Functional genomics identifies a requirement of pre-mRNA splicing factors for sister chromatid cohesion

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Abstract

Sister chromatid cohesion mediated by the cohesin complex is essential for chromosome segregation during cell division. Using functional genomic screening, we identify a set of 26 pre-mRNA splicing factors that are required for sister chromatid cohesion in human cells. Loss of spliceosome subunits increases the dissociation rate of cohesin from chromatin and abrogates cohesion after DNA replication, ultimately causing mitotic catastrophe. Depletion of splicing factors causes defective processing of the pre-mRNA encoding sororin, a factor required for the stable association of cohesin with chromatin, and an associated reduction of sororin protein level. Expression of an intronless version of sororin and depletion of the cohesin release protein WAPL suppress the cohesion defect in cells lacking splicing factors. We propose that spliceosome components contribute to sister chromatid cohesion and mitotic chromosome segregation through splicing of sororin pre-mRNA. Our results highlight the loss of cohesion as an early cellular consequence of compromised splicing. This may have clinical implications because SF3B1, a splicing factor that we identify to be essential for cohesion, is recurrently mutated in chronic lymphocytic leukaemia.

Keywords chromosome; cohesin; cohesion; mitosis; splicing

Subject Categories DNA Replication, Repair & Recombination; RNA Biology

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Introduction

The correct partitioning of sister genomes during cell division requires that sister kinetochores attach to microtubules emanating from opposite spindle poles. To facilitate this, sister chromatids are held together from their synthesis during DNA replication until their disjunction by a phenomenon called sister chromatid cohesion (Gucci et al., 1994; Nasmyth & Haering, 2009). Sister chromatid cohesion is mediated by a ring-shaped tripartite protein complex called cohesin that is composed of the core subunits SMC3, SMC1 and SCC1 and that is thought to topologically entrap sister chromatids (Haering et al., 2008).

A number of accessory factors regulate the function and properties of cohesin throughout the cell cycle. Cohesin loading onto chromatins happens during telophase/G1 and is performed by a cohesin-loading complex composed of SCC2 and SCC4 proteins (Ciosk et al., 2000). At this stage, cohesin interacts with chromatin transiently (Gerlich et al., 2006). A stable cohesin–chromatin liaison is prevented by the activity of the WAPL–PDSS protein complex (Tanaka et al., 2001; Gandhi et al., 2006; Kueng et al., 2006; Sutani et al., 2009). The cohesin release protein WAPL is thought to remove cohesin rings from chromatin by opening the SMC3–SCC1 interface of the complex (Chan et al., 2012; Buheitel & Stemmann, 2013; Eichinger et al., 2013). The subsequent establishment of sister chromatid cohesion requires DNA replication (Uhlmann & Nasmyth, 1998) as well as the opposition of WAPL–PDSS activity by the acetylation of cohesin’s SMC3 subunit at the hands of the acetyltransferases ESCO1/2 (Rolf Ben-Shahar et al., 2008; Unal et al., 2008; Zhang et al., 2008; Rowland et al., 2009; Chan et al., 2012). In vertebrate cells, sister chromatid cohesion and the stable association of cohesin with chromatin after DNA replication also require a protein called sororin (Rankin et al., 2005; Schmitz et al., 2007). DNA replication and SMC3 acetylation promote the binding of sororin to PDSS leading to the displacement of WAPL (Lafont et al., 2010; Nishiyama et al., 2010). By antagonizing the cohesin release activity of WAPL, sororin is thought to ensure the long-lived links that hold sister chromatids together until mitosis (Gerlich et al., 2006; Schmitz et al., 2007; Lafont et al., 2010; Nishiyama et al., 2010).

