps1 pool, we found that there was little if any change in whole cell Mps1 levels in cells transfected with either OAZ or AZI-specific siRNAs (Fig. 1A). When we examined these cells that lacked centrosomal OAZ or AZI, we found that depletion of AZI led to the loss of Mps1 from centrosomes (Fig. 2A) that was reversed by a 4 hr treatment with the proteasome inhibitor MG115 (Fig. 2B). Depletion of OAZ did not lead to any dramatic change in centrosomal Mps1 that could be easily observed in this experiment (Fig. 2C).

In order to quantify the effect of modulating OAZ activity on centrosomal Mps1 levels, we used our previously described comparative imaging techniques (Kasbek et al., 2007). Initially, we compared the intensity of centrosomal Mps1 staining between cells expressing GFP-OAZ or GFP-AZI and concurrently imaged adjacent untransfected cells; briefly, after arresting transfected cells in S-phase with a 24 hr HU treatment, cells were stained with antibodies against γ-tubulin and Mps1, and the fluorescence intensity of Mps1 antibody staining at centrosomes was determined as described in Materials and Methods using the previously described formula (Howell et al., 2000; Kasbek et al., 2007). The data are expressed as the ratio of Mps1 intensity at centrosomes in the GFP positive cell to that in an adjacent GFP negative cell from the same image, and we analyzed twenty five separate images for each construct described below. Decreasing OAZ activity by overexpressing GFP-AZI led to a 2.05±0.48-fold increase in centrosomal Mps1 compared to paired GFP negative cells (data are summarized in Table I). Figure 2D shows an example cell pair at left, with magnified images of centrosomes shown at right to demonstrate centrosome position (γ-Tubulin, cyan) and intensity of Mps1 staining (pseudocolor). In contrast, centrosomal Mps1 levels in cells expressing GFP-OAZ were just 0.47±0.19-fold that in paired GFP negative cells, suggesting that
increasing OAZ activity led to a roughly two-fold decrease in centrosomal Mps1. Both GFP-OAZ and GFP-AZI localized to centrosomes as verified by IIF (Supplemental Figure 2).

We next sought to determine the consequences for centrosomal Mps1 of depleting OAZ or AZI. In order to apply the same imaging technique, we depleted OAZ or AZI using transient transfection of plasmid-based gene-specific synthetic microRNA (miR) sequences embedded in the 3’UTR of the GFP mRNA (miR\textsubscript{GFP}). Although these constructs only depleted the AZI protein or OAZ mRNA roughly half as effectively as standard siRNAs, they nonetheless depleted the centrosomal pool of AZI and reduced (but did not eliminate) the centrosomal pool of OAZ (Supplemental Figure 3), and the depletion achieved with these reagents was biologically relevant (see Fig. 3 and Fig. 6 below). Moreover, this partial depletion of OAZ and AZI was sufficient to modulate centrosomal Mps1 levels, although in both cases the magnitude of the change was smaller than that achieved using GFP-OAZ or GFP-AZI. OAZ miR\textsubscript{GFP} caused a 1.54±0.26-fold increase in centrosomal Mps1 levels, while centrosomal Mps1 levels in cells expressing AZI-miR\textsubscript{GFP} were just 0.61±0.12-fold that in untransfected cells. Cells expressing lacZ miR\textsubscript{GFP} (which serves as a dual control for the expression of both miRs and GFP) showed no change in centrosomal Mps1 levels, which were 1.02±0.09-fold that in untransfected cells (Table I), suggesting that the changes in Mps1 levels were the consequences of modulating OAZ activity. Because this method is based on antibody staining, we cannot completely rule out an effect of OAZ on antigen accessibility. However, we observed no such effects on any other centrosome marker in our experiments (e.g. γ-Tubulin, Centrin, or CP110, data not shown), and a brief treatment with the proteasome inhibitor MG115 restores Mps1 antibody staining in AZI-depleted cells (e.g. Fig. 2). We also observe similar effects with GFP-Mps1 (see Fig. 7 below), making an effect on antigen accessibility unlikely. Therefore, this data suggest that reducing the levels or activity of OAZ increases centrosomal Mps1, while increasing OAZ activity or levels of decreases centrosomal Mps1.

**Antizyme activity modulates centrosome duplication in an Mps1-dependent manner**

Our previous studies have demonstrated that preventing the degradation of Mps1 at centrosomes causes centrosome reduplication in HeLa cells (Kasbek \textit{et al.}, 2007; Kasbek \textit{et al.}, 2009). The observation by Mangold \textit{et al.} (2008) that OAZ modulates centrosome number led us to hypothesize that OAZ affects centrosome duplication by modulating centrosomal Mps1. Our observations that decreasing OAZ activity increases centrosomal Mps1 with little effect on whole-cell Mps1 levels are consistent with this hypothesis, which predicts that reducing OAZ activity should cause centrosome reduplication in HeLa cells that requires Mps1. Depleting AZI with standard siRNAs had no effect on centrosome number, while roughly 20% of HeLa cells transfected with OAZ siRNAs had undergone centrosome reduplication (Fig. 3A). Figure 2C above shows a representative OAZ-siRNA transfected cell with more than two centrosomes. To test whether this reduplication requires Mps1, we sequentially transfected HeLa cells with control or Mps1-specific siRNAs (Kasbek \textit{et al.}, 2007), and either OAZ miR\textsubscript{GFP} or GFP-AZI to reduce OAZ levels or activity, then determined centrosome number in GFP positive cells after 48 hrs of S-phase arrest (Mps1 depletion was verified by immunoblotting). OAZ miR\textsubscript{GFP} caused centrosome reduplication in cells transfected with...
control siRNA (Fig. 3B, black bars, “OAZ miR”) at levels similar to that observed with OAZ siRNA (e.g. Fig. 3A). A similar level of centrosome reduplication was observed in GFP-AZI expressing cells transfected with control-siRNA (Fig. 3B, black bars, “AZI”). Because cells with excess γ-Tubulin foci also had excess centrioles (as judged by centrin staining), this observation reflects an effect of OAZ on centrosome duplication rather than on centrosome integrity, consistent with the observations of Mangold et al (2008) in U2OS cells. However, centrosome reduplication associated with either OAZ miR\textsuperscript{GFP} or GFP-AZI was abrogated by Mps1-specific siRNA (Fig. 3B, red bars), demonstrating that Mps1 is required for the centrosome reduplication caused by reducing OAZ activity. Because Mps1 may simply be required for all types of centrosome reduplication, this observation does not unambiguously place Mps1 and OAZ in the same pathway. However, a negative result would have indicated that OAZ influences centrosome duplication independently of Mps1.

Antizyme Targets the Centrosomal Pool of Mps1 through the Mps1 Degradation Signal

Both Mps1 and OAZ are found in the cytoplasm as well as at centrosomes, and the two proteins might interact at either location. The centrosomal pool of Mps1 is regulated by a degradation signal found within Mps1 amino acids 420-507 (encoded by exons 12 and 13) whose function has little or no effect on other Mps1 pools (Kasbek et al., 2007). The PACT domain has been frequently used to tether proteins exclusively to centrosomes in order to assess their centrosomal functions (Gillingham and Munro, 2000; Keryer et al., 2003; Mikule et al., 2007). In order to test whether OAZ can act on the centrosomal pool of Mps1, we utilized two previously described Mps1 constructs that are tethered to centrosomes via the AKAP450 PACT domain, GFP-Mps1-PACT and GFP-Mps1\textsuperscript{Δ12/13}-PACT that lacks the Mps1 degradation signal (Kasbek et al., 2007). While GFP-Mps1 is largely cytoplasmic (see e.g. Fig. 7), to the extent that can be determined by fluorescence microscopy GFP-Mps1-PACT (Fig. 4A) and GFP-Mps1\textsuperscript{Δ12/13}-PACT (Fig. 4B) are exclusively centrosomal. Despite being tethered to centrosomes via PACT binding sites, centrosomal accumulation of GFP-Mps1-PACT is still regulated by the Mps1 degradation signal in a proteasome-dependent manner; a 4 hr treatment with the proteasome inhibitor MG115 leads to a 5-fold increase in the percentage of cells where GFP-Mps1-PACT can be detected, but has no effect on GFP-Mps1\textsuperscript{Δ12/13}-PACT (Fig. 4C, gray bars). AZI-siRNA had no appreciable effect on either construct, while OAZ-siRNA caused an increase in the percentage of cells in which GFP-Mps1-PACT can be detected that was similar to that caused by MG115 treatment (Fig. 4C, “OAZ”). However, like DMSO treatment, OAZ-siRNA had very little effect on GFP-Mps1\textsuperscript{Δ12/13}-PACT (Fig. 4C). This demonstrates that OAZ can act on the centrosomal Mps1 pool through the Mps1 degradation signal. Because GFP-Mps1-PACT can accumulate at centrosomes by binding at PACT binding sites, its modulation by OAZ suggests that OAZ acts directly on Mps1, rather than by depleting a centrosomal Mps1 binding site. We did not examine centrosome number in these experiments, because overexpression of the PACT domain on its own can influence centrosome number and structure (Gillingham and Munro, 2000; Keryer et al., 2003; Kasbek et al., 2007; Mikule et al., 2007).