In vertebrate cells, cohesin is removed in two discrete steps during mitosis (Waizenegger et al., 2000). The first step takes place during
prophase and requires WAPL, as well as the phosphorylation of cohesin subunits and regulators by mitotic kinases (Losada et al., 2002; Sumara et al., 2002; Gimenez-Abian et al., 2004; Hauf et al., 2005; Kueng et al., 2006; Nishiyama et al., 2013; Tedeschi et al., 2013). This process, called the prophase pathway, removes cohesin from chromosome arms in a proteolysis-independent manner. Centromeric cohesin is rendered immune to the prophase pathway by the activity of the shugoshin protein Sgo1 (Kitajima et al., 2004; Salic et al., 2004; McGuinness et al., 2005). This maintains the connection between sister chromatids at the centromere and allows the correct attachment of chromosomes to the mitotic spindle. At anaphase onset, the remaining centromeric cohesin complexes are removed from chromatin through proteolytic cleavage of the SCC1 subunit leading to the segregation of sister chromatids towards opposite poles (Uhlmann et al., 2000; Hauf et al., 2001; Oliveira et al., 2010).

In addition to its function in chromosome segregation, cohesin plays important roles in other nuclear processes including the DNA damage response and DNA repair (Kim et al., 2002; Yazdi et al., 2002; Strom et al., 2004; Unal et al., 2004; Watrin & Peters, 2009), chromatin organization and gene regulation. Cohesin facilitates transcriptional termination between convergent genes in yeast (Gullerova & Proudfoot, 2008) and controls the developmentally regulated expression of genes in multiple systems (Rollins et al., 2004; Horsfield et al., 2007; Landeira et al., 2009; Pauli et al., 2010). Furthermore, cohesin acts in concert with the insulator protein CTCF (CCCTC binding factor) to promote long-range intra-chromatid interactions that govern gene expression (Pareloho et al., 2008; Rubio et al., 2008; Stedman et al., 2008; Wendt et al., 2008; Hadjur et al., 2009; Sofueva et al., 2013; Zuin et al., 2014). Thus, cohesin has emerged as a key regulator of gene expression.

Conversely, increasing evidence suggests that gene expression processes impact on the function and properties of cohesin. In budding yeast, cohesin concentrates in regions of convergent transcription (Lengronne et al., 2004). SRM160, a protein belonging to the SR family of pre-mRNA splicing regulators, was found to associate with core cohesin subunits (Mccracken et al., 2005). Also, depletion of UAP56 and URH49 helicases, members of the TREX complex that couple transcription and pre-mRNA processing with mRNA export (Lee & Tarn, 2013), led to alterations in cohesion in mitotic cells (Yamazaki et al., 2010).

In a search for novel factors controlling mitosis in human cells, we identify an essential and widespread role for pre-mRNA splicing factors in sister chromatid cohesion and provide a mechanistic explanation for their involvement during mitosis.

**Results**

**A screen for potential new mitotic regulators identifies MFAP1**

Genome-wide RNAi screens for cell cycle regulators (Mukherji et al., 2006; Kittler et al., 2007; Conery & Harlow, 2010; Neumann et al., 2010) and the proteomic analysis of the mammalian midbody (Skop et al., 2004) have highlighted candidate proteins that could represent novel factors involved in cell division. Using published data sets from these approaches, we compiled a list of 718 human candidate genes (Supplementary Fig S1A and Supplementary Table S1) for rescoring by RNA interference-mediated gene depletion in HeLa Kyoto cervical carcinoma cells. Established regulators of cell division and suspected pseudogenes were removed from the list (See Materials and Methods section for gene selection details). The screen utilized an arrayed siRNA library containing pools of four small interfering RNA (siRNA) duplexes for each of the 718 genes. We employed a fluorescence microscopy-based end point assay to detect nuclear defects in interphase cells that could be indicative for defects in cell division. In brief, HeLa Kyoto cells were transfected with siRNA pools in 96-well plates, grown for 52 h, fixed, labelled with a DNA dye and whole-cell stain, and subjected to automated image acquisition. Image data from three screen replicates were scored for the fraction of interphase cells displaying nuclear abnormalities, including multi-nucleation and nuclear fragmentation (Supplementary Fig S1B and Supplementary Table S1). siRNA pools targeting positive control genes, such as the essential cytokinesis regulator ECT2 (Yuce et al., 2005), scored with high phenotypic penetrance indicating effective target depletion in the screen (Supplementary Fig S1B and C).