Antizyme Binds to Mps1 via the Mps1 Degradation Signal
Because known substrates of OAZ such as ODC, cyclin D1, and Smad1 are targeted to the proteasome through a physical interaction with OAZ (Murakami et al., 1992; Lin et al., 2002; Newman et al., 2004), we explored a physical interaction between Mps1 and OAZ. As judged by immunoblotting with an antibody against GFP, a small fraction of GFP-OAZ co-immunoprecipitates with Mps1 from HeLa cells (Fig. 5A). This suggests that only a limited fraction of Mps1 interacts with OAZ, and vice versa, and is consistent with an interaction between Mps1 and OAZ that is limited to centrosomes. The interaction between Mps1 and GFP-OAZ was not altered when we treated S-phase arrested cells with the Cdk2 inhibitor Roscovitine and/or the proteasome inhibitor MG115 (data not shown). To determine whether OAZ interacts with Mps1 through the Mps1 degradation signal, we utilized the previously described HeLa-GFPMps1Δ12/13 cell line that expresses physiological levels of GFP-Mps1Δ12/13 from a doxycycline (Dox)-inducible promoter (Kasbek et al., 2009). This non-degradable mutant protein lacks the Mps1 degradation signal, and unlike wild type GFP-Mps1 causes centrosome reduplication in a variety of cell types (Kasbek et al., 2007; Kasbek et al., 2009). We transfected HeLa-GFPMps1Δ12/13 cells with DsRed-OAZ, and then performed immunoprecipitations with three different antibodies. As shown by immunoblot using the N1 monoclonal antibody against Mps1 (Fig. 5B, top panel), the N1 antibody immunoprecipitates both endogenous Mps1 and GFP-Mps1Δ12/13, while the MDS antibody (directed against the Mps1 degradation signal; Kasbek et al., 2009) only immunoprecipitates endogenous Mps1, and the GFP antibody only immunoprecipitates GFP-Mps1Δ12/13. A diagram of Mps1, Mps1Δ12/13, and the antibodies used in this experiment is shown in Fig. 5C. DsRed-OAZ co-immunoprecipitated with both N1 and MDS antibodies, but failed to co-immunoprecipitate with the GFP antibody, suggesting that it can bind to endogenous Mps1 but not to GFP-Mps1Δ12/13. The failure of DsRed-OAZ to bind to GFP-Mps1Δ12/13 does not appear to be due to the presence of the GFP tag, because in the previously described 293-Mps1Δ12/13 cell line (Kasbek et al., 2009), untagged Mps1Δ12/13 fails to interact with GFP-OAZ (Supplemental Figure 4). Together, these observations demonstrate reciprocal co-immunoprecipitation between Mps1 and GFP-OAZ, and suggest that the ability of OAZ to bind to Mps1 requires the presence of the Mps1 degradation signal.

**Antizyme Modulates Mps1 Function Through the Mps1 Degradation Signal**

The function of Mps1 in centrosome duplication is regulated by Cdk2-dependent phosphorylation within the Mps1 degradation signal at T468, which attenuates the removal of Mps1 from centrosomes (Kasbek et al., 2007; Kasbek et al., 2009). Our hypothesis that OAZ is responsible for the removal of Mps1 from centrosomes in the absence of Cdk2 suggests that both GFP-Mps1Δ12/13 that lacks the Mps1 degradation signal and the phosphomimetic GFP-Mps1T468D should be recalcitrant to OAZ activity. To test this suggestion, we utilized the ability of Mps1 to accelerate the onset of centrosome reduplication in U2OS cells. While U2OS cells naturally undergo centrosome reduplication, the presence of extra centrosomes only becomes apparent after 48 hrs of S-phase arrest. As we first demonstrated and others have since verified, overexpression of Mps1 accelerates the onset of this phenotype so that extra centrosomes are apparent after only 24 hrs of S-phase arrest (Fisk et al., 2003; Kanai et al., 2007; Kasbek et al., 2007; Kasbek et al., 2009). As expected, GFP-Mps1, GFP-Mps1Δ12/13, and GFP-Mps1T468D each
accelerated the onset of centrosome reduplication in U2OS cells doubly transfected with lacZ miR^{DsRed}. Consistent with the hypothesis that OAZ stimulates the degradation of Mps1 through the Cdk2-regulated degradation signal, both AZI miR^{DsRed} and DsRed-OAZ attenuated the ability of GFP-Mps1 to accelerate reduplication in U2OS cells, but had no effect on either GFP-Mps1^{Δ12/13} or GFP-Mps1^{T468D} (Fig. 6A). These observations suggest that the effect of OAZ on Mps1 function in this assay both requires the Mps1 degradation signal and is attenuated by T468 phosphorylation.

As previously reported (Kasbek et al., 2007; Kasbek et al., 2009), GFP-Mps1^{T468A} that cannot be phosphorylated at T468 had little effect on centrosome reduplication in U2OS cells expressing lacZ miR^{DsRed} (Fig. 6B). If OAZ is responsible for removing Mps1 from centrosomes in the absence of Cdk2 activity, reducing OAZ activity should allow GFP-Mps1^{T468A} to accelerate centrosome reduplication in U2OS cells. However, reducing OAZ activity was sufficient to accelerate centrosome reduplication in U2OS cells (Fig. 6B, DsRed-AZI), so we did not directly test this suggestion. Accordingly, we set out to examine the effects of OAZ on the centrosomal accumulation of the various Mps1 proteins in this assay. In S-phase arrested cells GFP-Mps1 is overexpressed by roughly 50-fold with respect to endogenous Mps1 and is largely cytoplasmic (Kasbek et al., 2007). This cytoplasmic signal is frequently granular in appearance with occasional aggregates (Fig. 7). Thus, the centrosomal accumulation of GFP-Mps1 constructs is apparent as a locally concentrated GFP signal in the immediate vicinity of centrosomes that is surrounded by a diffuse and frequently granular cytoplasmic signal. Wild type GFP-Mps1 was locally concentrated at centrosomes in 86% of U2OS cells doubly transfected with DsRed (Fig. 7A “DsRed”). However, GFP-Mps1 showed no such concentration at centrosomes in 60% of U2OS cells doubly transfected with DsRed-OAZ (Fig. 7A “DsRed-OAZ”). In contrast, GFP-Mps1^{T468A} showed no detectable concentration at centrosomes in 59% of U2OS cells doubly transfected with DsRed (Fig. 7B, “DsRed”), but was readily apparent at centrosomes in 87% of cells doubly transfected with DsRed-AZI (Fig. 7B, “DsRed-AZI”). GFP-Mps1^{T468D} was apparent at centrosomes in roughly 96% of U2OS cells doubly transfected with DsRed (Fig. 7C, “DsRed”), and this number was only modestly decreased to 88% upon double transfection with DsRed-OAZ (Fig. 7C, “DsRed-OAZ”). We noted that DsRed alone localized to centrosomes in these experiments. However, this does not affect conclusions regarding centrosomal localization of OAZ, because the OAZ antibody exhibits centrosomal staining that is depleted by OAZ siRNAs (e.g. Fig. 1), and GFP-OAZ localizes to centrosomes while GFP alone does not (e.g. Supplemental Figure 2). Moreover, DsRed alone had no effect on the centrosomal accumulation of GFP-Mps1 constructs or centrosome duplication (e.g. Fig. 6). Therefore, the observations that DsRed-AZI enhances centrosomal accumulation of GFP-Mps1^{T468A} while GFP-Mps1^{T468D} is insensitive to DsRed-OAZ suggest that OAZ participates in the removal of Mps1 from centrosomes, and that its ability to do so is attenuated by T468 phosphorylation. Together, these observations provide further support for the hypothesis that OAZ modulates centrosomal Mps1 via the Mps1 degradation signal. Moreover, the observation that GFP-Mps1^{T468D} and GFP-Mps1^{Δ12/13} are insensitive to OAZ provides additional evidence that OAZ acts directly on Mps1 rather than on an Mps1 binding partner, because if OAZ removed an Mps1 binding site, modulating OAZ levels and/or activity should affect all Mps1 proteins equally.
**Antizyme Regulates the Function of Mps1 During the Canonical Duplication Cycle**