Fifty candidate genes whose siRNA pools induced abnormal nuclear morphology in more than 20% of interphase cells were selected for further analysis. The selected genes were subjected to deconvolution analysis in which cells were transfected with the individual siRNA duplexes that constituted the pool of four duplexes used in the primary screen. For seven genes, two or more out of four siRNA duplexes recapitulated the phenotype observed in the primary screen (Supplementary Table S2). Transfection of the siRNA pool and of all four individual siRNA duplexes targeting the gene MFAP1 microfibrillar-associated protein 1) caused severe nuclear fragmentation characterized by the formation of small and large karyomeres and an increase in DNA content (Fig 1A and Supplementary Fig S1C). Consistent with an on-target effect, we found that the 4 siRNA duplexes also decreased MFAP1 protein levels (Fig 1A). MFAP1 siRNA #3 was selected for further analyses. MFAP1 is a conserved 52 kDa nuclear protein that has been purified in human spliceosomal fractions (Jurica & Moore, 2003). The Drosophila orthologue of MFAP1 associates with factors of the spliceosomal tri-small nuclear ribonucleoprotein (tri-snRNP) complex and has been implicated in pre-mRNA processing (Andersen & Tapon, 2008). The nuclear defects observed upon depletion of MFAP1 in human cells (Fig 1A) raise the possibility that this splicing factor is required for the segregation of chromosomes during cell division.

**Loss of MFAP1 prevents chromosome alignment and causes a spindle assembly checkpoint-dependent mitotic arrest**

To investigate the effect of MFAP1 loss on cell division, we first measured the duration of mitosis by tracking cells using bright-field microscopy. Control cells entered mitosis and underwent the metaphase-to-anaphase transition within 46 min on average (Fig 1B). Subsequently, control cells underwent cytokinesis and exited mitosis. In contrast, the majority of cells lacking MFAP1 remained arrested in mitosis for several hours before undergoing cell death or aberrantly exiting mitosis without attempting cytokinesis (Fig 1B). The increased duration of mitosis observed in cells depleted of MFAP1 was abrogated upon co-depletion of the spindle assembly checkpoint component Mad2 (Lara-Gonzalez et al., 2012) (Fig 1C). These results indicate that loss of MFAP1 causes a mitotic arrest that is dependent on the spindle assembly checkpoint.
The spindle assembly checkpoint dependency of the arrest in MFAP1-depleted cells suggested that chromosomes might fail to bi-orient on the mitotic spindle. To test this, we used live imaging of cells co-expressing histone H2B-mCherry and α-tubulin-EGFP. Control cells achieved chromosome alignment at the metaphase plate within an hour before initiating anaphase and exiting mitosis (8/9 cells) (Fig 1D). In contrast, cells lacking MFAP1 were unable to form a metaphase plate upon mitotic entry and remained arrested in mitosis with a scattered chromosome configuration (9/9 cells) (Fig 1D). To corroborate this observation, we treated cells with the proteasomal inhibitor MG132 that prevents anaphase onset and arrests cells in metaphase. Depletion of MFAP1 completely abolished...
the chromosome alignment that was observed in the majority of control siRNA transfected cells after a 3-h arrest in MG132 (Fig 1E). These results indicate that MFAP1 is required for the alignment of chromosomes at the metaphase plate. This function of MFAP1 could contribute to the prometaphase arrest and mitotic catastrophe observed in cells lacking the protein.

**MFAP1 is required for sister chromatid cohesion**

A failure to align chromosomes could be caused by defects in mitotic spindle dynamics, kinetochore function or sister chromatid cohesion. Loss of sister chromatid cohesion prevents the stable attachment of microtubules to kinetochores and the bi-orientation of chromatids (Tanaka et al., 2000). We observed that centromeres in MFAP1-depleted prometaphase cells appeared to lack the pairwise arrangement normally found in control cells (Fig 1E). Therefore, we decided to test the status of sister chromatid cohesion in nocodazole-arrested mitotic cells by chromosome spreading. Chromosomes from control siRNA-transfected cells were connected at the central constriction reflecting intact cohesion between sister centromeres (Fig 2A). In contrast, the vast majority of chromosomes from MFAP1-depleted cells had lost their characteristic X shape and instead appeared as single chromatids (Fig 2A). Fluorescence *in situ* hybridization (FISH) experiments confirmed the loss of sister chromatid cohesion upon depletion of MFAP1 in intact mitotic cells (Fig 2B). These results suggest that MFAP1 is required for sister chromatid cohesion in mitosis. Remarkably, the severity of the sister chromatid cohesion loss phenotype in MFAP1-depleted cells was comparable to the loss of the centromeric cohesion protector SGOL1 (Fig 2A). To test whether loss of MFAP1 protein is responsible for the observed defects, we generated a cell line stably expressing a transgenic and siRNA-resistant version of MFAP1 that was tagged with AcGFP (*Aequora coeruleus* green fluorescent protein) and a FLAG epitope (AcFL-MFAP1-r) at a level close to the endogenous counterpart (Fig 2C, right panel). Expression of the RNAi-resistant transgene suppressed both the mitotic loss of sister chromatid cohesion and the interphase nuclear defect in cells transfected with the corresponding siRNA duplex targeting MFAP1 (Fig 2C). Thus, we have identified a role for the splicing factor MFAP1 in sister chromatid cohesion, the crucial connection between DNA copies that allows the bi-orientation and subsequent accurate segregation of chromosomes in mitosis.

**A multitude of spliceosome components are required for sister chromatid cohesion in human cells**

The requirement of MFAP1 for sister chromatid cohesion raised the question whether MFAP1 has a dedicated second role in cohesion independent of pre-mRNA splicing or whether there is a much more widespread requirement of spliceosome components for sister chromatid cohesion. Analysis of genome-wide RNAi screen data sets revealed an enrichment of spliceosome components in the category of genes whose depletion resulted in mitotic defects (Kittler et al., 2007; Hofmann et al., 2010; Neumann et al., 2010). A comparison of the proteomic composition of the human spliceosome (Zhou et al., 2002; Jurica & Moore, 2003) with genes classified as mitosis-defective in a genome-wide screen (Neumann et al., 2010) revealed an overlap of 33 factors. By transfecting siRNA pools and performing chromosome spreading analysis, we tested whether depletion of these 33 proteins had an effect on sister chromatid cohesion. Transfection of 26 siRNA pools targeting splicing genes resulted in a loss of sister chromatid cohesion in more than half of the cells (Fig 3A and B; for protein depletion data see Supplementary Fig S3A). Remarkably, siRNA pools targeting 18 splicing factors caused an almost complete disruption of sister chromatid cohesion that was comparable to the loss of SGOL1 or the loss of the core cohesion subunit SCC1 (Fig 3A and B). These included siRNA pools targeting SF3B1 (splicing factor 3B subunit 1), NHP2L1 (non-histone protein 2-like protein 1), SART1 (squamous cell carcinoma antigen recognized by T cells) and CDC5L (cell division cycle 5-like protein). Deconvolution experiments using individual siRNA duplexes for splicing factors whose siRNA pools caused the most severe defects in sister chromatid cohesion supported a causal relationship between loss of the target protein and the observed chromosomal phenotype (Supplementary Fig S2). Stable transgenic expression of tagged and RNAi-resistant versions of the spliceosome components NHP2L1 and SART1 fully restored sister chromatid cohesion in cells depleted of the endogenous counterparts (Fig 3C and Supplementary Fig S3B). Transfection of siRNA pools targeting MFAP1, NHP2L1 and SART1 also caused defects in sister chromatid cohesion in the human diploid colon carcinoma cell line HCT116 (Supplementary Fig S4) suggesting a general requirement of these splicing factors for cohesion in human cells.