Data presented thus far demonstrate that OAZ prevents centrosome reduplication by targeting Mps1 for degradation. However, it does not address whether OAZ has a general role in centrosome duplication, or if it acts solely to prevent the aberrant execution of excessive rounds of duplication. Accordingly, we also tested whether OAZ regulates the function of Mps1 in the canonical centrosome duplication event that occurs as cells progress from G1 into S-phase. To this end, HeLa cells transfected with GFP or GFP-OAZ were enriched in G1 and stimulated to rapidly enter S-phase according to our previously described protocol (Fisk et al., 2003), as described in Materials and Methods. Eight hrs after release from G1 enrichment, centriole number was assessed with antibodies against Centrin2 and CP110 in cells that were in S-phase. Roughly 95% of cells expressing GFP alone that were in S-phase (as judged by BrdU staining) had completed centrosome duplication and had four centrioles, while only 5% still had two centrioles (as judged by Centrin2 staining; Fig. 8A). However, roughly 20% of GFP-OAZ expressing cells that were in S-phase had two centrioles (Fig. 8A,B). This increase in cells with two centrioles is unlikely to be due to an effect of OAZ on cell cycle progression because there was no difference in the percentage of BrdU positive cells between GFP and GFP-OAZ transfections, and because we examined centrioles only in cells that were in S-phase. Moreover, overexpression of the dominant negative GFP-Mps1KD (Fisk and Winey, 2001; Fisk et al., 2003) led to a similar level of S-phase cells with two centrioles (Fig. 8A,C), suggesting that the increased percentage of cells with two centrioles may be a consequence of inhibiting Mps1 activity. We made similar observations in MCF7-GFPCentrin2 cells (D'Assoro et al., 2001) after their release into the cell cycle from G0 arrest (Fig. 8A), suggesting that this effect of OAZ can be generalized to other cell types.

We further analyzed HeLa cells expressing GFP-OAZ and GFP-Mps1KD with an antibody against HsSas-6, a marker of procentriole assembly. All BrdU positive cells expressing GFP alone, including those that had only two Centrin2- or CP110-positive centrioles, had formed two procentrioles as judged by HsSas-6 staining (Fig. 8D). Moreover, all cells expressing GFP alone had either zero or two HsSas-6 foci (for comparison, only cells with two centrioles are shown in Fig. 8). In contrast, none of the GFP-OAZ (Fig. 8E) or GFP-Mps1KD (Fig. 8F) expressing cells with two Centrin2- or CP110-positive centrioles had two HsSas-6 positive structures; 88% and 95%, respectively, had a single HsSas-6 positive structure that did not contain either CP110 or Cetn2 and was positioned roughly between the two centrioles (the remaining GFP-OAZ or GFP-Mps1KD expressing cells with two centrioles had zero HsSas-6 foci). These observations suggest that cells expressing GFP-OAZ or GFP-Mps1KD had initiated centrosome duplication but were blocked or delayed at a very early stage.

Data described above support the suggestion that the effect of OAZ on centriole assembly is due to increased degradation of centrosomal Mps1. In order to rule out any unintended consequence of serum starvation, we analyzed centriole number in BrdU positive cells from asynchronously growing cultures of HeLa cells. After transfection with either DsRed or DsRed-OAZ, HeLa cells were labeled with BrdU for 4h, and centriole number was assessed by IIF with an antibody against Centrin2. Approximately 0.7% of BrdU-positive cells expressing DsRed alone had only two centrioles, while 6.7%
of BrdU-positive DsRed-OAZ expressing cells had only two centrioles (Fig. 9). The lower percentage of cells with two centrioles in asynchronous cells suggests that the phenotype of OAZ overexpression reflects a delay in procentriole formation, rather than an outright block. In order to determine whether this delay is mediated by decreased Mps1 function we repeated the experiment in the previously described HeLa-GFP-Mps1 cell line that overexpresses GFP-Mps1 by roughly 5-fold with respect to endogenous Mps1 (Kasbek et al., 2009). Approximately 1.3% of BrdU positive HeLa-GFP-Mps1 cells expressing DsRed alone had two centrioles, and this percentage increased only slightly to 2% in HeLa-GFP-Mps1 cells expressing DsRed-OAZ (Fig. 9), suggesting that increasing Mps1 levels compensates for the effect of OAZ on centriole assembly. The observation that GFP-Mps1 overexpression can rescue the centriole assembly defect caused by OAZ overexpression suggests that a reduction in Mps1 levels and/or activity is responsible for the OAZ-mediated delay. This observation also provides additional support for the suggestion that OAZ targets Mps1 directly, because if OAZ removed a protein or proteins responsible for the binding of Mps1 to centrosomes, GFP-Mps1 on its own would not be sufficient to rescue the delay in centriole assembly caused by OAZ. Together, our data suggest that OAZ regulates the function of Mps1 in the canonical centrosome duplication pathway at the G1/S transition in multiple cell types, and that suppressing OAZ activity can increase the levels of centrosomal Mps1 and lead to Mps1-dependent centrosome reduplication.
Discussion:

Based on their data that OAZ prevents centrosome reduplication, Mangold et al. suggested that OAZ promotes the proteasome-mediated degradation of some factor required for centrosome amplification (Mangold et al., 2008). Because preventing Mps1 degradation causes centrosome reduplication, we hypothesized that this factor might be Mps1. Data presented here supports this hypothesis and further suggest that OAZ regulates Mps1 early in procentriole assembly, as evidenced by our observations of an aberrant HsSas-6 containing structure that lacks Centrin2 and CP110. A similar consequence of inhibiting Mps1 was observed in yeast; overexpressed Spc42p is not properly assembled into the Spindle Pole Body in the mps1-1 mutant, but forms an aberrant structure adjacent to the Spindle Pole Body (Castillo et al., 2002).

We considered the possibility that rather than suppressing centrosome duplication per se, OAZ overexpression causes a cell cycle defect through another of its substrates, cyclin D1. Spermidine upregulates OAZ causing G0 arrest due to degradation of cyclin D1 (Newman et al., 2004). Because Mangold et al. found that neither cyclin D1 nor ODC were responsible for the effect of OAZ on centrosome duplication (Mangold et al., 2008), and because we found that GFP-OAZ and GFP-AZI had no effect on either S-phase entry or cyclin D1 levels (Supplementary Figure 5), we conclude that OAZ affects centrosome duplication directly. We also considered the possibility that OAZ affects centrosomal Mps1 indirectly by acting on a protein(s) responsible for targeting Mps1 to centrosomes, or by affecting the maturation state or duplication potential of centrosomes. If OAZ removed a centrosomal Mps1 binding site, increasing OAZ should affect all forms of Mps1 identically, increasing Mps1 should be insufficient to counteract for loss of this binding site, and tethering Mps1 to centrosomes independently of this binding site should render it resistant to OAZ. Furthermore, if OAZ restrained centrosome duplication by affecting the maturation state of centrosomes or otherwise affected their ability to duplicate, Mps1$^{A12/13}$ and Mps1$^{T468D}$ would be unable to promote centrosome reduplication in cells overexpressing OAZ. However, our data contradict each of these predictions; the centrosomal accumulation of GFP-Mps1$^{A12/13}$ and GFP-Mps1$^{T468D}$ are insensitive to OAZ, overexpression of Mps1 is sufficient to reverse the effect of OAZ overexpression, and OAZ can modulate GFP-Mps1-PACT, and GFP-Mps1$^{A12/13}$ and GFP-Mps1$^{T468D}$ cause centrosome reduplication in cells overexpressing OAZ. The simplest explanation of these observations is that OAZ acts directly on Mps1 to modulate centrosome duplication. Moreover, OAZ appears to specifically affect centrosomal Mps1. This suggestion is supported by the observations that modulating OAZ has little noticeable effect on whole cell Mps1 levels or cytoplasmic GFP-Mps1, that only small fractions of Mps1 and OAZ interact, and that OAZ can modulate GFP-Mps1-PACT. Interestingly, OAZ-PACT and AZI-PACT constructs had no effect on centrosome numbers. We assume that the PACT domain perturbs folding or activity of OAZ and AZI, but it is also possible that the constructs are non-functional because the OAZ-Mps1 interaction occurs in the cytoplasm. However, in that case the interaction cannot be required for centrosomal targeting, because Mps1 accumulates at centrosomes in OAZ-depleted cells, and OAZ accumulates at centrosomes in Mps1 depleted cells (data not shown). While it is difficult to prove that the interaction occurs at centrosomes, or that Mps1 is actually degraded at centrosomes, the simplest hypothesis that incorporates all of our observations is that OAZ targets Mps1 for degradation at centrosomes. But regardless
how OAZ controls centrosomal Mps1 levels, our data strongly suggest that OAZ exerts control over centrosome duplication by regulating the accumulation of Mps1 at centrosomes.

In yeast the cyclin-dependent kinase Cdc28p suppresses the proteasome-dependent APC/C-mediated degradation of Mps1 (Palframan et al., 2006). However, this degradation is responsible for turning off the spindle checkpoint; its relevance to the control of spindle pole body duplication is not clear, and the Cdk2-regulated degradation signal in human Mps1 lacks all known APC/C recognition motifs. The identification of Mps1 as an OAZ target suggests that the function of vertebrate Mps1 in centrosome duplication is regulated by an OAZ-mediated ubiquitin-independent proteasomal mechanism. This suggestion is supported by four key observations; Modulating OAZ activity modulates the levels of Mps1 at centrosomes in a proteasome-dependent manner; OAZ participates in the constitutive removal of GFP-Mps1T468A from centrosomes; the binding of OAZ to Mps1 requires the Mps1 degradation signal; and the biological effects of OAZ on both Mps1 and centrosome duplication are regulated by T468. We attempted to test whether T468 phosphorylation blocks the OAZ-Mps1 interaction. Unfortunately, technical considerations in transient transfections hampered co-immunoprecipitations with GFP-Mps1T468A and GFP-Mps1T468D, and recombinant OAZ and Mps1 (wild type, or T468A/T468D mutants) did not interact in vitro (data not shown). While this later observation suggests the possibility that OAZ may not interact directly with Mps1, the interaction may simply require post translational modifications not present in bacterially expressed proteins. Regardless, our findings strongly suggest that OAZ regulates centrosome duplication by targeting Mps1 for degradation through the Cdk2-regulated Mps1 degradation signal. There are a growing number of examples in biology of ubiquitin-independent proteasomal degradation (Jariel-Encontre et al., 2008; Yuksek et al., 2009) such as that mediated by OAZ (Murakami et al., 1992; Lin et al., 2002; Newman et al., 2004). In the case of the ubiquitin-independent degradation of Aurora-A, OAZ also cooperates with AURKAIP1 (Lim and Gopalan, 2007), suggesting the possibility that additional factors may cooperate with OAZ to promote Mps1 degradation.

We hypothesize that the accumulation of Mps1 at centrosomes is regulated by antagonistic OAZ binding and T468 phosphorylation. In this model, phosphorylation is transient because it is entrained to Cdk2, and OAZ restrains centrosome duplication when Cdk2 activity is low by binding to unphosphorylated Mps1 to promote its proteasome-dependent removal from centrosomes (see Supplemental Figure 6). Although we assume this binding occurs at centrosomes and is blocked by T468 phosphorylation, we have been unable to directly test these assumptions. The observation that GFP-AZI enhances the centrosomal accumulation of Mps1T468A suggests that there is sufficient constitutive OAZ activity at centrosomes to promote removal of un-phosphorylated Mps1, although the presence of AZI at centrosomes suggests that OAZ is regulated. Although OAZ has a yeast orthologue, the regulation of Mps1 by OAZ is not likely to be evolutionarily conserved because the Mps1 degradation signal is not conserved between yeast and humans.

Interestingly, Mps1 has been shown to be stabilized by other kinases in addition to Cdk2. CHK2 stabilizes Mps1 by phosphorylation of T288 in response to DNA damage (Wei et al., 2005), and oncogenic B-Raf stabilizes Mps1 via Erk (Cui and Guadagno, 2008), suggesting that multiple pathways contribute to the control of Mps1 stability.
Together with the observation that the failure to degrade Mps1 at centrosomes causes reduplication in a variety of cell types and correlates with reduplication in tumor-derived cells (Kasbek et al., 2007; Kasbek et al., 2009), the observation that OAZ, a potential tumor suppressor, controls Mps1 degradation suggests that the failure to degrade Mps1 might have oncogenic consequences. We look forward to determining to what extent the tumor suppressive effects of OAZ relate to the centrosomal functions of Mps1, as well as identifying other factors involved in controlling the removal of Mps1 from centrosomes.
Acknowledgements

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References:


Cui, Y., and Guadagno, T.M. (2008). B-Raf(V600E) signaling deregulates the mitotic spindle checkpoint through stabilizing Mps1 levels in melanoma cells. Oncogene 27, 3122-3133.


Tables

Table I. Antizyme activity controls centrosomal Mps1 Levels.
Cells expressing GFP-AZI, GFP-OAZ, OAZ miRGFP, AZI miRGFP, or lacZ miRGFP were arrested in S-phase and analyzed by IIF using antibodies against Mps1 and γ-tubulin. The intensity of Mps1 staining was determined as described in the materials and methods, and compared between paired GFP positive and GFP negative cells that were contained within the same image. Numbers indicate the average fold change in centrosomal Mps1 levels ± standard deviation in the GFP positive cell compared to adjacent GFP negative cell for 25 cell pairs (as pictured in Fig. 2D). The differences between all pairwise combinations were highly significant (p<0.0001), as judged by one-way ANOVA followed by Tukey’s HSD post hoc test, with the exception of GFP-OAZ and AZI miRGFP (both of which lead to increased OAZ activity) for which the difference was not significant (p<0.37). The difference between GFP-AZI and OAZ miRGFP that should both lead to decreased OAZ activity was statistically significant, most likely due the inefficient depletion achieved with OAZ miRGFP.

<table>
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<th>Plasmid</th>
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<th>GFP-OAZ</th>
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<th>AZI miRGFP</th>
<th>lacZ miRGFP</th>
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<td>Mps1 level; Fold Change ± S.D.</td>
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<td>0.47 ± 0.19</td>
<td>1.54 ± 0.26</td>
<td>0.61 ± 0.12</td>
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Figure Legends:

Figure 1. OAZ- and AZI-specific siRNAs deplete the centrosomal OAZ and AZI pools but do not affect the whole cell levels of Mps1.
(A) AZI siRNA results in an approximately 5-fold decrease in the AZI protein (relative levels are indicated below each lane), but neither AZI nor OAZ siRNA significantly affected Mps1 whole cell levels, as compared to control siRNA. Shown are immunoblots of siRNA transfected cells; Mps1 and the α-tubulin loading control were stained on the same blot and imaged using the LI-COR Odyssey, while AZI was stained on a separate blot and imaged with ECL. (B) RT-PCR of total RNA isolated from HeLa cells treated with control or OAZ siRNA shows the OAZ message is depleted by OAZ siRNA. The negative control (-RT) shows the product is not due to contaminating chromosomal DNA. Final concentrations of siRNA in µM used are listed above each lane. Lower panel shows amplification with GAPDH-specific primers as a loading control. (C, D) The centrosomal pools of OAZ and AZI are depleted by siRNA treatment. IIF of HeLa cells transfected with control (siControl), OAZ-specific (siOAZ), or AZI-specific (siAZI) siRNAs for 48 hours. Shown are representative cells stained with γ-tubulin (green), Mps1 (red), and Hoechst (blue); bar = 5 µm. Panels at the right of each cell show digitally magnified images of the box surrounding the centrosomes; bar = 1 µm.

Figure 2. Antizyme activity modulates centrosomal Mps1 in a proteasome-dependent manner.
(A) Increasing Antizyme activity by depleting its inhibitor AZI decreases centrosomal Mps1. (B) The loss of Mps1 from centrosomes in AZI-siRNA transfected cells is reversed by inhibiting the proteasome. (A,B) siRNA-transfected HeLa cells were arrested in S-phase with a 24 HU treatment, then treated with DMSO (A) or MG115 (B) for 4 hr. Shown are representative cells stained with γ-tubulin (green), Mps1 (red), and Hoechst (blue); bar = 5 µm. Panels to the right of each cell show digitally magnified images of the box surrounding the centrosomes; bar = 1 µm. (C) Decreasing Antizyme activity by siRNA depletion causes centrosome reduplication, along with a slight increase in Mps1 levels. siRNA-transfected HeLa cells were arrested in S-phase with a 24 HU treatment. Shown are representative cells stained with γ-tubulin (green), Mps1 (red), and Hoechst (blue); bar = 5 µm. Panels to the right of each cell show digitally magnified images of the box surrounding the centrosomes; bar = 1 µm. (D) Decreasing Antizyme activity by overexpressing GFP-AZI increases centrosomal Mps1 levels. HeLa cells were transfected with GFP-AZI, arrested in S-phase with a 24 hr HU treatment, then 25 cell pairs were imaged as described in materials and methods. Shown at left is a representative cell pair showing GFP-AZI (green), Mps1 (red), DNA (blue) and γ-Tubulin (cyan), with arrows indicating the corresponding panel at right for each cell; bar = 5 µm. The panels at right show magnified images of the box surrounding centrosomes for the cells in the left panel to illustrate centrosome position (γ-tubulin staining, cyan) and Mps1 levels (shown in pseudocolor to highlight differences between the two cells); arrow heads indicate centrosome position; bar = 1 µm.

Figure 3. Antizyme depletion causes Mps1 dependent Centrosome reduplication in
HeLa cells.
(A) OAZ siRNA caused an approximately 5-fold increase in cells with more than two centrosomes (and more than 4 centrioles). HeLa cells were transfected with the indicated siRNA then arrested in S-phase for 48 hr. (B) Centrosome reduplication in OAZ-depleted cells requires Mps1. HeLa cells were sequentially transfected with control (black bars) or Mps1-specific siRNA (red bars) followed by lacZ miR<sup>GFP</sup> (lacZ miR), OAZ miR<sup>GFP</sup> (OAZ miR), or GFP-AZI (AZI), then arrested in S-phase for 48 hr. (A and B) Centrosome number was determined by IIF with antibodies against γ-Tubulin and centrin. Values represent the mean ± standard deviation for 3 independent experiments where at least 100 cells were counted per replicate. Mps1 depletion was verified in (B) by immunoblotting with antibodies against Mps1 and the α-Tubulin loading control, as shown below the bar graph.