Our observations demonstrate that the connection between pre-mRNA splicing and sister chromatid cohesion is not restricted to MFAP1 but extends well beyond. They suggest a widespread association of spliceosome component depletion with a loss of sister chromatid cohesion in human cells. Splicing proteins, whose depletion had a severe impact on sister chromatid cohesion, act in different subcomplexes and at different steps within the spliceosome assembly pathway (Supplementary Fig S5). SF3B1 is an essential component of the spliceosomal U2 snRNP complex and helps to target the complex to the branch point of introns in the initial phase of the splicing reaction (Wahl et al., 2009; Folco et al., 2011). NHP2L1 and SART1 are both components of the US and U4/U6 tri-snRNP complex, which associates with the spliceosome in a subsequent step (Makarova et al., 2001). CDC5L is a subunit of the Prp19 complex that is required for the activation of the spliceosome before the first step of the splicing reaction (Ajuh et al., 2000; Makarova et al., 2004; Grote et al., 2010). The fact that spliceosome components from distinct subcomplexes that act at different steps of spliceosome assembly are required for cohesion suggests that compromised pre-mRNA splicing in general could be responsible for the loss of sister chromatid cohesion. To investigate the link between splicing and sister chromatid cohesion, we decided to focus on the analysis of cells lacking the spliceosome components MFAP1, NHP2L1, SART1 and CDC5L.

**Loss of spliceosome components disrupts sister chromatid cohesion in interphase**

We first aimed to determine whether the loss of sister chromatid cohesion upon depletion of splicing factors occurred already during interphase or only upon entry into mitosis. To assess the status of sister chromatid cohesion in interphase, we used FISH probes for the *tff1* locus on chromosome 21 in post-replicative cells (Schmitz et al., 2007). Depletion of NHP2L1, SART1 and MFAP1 increased the
distance between sister chromatid signals to a similar extent as depletion of sororin, a protein essential for sister chromatid cohesion in interphase (Rankin et al., 2005; Schmitz et al., 2007) (Fig 4A). In contrast, loss of SGOL1, a protein essential only for maintaining centromeric cohesion upon entry into mitosis (Salic et al., 2004; McGuinness et al., 2005), did not alter the distance between sister chromatid signals (Fig 4A). These results indicate that the loss of splicing factors disrupts sister chromatid cohesion in interphase. This defect could then result and culminate in the emergence of single chromatids in mitotic chromosome spreads.

**Figure 2. MFAP1 is required for sister chromatid cohesion in mitosis.**

A Representative images of chromosome spreads (left) and quantification of the different states of sister chromatid cohesion (right) in cells that were transfected with the indicated siRNA duplexes 52 h prior to the analysis (n = 100 cells). Depletion of SGOL1 was used as a positive control for premature loss of sister chromatid cohesion.

B Fluorescence in situ hybridization (FISH) analysis performed using centromeric probes for chromosome 6 (green) and chromosome 8 (red) in cells transfected with the indicated siRNAs 48 h prior to analysis. Quantification of the number of centromere pairs that are more than 2 μm apart and were classified as split is shown (n > 30 cells).

C Chromosome spread analysis (52 h post-siRNA transfection) and IF analysis of interphase cells (72 h post-siRNA transfection) of cell lines stably expressing the AcFL tag alone or an siRNA-resistant version of MFAP1 (AcFL-MFAP1-r) following transfection with MFAP1 siRNA (left) (n > 100 cells, 3 experiments). Immunoblot of whole-cell extracts from the stable cell lines transfected with MFAP1 siRNA (+) or control siRNA (−) 72 h prior to analysis. Endogenous MFAP1 (filled arrowhead) and transgenic AcFL-MFAP1-r (open arrowhead) are indicated.

Data information: Scale bars, 10 μm.

Source data are available online for this figure.

**Spliceosome components are dispensable for cohesin loading onto chromatin and SMC3 acetylation**

To define the molecular mechanism that underlies the link between splicing and sister chromatid cohesion, we scrutinized the properties of cohesin components in interphase nuclei. To interrogate the loading of cohesin onto chromatin, we quantified the signal intensity of extraction-resistant SCC1 staining in fixed interphase nuclei (Schmitz et al., 2007). To facilitate comparative analysis and provide an internal experimental standard, we mixed untreated cells that...
were labelled with H2B-mCherry and unlabelled cells that were transfected with siRNA duplexes. Although depletion of the cohesin-loading factor SCC4 (Ciosk et al, 2000; Watrin et al, 2006) reduced the intensity of extraction-resistant SCC1, depletion of the splicing proteins NHP2L1, MFAP1 and SART1 had no discernible effect (Fig 4B). Thus, loss of spliceosome components does not interfere with the bulk loading of cohesin onto chromatin in interphase cells.