Figure 4. Centrosomally localized Mps1-PACT is modulated by Antizyme and proteasome activities through the Mps1 degradation signal.
(A and B) GFP-Mps1-PACT (A) and GFP-Mps1<sup>Δ12/13</sup>-PACT (B) can only be detected at centrosomes. Shown are the GFP-Mps1-PACT or GFP-Mps1<sup>Δ12/13</sup>-PACT signal (green) from representative S-phase arrested HeLa cells stained with Centrin (red), and Hoechst (blue); bar = 5 µm. Lower panels show digitally magnified images of the box surrounding the centrosomes for each cell; bar = 1 µm. (C) OAZ and proteasome activity modulate GFP-Mps1-PACT but not GFP-Mps1<sup>Δ12/13</sup>-PACT that lacks the Mps1 degradation signal. HeLa cells were sequentially transfected with control (Con), OAZ-, or AZI-specific siRNAs, followed by GFP-Mps1-PACT (Mps1-PACT) or GFP-Mps1<sup>Δ12/13</sup>-PACT (Mps1<sup>Δ12/13</sup>-PACT), arrested in S-phase with a 24 hr HU treatment, then treated with either DMSO (black bars) or MG115 (gray bars) for 4 hrs. The percentage of cells in which a GFP signal could be detected was determined in 250-300 cells for each sample, and normalized to that in the respective DMSO-treated control siRNA sample.

Figure 5. Antizyme binds to the Mps1 degradation signal.
OAZ binds to Mps1 in HeLa cells, but fails to bind to a form of Mps1 lacking the degradation signal. (A) GFP-OAZ co-immunoprecipitates with Mps1. HeLa cells were transfected with either GFP or GFP-OAZ, then arrested in S-phase. Mps1 complexes were immunoprecipitated from lysates with the Mps1 N1 antibody, and analyzed by dual color immunoblotting with rabbit antibodies against GFP or Mps1. Input (I) shows 2.5% of the lysate used in the preclear (PC, showing any proteins that bind to beads independently of antibody) and immunoprecipitation (IP). (B) DsRed-OAZ (open arrow) binds to endogenous Mps1 (closed arrow) but not GFP-Mps1<sup>Δ12/13</sup> (arrowhead). HeLa-GFP-Mps1<sup>Δ12/13</sup> cells induced with doxycycline were transfected with DsRed-OAZ, then arrested in S-phase. Mps1 complexes were immunoprecipitated from lysates with three different antibodies and analyzed by dual color immunoblotting with antibodies against Mps1 (N1) and DsRed. Input (I) shows 2.5% of the total lysate used in the preclear (PC) and immunoprecipitations; PC shows proteins bound to beads in the absence of antibody, while N1, MDS, and GFP shows proteins that immunoprecipitate with those antibodies. (C) Diagram of Mps1 and Mps1<sup>Δ12/13</sup> showing the location of the Mps1 degradation signal and the epitopes for the antibodies used in (A) and (B).
Figure 6. The effect of Antizyme on Mps1 function is mediated by the Mps1 degradation signal and regulated by T468 phosphorylation.

(A) Increasing Antizyme activity abrogates the ability of GFP-Mps1 to accelerate the onset of centrosome reduplication in U2OS cells, but has no effect on GFP-Mps1\textsuperscript{A12/13} or GFP-Mps1\textsuperscript{T468D}. U2OS cells were sequentially transfected with DsRed (black bars), AZI miR\textsuperscript{DsRed} (gray bars), or DsRed-OAZ (white bars) followed by GFP-Mps1, GFP-Mps1\textsuperscript{A12/13}, or GFP-Mps1\textsuperscript{T468D} as indicated. After 24 hr of S-phase arrest centrosome number was determined by IIF with antibodies against \(\gamma\)-Tubulin and centrin. Values represent the mean \(\pm\) standard deviation for 3 independent experiments where at least 100 cells were counted per replicate. (B) DsRed-AZI accelerates the onset of centrosome reduplication in U2OS cells. Centrosome number was determined as described in (A) in U2OS cells transfected with DsRed-AZI (white bars), or in U2OS cells sequentially transfected with DsRed (black bars), OAZ miR\textsuperscript{DsRed} (gray bars), or DsRed-AZI followed by GFP-Mps1\textsuperscript{T468A}. GFP-Mps1\textsuperscript{T468A} does not accelerate the onset of reduplication on its own, but DsRed-AZI does.

Figure 7. Antizyme modulates the centrosomal accumulation of Mps1 in a T468-dependent manner.

(A) OAZ overexpression leads to reduced centrosomal accumulation of GFP-Mps1. (B) AZI overexpression leads to increased centrosomal accumulation of GFP-Mps1\textsuperscript{T468A}. (C) GFP-Mps1\textsuperscript{T468D} is insensitive to OAZ overexpression. (A-C) U2OS cells were sequentially transfected with DsRed, DsRed-OAZ, or DsRed-AZI, followed by (A) GFP-Mps1, (B) GFP-Mps1\textsuperscript{T468A}, or (C) GFP-Mps1\textsuperscript{T468D}, then arrested in S-phase and stained with an antibody against \(\gamma\)-Tubulin. The accumulation of GFP-Mps1 constructs at centrosomes was determined by visual inspection in 30-50 cells. Shown are representative cells displaying the majority phenotype, with the number in the upper right corner representing the percentage of cells showing the demonstrated phenotype; Hoechst (blue), GFP (green), DsRed (red); bar = 5 \(\mu\)m. Panels to the right show digitally magnified images of the box surrounding the centrosomes for each cell; GFP (green), DsRed (red), \(\gamma\)-Tubulin (cyan); bar = 1 \(\mu\)m.

Figure 8. Antizyme inhibits Mps1-dependent formation of procentrioles during the canonical duplication cycle.

HeLa cells were transfected with GFP, GFP-Mps1KD, or GFP-OAZ, enriched in G1 by a 48 hr serum starvation, then stimulated to enter S-phase by the addition of serum in the presence of BrdU. MCF7-GFPCentrin2 cells were transfected with DsRed or DsRed-OAZ followed by G0 arrest with a 48 hr serum starvation, then stimulated to enter the cell cycle by the addition of serum and growth factors in the presence of BrdU. HeLa cells were analyzed by IIF eight hrs after stimulation with antibodies against BrdU, HsSas-6, Centrin2, and CP110. MCF7-GFPCentrin2 cells were analyzed by IIF twelve hrs after stimulation with an antibody against BrdU. (A) OAZ and Mps1KD suppress centriole formation in HeLa and MCF7 cells. Graph shows the percentage of BrdU positive cells with two centrioles as judged by Centrin2 staining (HeLa) or GFPCentrin2 foci (MCF7). Values represent mean \(\pm\) standard deviation for 3 independent experiments where at least 100 BrdU positive cells were counted per replicate. (B-F) Left hand panels show representative BrdU-positive cells from the experiments described in (A) with
expression constructs and colors as indicated below, bar = 5 μm; right hand panels show digitally magnified images of centrioles, bar = 1μm. (B, C) Representative BrdU positive GFP-OAZ (B) or GFP-Mps1KD (C) expressing cells with two centrioles as judged by Centrin2 and CP110. (D) A representative BrdU positive cell expressing GFP alone with two centrioles and two procentrioles as judged by Centrin2 and HsSas-6, respectively. (E, F) Representative BrdU positive GFP-OAZ (E) or GFP-Mps1KD (F) expressing cells with two centrioles as judged by Centrin2 and a single HsSas-6-positive structure; (B, C) BrdU, blue; Cetn2, red; CP110, green. (D-F) BrdU, blue; Centrin2 (Cetn2), red; HsSas-6 (Sas-6), green.

**Figure 9. Overexpression of Mps1 negates the OAZ-mediated centriole-duplication defect.**

Asynchronously grown HeLa or HeLa-GFP-Mps1 cells were transfected with DsRed or DsRed-OAZ, BrdU was added for 4 hrs, and IIF was performed using antibodies against Centrin2. OAZ overexpression causes approximately a 10-fold increase in BrdU positive cells with two centrioles in HeLa cells, but has very little effect in HeLa-GFP-Mps1 cells. Bars represent the mean ± standard deviation in 3 independent experiments where at least 100 BrdU positive cells were counted per replicate.
Supplemental Information

Supplemental Methods

The following overlapping single stranded oligonucleotides were inserted into the miR\textsubscript{GFP} or miR\textsubscript{DsRed} plasmids to generate expression constructs with gene-specific synthetic miRs embedded in the 3’UTR of the GFP or DsRed mRNA:

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<td>AZI miR Bottom</td>
<td>CCTGTTTGCTATAGTATATCACTGCGGTTCAGTGCCACTG AAAACGCGAGTATATCATAGCAA</td>
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Supplemental Figure Legends

Supplemental Figure 1. Antizyme (OAZ) and its inhibitor (AZI) localize to centrosomes throughout the cell cycle in HeLa.