Acetylation of the core cohesin subunit SMC3 acts as a marker for the establishment of sister chromatid cohesion during DNA replication (Rolef Ben-Shahar et al, 2008; Unal et al, 2008; Zhang et al, 2008; Rowland et al, 2009; Nishiyama et al, 2010). Depletion of the spliceosome components NHP2L1, MFAP1, SART1 and CDC5L caused a minor reduction of SMC3 acetylation in replicating cells (Fig 4C). In contrast, loss of the acetyltransferases ESCO1 and ESCO2 (Hou & Zou, 2005) largely abolished SMC3 acetylation (Fig 4C). The minor reduction of SMC3 modification cannot explain the dramatic loss of sister chromatid cohesion detected in cells lacking splicing components. Even the sevenfold reduction in

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**Figure 3. Identification of spliceosome components that are required for sister chromatid cohesion.**

A Status of sister chromatid cohesion in HeLa Kyoto cells transfected with siRNA pools targeting the indicated spliceosome components 52 h before analysis (n = 100 cells). Cohesion status was determined by chromosome spreading.

B Chromosome spread images corresponding to the data shown in (A). Scale bar, 10 μm.

C Immunoblot analysis of whole-cell extracts prepared from cell lines stably expressing the AcFL tag or an siRNA-resistant NHP2L1 (AcFL-NHP2L1-r). Extracts were prepared 72 h after transfection with control (−) or NHP2L1 siRNA. Endogenous NHP2L1 (filled arrowhead) and transgenic AcFL-NHP2L1-r (open arrowhead) are indicated. Percentages of cells exhibiting loss of sister chromatid cohesion are indicated (n = 100 cells each, mean of three independent experiments).

D Transgenic cell lines stably expressing AcFL or an siRNA-resistant version of SART1 (AcFL-SART1-r) were transfected with control (−) and SART1 siRNA (+) and analysed as in (C).

Source data are available online for this figure.
SMC3 acetylation observed in cells co-depleted of ESCO1 and ESCO2 is not associated with a comparably penetrant disruption of cohesion (data not shown). Our results indicate that cells lacking spliceosome components lose sister chromatid cohesion during interphase despite loading cohesin onto chromatin and the occurrence of SMC3 acetylation, an indicator for cohesion establishment.

**Figure 4. Depletion of splicing factors abrogates sister chromatid cohesion in interphase despite cohesin loading and SMC3 acetylation.**

A Interphase FISH of G2 synchronized HeLa Kyoto cells that were transfected with control siRNA or siRNA duplexes targeting MFAP1, 53 h prior to analysis, sororin, SGOL1 or NHP2L1 (29 h prior to analysis). FISH was performed with a probe specific for the tff1 locus on trisomic chromosome 21 (yellow). DNA was stained with DAPI (blue). Magnified images of single pairs of FISH signals are displayed in the insets. Graph depicts the distance between the paired FISH signals measured in each of the indicated siRNA treatments. Bars represent mean ± SEM. Asterisks indicate a significant difference according to Student’s t-test (P < 0.05) (n = 30 paired signals). Scale bar, 10 µm.

B IF analysis of extraction-resistant SCC1 in H2B-mCherry-expressing cells that were untreated or in mCherry-negative cells that were transfected with the indicated siRNA duplexes (white arrowheads). Graph shows SCC1 fluorescence intensity quantification normalized to DAPI intensity and expressed as ratio between untransfected and siRNA transfected cells. Bars represent mean ± SD (n > 25 cells). Scale bar, 10 µm.

C Immunoblot analysis of whole-cell extracts from S phase synchronized cells that were previously transfected with sororin or NHP2L1 siRNAs (for 33 h) or control, MFAP1, SART1 or CDC5L siRNAs (for 51 h). The ratio of SMC3 acetylated at Lys105 and Lys106 (AcSMC3), and total SMC3 was measured by fluorescent detection and normalized to the ratio determined for cells that were transfected with control siRNA. α-tubulin serves as a loading control.

Source data are available online for this figure.