HeLa cells were grown in culture for 24 hrs and immunofluorescence was performed using antibodies against γ-tubulin (green) and either OAZ (A) or AZI (B) (red). Shown are representative interphase and mitotic cells (as determined by DNA staining, blue), with centrosomes indicated by arrows; bars = 5\(\mu\)m.

Supplemental Figure 2. Expression and localization of fluorescently tagged versions of AZI and OAZ.

(A) HeLa cells were analyzed by dual-color immunoblot with a mouse antibody against GFP (red) and a rabbit antibody against AZI (green) 24 hrs after transfection with GFP alone or GFP-AZI. The sizes of molecular weight markers (MW), which are visible in both channels using the LI-COR Odyssey scanner, are indicated in KDa (the 72 KDa marker is only visible in the green channel). Similar results were obtained using DsRed-AZI (not shown). (B) Identical U2OS cell lysates were analyzed on separate immunoblots with rabbit antibodies against DsRed or OAZ, as indicated, 24 hrs after transfection with DsRed alone or DsRed-OAZ. The sizes of molecular weight markers (MW) are indicated in KDa. Similar results were obtained using GFP and GFP-OAZ (not shown). (C) GFP-AZI and GFP-OAZ localize to centrosomes. HeLa cells were analyzed...
by IIF with an antibody against γ-tubulin 24 hrs after transfection with GFP, GFP-AZI, or GFP-OAZ; shown are representative cells expressing GFP, GFP-AZI, or GFP-OAZ, as indicated (bar = 5 µm) above digitally magnified images of the box surrounding the centrosome (bar = 1 µm); GFP, green; γ-tubulin, red; DNA, blue.

Supplemental Figure 3. miR-dependent depletion of OAZ and AZI.
(A and B) miR constructs reduce the AZI protein and the OAZ message by roughly 50%. Cells transfected with lacZ miR<sub>GFP</sub>, AZI miR<sub>GFP</sub>, or OAZ miR<sub>GFP</sub> were treated with 5 µg/ml blasticidin for three days to kill non-transfected cells. (A) The level of AZI protein was determined in pooled drug resistant cells as described in Figure 1. Numbers below the tubulin loading control indicate the relative normalized AZI levels. (B) The level of the OAZ and GAPDH messages were determined in pooled drug resistant cells by RT-PCR as described in Figure 1. Numbers below the GAPDH loading control indicate the relative normalized levels of the OAZ message. (C) OAZ miR<sub>GFP</sub> only partially depletes centrosomal OAZ. HeLa cells transfected with lacZ miR<sub>GFP</sub> or OAZ miR<sub>GFP</sub> were arrested in S-phase for 48h and analyzed by IIF with antibodies against OAZ (red) and γ-tubulin (green). Shown at left is a representative pair of OAZ miR<sub>GFP</sub>-expressing (GFP positive) and GFP negative cells showing GFP (green), OAZ (red), DNA (blue) and γ-Tubulin (cyan), with boxes surrounding the centrosomes and arrows indicating the corresponding panel at right for each cell; bar = 5 µm. Panels to the right are digitally magnified images of the boxes surrounding the centrosomes for each cell in the image at left; OAZ, red; γ-tubulin, green; bar = 1 µm. OAZ staining is reduced but not eliminated in GFP positive (GFP+) OAZ miR-expressing cells compared to adjacent GFP negative (GFP-) cells. (D) LacZ miR<sub>GFP</sub> does not affect centrosomal OAZ. Shown are magnified images of centrosomes from a representative lacZ miR<sub>GFP</sub>-expressing (GFP+) cell and its paired GFP- control cell; colors as in (A), bar = 1 µm. (E) AZI miR<sub>DsRed</sub> effectively depletes centriolar AZI. Shown at left is a representative pair of AZI miR<sub>DsRed</sub>-expressing (DsRed positive) and DsRed negative cells, showing DsRed (red), AZI (green), DNA (blue), and γ-tubulin (cyan), with boxes surrounding the centrosomes and arrows indicating the corresponding panel at right for each cell; bar = 5 µm. Panels to the right are digitally magnified images of the boxes surrounding the centrosomes for each cell in the image at left; AZI, green; γ-tubulin, red; bar = 1 µm. AZI can be detected in foci characteristic of centrioles (arrows) in DsRed negative (DsRed-) cells, but this staining is significantly reduced or absent in the DsRed positive (DsRed+) AZI miR<sub>DsRed</sub>-expressing cell (carets indicate centrosome position as judged by γ-tubulin staining in the DsRed+ cell).

Supplemental Figure 4. The form of Mps1 that co-immunoprecipitates with GFP-OAZ contains the degradation signal.
(A) GFP-OAZ binds to a single Mps1 isoform in 293 cells that contains the Mps1 degradation signal, but fails to bind to Mps1<sub>△12/13</sub> that lacks the Mps1 degradation signal. After transfection of 293-Mps1<sub>△12/13</sub> cells with GFP or GFP-OAZ, cells were arrested in S-phase with a 24 hr HU treatment. GFP-OAZ complexes were immunoprecipitated with a mouse antibody against GFP and analyzed by dual channel immunoblotting on the LI-COR Odyssey scanner with the M02 and MDS antibodies against Mps1 (see Fig. 5 for a description of these antibodies) and an antibody against GFP. In the presence of
doxycycline (+Dox) untagged Mps1<sup>Δ12/13</sup> (indicated by an arrowhead) is expressed at roughly 50% the level of endogenous full-length Mps1, but Mps1<sup>Δ12/13</sup> is not detectable in uninduced cells (-Dox) (second panel, “WB: M02”). The MDS antibody recognizes full-length endogenous Mps1 (closed arrow) and a 293-specific Mps1 isoform (open arrow) that migrates faster than full-length Mps1, but does not bind to Mps1<sup>Δ12/13</sup> (MDS; top panel). The same Mps1 isoform binds to GFP-OAZ in both uninduced cells (where Mps1<sup>Δ12/13</sup> is not present) and induced cells, and is recognized by the Mps1 Degradation Signal antibody that binds to the region of Mps1 that is missing in Mps1<sup>Δ12/13</sup> (MDS; top panel). The failure of the MDS antibody to bind to Mps1<sup>Δ12/13</sup> is most notable in the “Merge” panel.

(B-D) GFP-OAZ complexes prepared as described in (A) were analyzed by dual color immunoblotting on the LI-COR Odyssey scanner with the MDS, N1, C-19, and M02 antibodies (see Fig. 5 for a description of these antibodies). All blots were incubated with Alexa680-donkey anti-rabbit and IRDye800-donkey anti-mouse secondary antibodies. Shown are the signals from both the 680 and 800 channels (only immunoprecipitates are shown, see (A) for controls). (B) Shown is a blot incubated with MDS as the only primary antibody followed by both anti-rabbit and anti-mouse secondary antibodies. Asterisks indicate a non-specific band (presumably non-denatured IgG from the immunoprecipitation) generated by IRDye800 anti-mouse. (C) The co-precipitated form of Mps1 is also recognized by N1. (D) The co-precipitated form of Mps1 is additionally recognized by C19 and M02. Although the N1 and M02 signals are weak, the absence of signal from the anti-mouse antibody in panel (A) shows that these signal are due to the presence of the N1 or M02 antibody.

**Supplemental Figure 5. Modulating Antizyme activity does not affect the levels of cyclin D1 or cyclin A2.**

(A) Overexpression of OAZ or AZI has no detectable effect on Cyclin D1. HeLa cells were transfected with GFP, GFP-OAZ, or GFP-AZI and collected 48 hrs after transfection. Immunoblot was performed on cell lysates using antibodies against GFP, α-tubulin, and cyclin D1. Quantification using the Licor-Odyssey showed no difference in cyclin D1 levels after normalization to α-tubulin. (B) Depletion of OAZ or AZI has no detectable effect on Cyclin A2. HeLa cells were transfected with control (siCon), OAZ-specific (siOAZ), or AZI-specific (siAZI) siRNAs, and collected 48 hrs after transfection. Immunoblot was performed on cell lysates using antibodies against α-tubulin and cyclin A2. Quantification using the Licor-Odyssey showed no difference in cyclin A levels after normalization to α-tubulin.

**Supplemental Figure 6. Model for the Control of Mps1 Degradation by Antizyme.**

OAZ binds to the Mps1 degradation signal (D.S), and the Mps1-OAZ complex is targeted to the proteasome for degradation. Cdk2 phosphorylates Mps1 at T468 within the degradation signal, and we hypothesize that this phosphorylation prevents OAZ from binding to Mps1. We further hypothesize that both the Mps1-OAZ interaction and Mps1 degradation occur at the centrosome. OAZ might prevent both premature initiation of centrosome duplication and centrosome reduplication by removing Mps1 from centrosomes when Cdk2 activity is low. When Cdk2 levels rise, Mps1 is protected from degradation, allowing centrosome duplication to proceed. An as yet unidentified
phosphatase (indicated by a question mark) removes T468 phosphorylation, returning Mps1 to a state permissive for OAZ binding. Mps1 is represented by a white rectangle, with a black rectangle representing the Mps1 degradation signal (D.S; amino acids 420-507, encoded by exons 12 and 13 that are missing in Mps1\(^{\Delta12/13}\)), and a grey rectangle representing the kinase domain. OAZ is represented as a light grey shape that binds to the Mps1 degradation signal.
Figure 1. OAZ- and AZI-specific siRNAs deplete the centrosomal OAZ and AZI pools but do not affect the whole cell levels of Mps1.

(A) AZI siRNA results in an approximately 5-fold decrease in the AZI protein (relative levels are indicated below each lane), but neither AZI nor OAZ siRNA significantly affected Mps1 whole cell levels, as compared to control siRNA. Shown are immunoblots of siRNA transfected cells; Mps1 and the α-tubulin loading control were stained on the same blot and imaged using the LI-COR Odyssey, while AZI was stained on a separate blot and imaged with ECL. (B) RT-PCR of total RNA isolated from HeLa cells treated with control or OAZ siRNA shows the OAZ message is depleted by OAZ siRNA. The negative control (-RT) shows the product is not due to contaminating chromosomal DNA. Final concentrations of siRNA in µM used are listed above each lane. Lower panel shows amplification with GAPDH-specific primers as a loading control. (C, D) The centrosomal pools of OAZ and AZI are depleted by siRNA treatment. IIF of HeLa cells transfected with control (siControl), OAZ-specific (siOAZ), or AZI-specific (siAZI) siRNAs for 48 hours. Shown are representative cells stained with γ-tubulin (green), Mps1 (red), and Hoechst (blue); bar = 5 µm. Panels at the right of each cell show digitally magnified images of the box surrounding the centrosomes; bar = 1 µm.
Figure 2. Antizyme activity modulates centrosomal Mps1 in a proteasome-dependent manner.

(A) Increasing Antizyme activity by depleting its inhibitor AZI decreases centrosomal Mps1. (B) The loss of Mps1 from centrosomes in AZI-siRNA transfected cells is reversed by inhibiting the proteasome. (A,B) siRNA-transfected HeLa cells were arrested in S-phase with a 24 HU treatment, then treated with DMSO (A) or MG115 (B) for 4 hr. Shown are representative cells stained with γ-tubulin (green), Mps1 (red), and Hoechst (blue); bar = 5 µm. Panels to the right of each cell show digitally magnified images of the box surrounding the centrosomes; bar = 1 µm. (C) Decreasing Antizyme activity by siRNA depletion causes centrosome reduplication, along with a slight increase in Mps1 levels. siRNA-transfected HeLa cells were arrested in S-phase with a 24 HU treatment. Shown are representative cells stained with γ-tubulin (green), Mps1 (red), and Hoechst (blue); bar = 5 µm. Panels to the right of each cell show digitally magnified images of the box surrounding the centrosomes; bar = 1 µm. (D) Decreasing Antizyme activity by overexpressing GFP-AZI increases centrosomal Mps1 levels. HeLa cells were transfected with GFP-AZI, arrested in S-phase with a 24 hr HU treatment, then twenty-five cell pairs were imaged as described in materials and methods. Shown at left is a representative cell pair showing GFP-AZI (green), Mps1 (red), DNA (blue) and γ-Tubulin (cyan), with arrows indicating the corresponding panel at right for each cell; bar = 5 µm. The panels at right show magnified images of the box surrounding centrosomes for the cells in the left panel to illustrate centrosome position (γ-tubulin staining, cyan) and Mps1 levels (shown in pseudocolor to highlight differences between the two cells); arrow heads indicate centrosome position; bar = 1 µm.
Figure 3. Antizyme depletion causes Mps1 dependent Centrosome re-duplication in HeLa cells.

(A) OAZ siRNA caused an approximately 5-fold increase in cells with more than two centrosomes (and more than 4 centrioles). HeLa cells were transfected with the indicated siRNA then arrested in S-phase for 48 hr. (B) Centrosome re-duplication in OAZ-depleted cells requires Mps1. HeLa cells were sequentially transfected with control (black bars) or Mps1-specific siRNA (red bars) followed by lacZ miR<sup>GFP</sup> (lacZ miR), OAZ miR<sup>GFP</sup> (OAZ miR), or GFP-AZI (AZI), then arrested in S-phase for 48 hr. (A and B) Centrosome number was determined by IIF with antibodies against γ-Tubulin and centrin. Values represent the mean ± standard deviation for 3 independent experiments where at least 100 cells were counted per replicate. Mps1 depletion was verified in (B) by immunoblotting with antibodies against Mps1 and the α-Tubulin loading control, as shown below the bar graph.
Figure 4. Centrosomally localized Mps1-PACT is modulated by Antizyme and proteasome activities through the Mps1 degradation signal. (A and B) GFP-Mps1-PACT (A) and GFP-Mps1 Δ12/13-PACT (B) can only be detected at centrosomes. Shown are the GFP-Mps1-PACT or GFP-Mps1 Δ12/13-PACT signal (green) from representative S-phase arrested HeLa cells stained with Centrin (red), and Hoechst (blue); bar = 5 µm. Lower panels show digitally magnified images of the box surrounding the centrosomes for each cell; bar = 1 µm. (C) OAZ and proteasome activity modulate GFP-Mps1-PACT but not GFP-Mps1 Δ12/13-PACT that lacks the Mps1 degradation signal. HeLa cells were sequentially transfected with control (Con), OAZ-, or AZI-specific siRNAs, followed by GFP-Mps1-PACT (Mps1-PACT) or GFP-Mps1 Δ12/13-PACT (Mps1 Δ12/13-PACT), arrested in S-phase with a 24 hr HU treatment, then treated with either DMSO (black bars) or MG115 (gray bars) for 4 hrs. The percentage of cells in which a GFP signal could be detected was determined in 250-300 cells for each sample, and normalized to that in the respective DMSO-treated control siRNA sample.
Figure 5. Binding of Antizyme to Mps1 requires the Mps1 degradation signal.
OAZ binds to Mps1 in HeLa cells, but fails to bind to a form of Mps1 lacking the degradation signal. (A) GFP-OAZ co-immunoprecipitates with Mps1. HeLa cells were transfected with either GFP or GFP-OAZ, then arrested in S-phase. Mps1 complexes were immunoprecipitated from lysates with the Mps1 N1 antibody, and analyzed by dual color immunoblotting with rabbit antibodies against GFP or Mps1. Input (I) shows 2.5% of the lysate used in the preclear (PC, showing any proteins that bind to beads independently of antibody) and immunoprecipitation (IP). (B) DsRed-OAZ (open arrow) binds to endogenous Mps1 (closed arrow) but not GFP-Mps1Δ12/13 (arrowhead). HeLa-GFP-Mps1Δ12/13 cells induced with doxycycline were transfected with DsRed-OAZ, then arrested in S-phase. Mps1 complexes were immunoprecipitated from lysates with three different antibodies and analyzed by dual color immunoblotting with antibodies against Mps1 (N1) and DsRed. Input (I) shows 2.5% of the total lysate used in the preclear (PC) and immunoprecipitations; PC shows proteins bound to beads in the absence of antibody, while N1, MDS, and GFP shows proteins that immunoprecipitate with those antibodies. (C) Diagram of Mps1 and Mps1Δ12/13 showing the location of the Mps1 degradation signal and the epitopes for the antibodies used in (A) and (B).
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Asynchronously grown HeLa or HeLa-GFPMps1 cells were transfected with DsRed or DsRed-OAZ, BrdU was added for 4 hrs, and IIF was performed using antibodies against Centrin2. OAZ overexpression causes approximately a 10-fold increase in BrdU positive cells with two centrioles in HeLa cells, but has very little effect in HeLa-GFPMps1 cells. Bars represent the mean ± standard deviation in 3 independent experiments where at least 100 BrdU positive cells were counted per replicate.
Supplemental Figure 1. Antizyme (OAZ) and its inhibitor (AZI) localize to centrosomes throughout the cell cycle in HeLa.

HeLa cells were grown in culture for 24 hrs and immunofluorescence was performed using antibodies against γ-tubulin (green) and either OAZ (A) or AZI (B) (red). Shown are representative interphase and mitotic cells (as determined by DNA staining, blue), with centrosomes indicated by arrows; bars = 5µm.
Supplemental Figure 2. Expression and localization of fluorescently tagged versions of AZI and OAZ.

(A) HeLa cells were analyzed by dual-color immunoblot with a mouse antibody against GFP (red) and a rabbit antibody against AZI (green) 24 hrs after transfection with GFP alone or GFP-AZI. The sizes of molecular weight markers (MW), which are visible in both channels using the LI-COR Odyssey scanner, are indicated in KDa (the 72 KDa marker is only visible in the green channel). Similar results were obtained using DsRed-AZI (not shown). (B) Identical U2OS cell lysates were analyzed on separate immunoblots with rabbit antibodies against DsRed or OAZ, as indicated, 24 hrs after transfection with DsRed alone or DsRed-OAZ. The sizes of molecular weight markers (MW) are indicated in KDa. Similar results were obtained using GFP and GFP-OAZ (not shown). (C) GFP-AZI and GFP-OAZ localize to centrosomes. HeLa cells were analyzed by IIF with an antibody against γ-tubulin 24 hrs after transfection with GFP, GFP-AZI, or GFP-OAZ; shown are representative cells expressing GFP, GFP-AZI, or GFP-OAZ, as indicated (bar = 5 µm) above digitally magnified images of the box surrounding the centrosome (bar = 1 µm); GFP, green; γ-tubulin, red; DNA, blue.
Supplemental Figure 3. miR-dependent depletion of OAZ and AZI.

(A and B) miR constructs reduce the AZI protein and the OAZ message by roughly 50%. Cells transfected with lacZ miR<sup>GFP</sup>, AZI miR<sup>GFP</sup>, or OAZ miR<sup>GFP</sup> were treated with 5 µg/ml blasticidin for three days to kill non-transfected cells. (A) The level of AZI protein was determined in pooled drug resistant cells as described in Figure 1. Numbers below the tubulin loading control indicate the relative normalized AZI levels. (B) The level of the OAZ and GAPDH messages were determined in pooled drug resistant cells by RT-PCR as described in Figure 1. Numbers below the GAPDH loading control indicate the relative normalized levels of the OAZ message. (C) OAZ miR<sup>GFP</sup> only partially depletes centrosomal OAZ. HeLa cells transfected with lacZ miR<sup>GFP</sup> or OAZ miR<sup>GFP</sup> were arrested in S-phase for 48h and analyzed by IIF with antibodies against OAZ (red) and γ-Tubulin (green). Shown at left is a representative pair of OAZ miR<sup>GFP</sup>-expressing (GFP positive) and GFP negative cells showing GFP (green), OAZ (red), DNA (blue) and γ-Tubulin (cyan), with boxes surrounding the centrosomes and arrows indicating the corresponding panel at right for each cell; bar = 5 µm. Panels to the right are digitally magnified images of the boxes surrounding the centrosomes for each cell in the image at left; OAZ, red; γ-Tubulin, green; bar = 1 µm. OAZ staining is reduced but not eliminated in GFP positive (GFP+) OAZ miR-expressing cells compared to adjacent GFP
negative (GFP-) cells. (D) LacZ miR<sup>GFP</sup> does not affect centrosomal OAZ. Shown are magnified images of centrosomes from a representative lacZ miR<sup>GFP</sup>-expressing (GFP+) cell and its paired GFP- control cell; colors as in (A), bar = 1 µm. (E) AZI miR<sup>DsRed</sup> effectively depletes centriolar AZI. Shown at left is a representative pair of AZI miR<sup>DsRed</sup>-expressing (DsRed positive) and DsRed negative cells, showing DsRed (red), AZI (green), DNA (blue), and γ-tubulin (cyan), with boxes surrounding the centrosomes and arrows indicating the corresponding panel at right for each cell; bar = 5 µm. Panels to the right are digitally magnified images of the boxes surrounding the centrosomes for each cell in the image at left; AZI, green; γ-tubulin, red; bar = 1 µm. AZI can be detected in foci characteristic of centrioles (arrows) in DsRed negative (DsRed-) cells, but this staining is significantly reduced or absent in the DsRed positive (DsRed+) AZI miR<sup>DsRed</sup>-expressing cell (carets indicate centrosome position as judged by γ-tubulin staining in the DsRed+ cell).
Supplemental Figure 4. The form of Mps1 that co-immunoprecipitates with GFP-OAZ contains the degradation signal.

(A) GFP-OAZ binds to a single Mps1 isoform in 293 cells that contains the Mps1 degradation signal, but fails to bind to Mps1\(^{\Delta 12/13}\) that lacks the Mps1 degradation signal. After transfection of 293-Mps1\(^{\Delta 12/13}\) cells with GFP or GFP-OAZ, cells were arrested in S-phase with a 24 hr HU treatment. GFP-OAZ complexes were immunoprecipitated with a mouse antibody against GFP and analyzed by dual channel immunoblotting on the LI-COR Odyssey scanner with the M02 and MDS antibodies against Mps1 (see Fig. 5 for a description of these antibodies) and an antibody against GFP. In the presence of doxycycline (+Dox) untagged Mps1\(^{\Delta 12/13}\) (indicated by an arrowhead) is expressed at roughly 50% the level of endogenous full-length Mps1, but Mps1\(^{\Delta 12/13}\) is not detectable in uninduced cells (-Dox) (second panel, “WB: M02”). The MDS antibody recognizes full-length endogenous Mps1 (closed arrow) and a 293-specific Mps1 isoform (open arrow) that migrates faster than full-length Mps1, but does not bind to Mps1\(^{\Delta 12/13}\) (MDS; top panel). The same Mps1 isoform binds to GFP-OAZ in both uninduced cells (where Mps1\(^{\Delta 12/13}\) is not present) and induced cells, and is recognized by the Mps1 Degradation
Signal antibody that binds to the region of Mps1 that is missing in Mps1Δ12/13 (MDS; top panel). The failure of the MDS antibody to bind to Mps1Δ12/13 is most notable in the “Merge” panel.

(B-D) GFP-OAZ complexes prepared as described in (A) were analyzed by dual color immunoblotting on the LI-COR Odyssey scanner with the MDS, N1, C-19, and M02 antibodies (see Fig. 5 for a description of these antibodies). All blots were incubated with Alexa680-donkey anti-rabbit and IRDye800-donkey anti-mouse secondary antibodies. Shown are the signals from both the 680 and 800 channels (only immunoprecipitates are shown, see (A) for controls). (B) Shown is a blot incubated with MDS as the only primary antibody followed by both anti-rabbit and anti-mouse secondary antibodies. Asterisks indicate a non-specific band (presumably non-denatured IgG from the immunoprecipitation) generated by IRDye800 anti-mouse. (C) The co-precipitated form of Mps1 is also recognized by N1. (D) The co-precipitated form of Mps1 is additionally recognized by C19 and M02. Although the N1 and M02 signals are weak, the absence of signal from the anti-mouse antibody in panel (A) shows that these signal are due to the presence of the N1 or M02 antibody.
Supplemental Figure 5. Modulating Antizyme activity does not affect the levels of cyclin D1 or cyclin A2.

(A) Overexpression of OAZ or AZI has no detectable effect on Cyclin D1. HeLa cells were transfected with GFP, GFP-OAZ, or GFP-AZI and collected 48 hrs after transfection. Immunoblot was performed on cell lysates using antibodies against GFP, α-tubulin, and cyclin D1. Quantitation using the Licor-Odyssey showed no difference in cyclin D1 levels after normalization to α-tubulin. (B) Depletion of OAZ or AZI has no detectable effect on Cyclin A2. HeLa cells were transfected with control (siCon), OAZ-specific (siOAZ), or AZI-specific (siAZI) siRNAs, and collected 48 hrs after transfection. Immunoblot was performed on cell lysates using antibodies against α-tubulin and cyclin A2. Quantitation using the Licor-Odyssey showed no difference in cyclin A levels after normalization to α-tubulin.
Supplemental Figure 6. Model for the Control of Mps1 Degradation by Antizyme.
OAZ binds to the Mps1 degradation signal (D.S), and the Mps1-OAZ complex is targeted to the proteasome for degradation. Cdk2 phosphorylates Mps1 at T468 within the degradation signal, and we hypothesize that this phosphorylation prevents OAZ from binding to Mps1. We further hypothesize that both the Mps1-OAZ interaction and Mps1 degradation occur at the centrosome. OAZ might prevent both premature initiation of centrosome duplication and centrosome reduplication by removing Mps1 from centrosomes when Cdk2 activity is low. When Cdk2 levels rise, Mps1 is protected from degradation, allowing centrosome duplication to proceed. An as yet unidentified phosphatase (indicated by a question mark) removes T468 phosphorylation, returning Mps1 to a state permissive for OAZ binding. Mps1 is represented by a white rectangle, with a black rectangle representing the Mps1 degradation signal (D.S; amino acids 420-507, encoded by exons 12 and 13 that are missing in Mps1Δ12/13), and a grey rectangle representing the kinase domain. OAZ is represented as a light grey shape that binds to the Mps1 degradation signal.