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Spliceosome components are required for the stable association of cohesin with chromatin in G2 cells

Following DNA replication, a subpopulation of cohesin complexes binds to chromatin with a mean residence time of around 6 h (Gerlich et al, 2006). This stably associated pool requires the protein sororin (Schmitz et al, 2007) and is thought to provide the
long-lived links mediating cohesion from S phase into mitosis. Our observation that the loss of splicing factors disrupts sister chromatid cohesion in G2 phase raises the possibility that the dynamics of cohesin association with chromatin are altered. To test this possibility, we used fluorescence recovery after photobleaching (FRAP) experiments. Cells stably expressing SMC1-EGFP were released from an S phase arrest for 5 h to allow entry into G2 phase. Following photobleaching of SMC1-EGFP in one half of the nucleus, the loss of the fluorescent signal in the unbleached nuclear area and the recovery of the signal in the bleached nuclear area were measured over 120 min (Fig 5). The redistribution kinetics of chromatin-bound cohesin were analysed by plotting the intensity difference between the two areas over time. Consistent with published observations (Gerlich et al, 2006; Schmitz et al, 2007), the SMC1-EGFP intensity difference dropped slowly and persisted until 120 min after photobleaching in control siRNA transfected cells (Fig 5). In contrast, in cells transfected with siRNA duplexes targeting the splicing proteins MFAP1, CDC5L and SART1, the intensity difference decreased rapidly and the fluorescence in the bleached and unbleached area equilibrated within about 45 min (Fig 5). Depletion of spliceosome components had a similar effect on cohesin dynamics as loss of sororin, a protein that antagonizes the activity of the cohesin release mechanism (Nishiyama et al, 2010) (Fig 5). Our results suggest that cohesin complexes dissociate from chromatin faster in cells lacking spliceosome components than in control cells. It is thus possible that splicing factors contribute to sister chromatid cohesion by

![Figure 5](image-url)

**Figure 5. Stable association of cohesin with chromatin requires splicing factors.**
Representative frames from fluorescence recovery after photobleaching (FRAP) experiments in SMC1-EGFP-expressing cells that were synchronized in G2 and that were previously transfected with control, MFAP1, SART1 and CDC5L siRNA duplexes (for 53 h) or with control and sororin siRNA (for 29 h) (upper panel). The graph shows the difference between SMC1-EGFP fluorescence intensity in the unbleached area and the intensity in the bleached area of the nucleus normalized to the first post-bleach frame and plotted over time (lower panel). Bars represent mean ± SEM, n = 6 cells for each condition from two independent experiments.
ensuring the stable association of cohesin with chromatin. Failure to do so may explain why cells that lack spliceosome subunits lose sister chromatid cohesion soon after DNA replication.

**Expression of RNase H1 does not restore cohesion in splicing factor-depleted cells**

Next, we investigated the molecular basis for how depletion of splicing factors perturbed the stable association of cohesin with chromatin and sister chromatid cohesion. Defective splicing of pre-mRNAs could abrogate sister chromatid cohesion through the formation of R loops, DNA:RNA hybrid structures that can threaten genome stability (Li & Manley, 2005; Aguilera & Garcia-Muse, 2012). To test this possibility, we generated a cell line overexpressing RNase H1, an enzyme that cleaves the RNA moiety of DNA:RNA hybrids (Cerritelli & Crouch, 2009; Skouri-Stathaki et al., 2011). Expression of nuclear-targeted RNase H1-EGFP failed to restore sister chromatid cohesion upon depletion of spliceosome components (Supplementary Fig S6) indicating that R loops are not responsible for the loss of cohesion.

**Loss of spliceosome components reduces the protein level and impairs the splicing of sororin but not of core cohesin subunits**

Alternatively, we hypothesized that compromised splicing and defective pre-mRNA processing could lead to the acute loss of one or multiple proteins that are essential for sister chromatid cohesion. Profiling of protein levels of cohesin subunits or regulators revealed that splicing factor depletion did not severely reduce the steady-state abundance of the cohesin subunits SMC1, SMC3, SCC1 and SA2 (Fig 6A). In contrast, depletion of all splicing factors tested (NHP2L1, MFA1, SART1 and CDC5L) resulted in a fourfold or higher drop in the protein levels of sororin, a protein required for maintenance of cohesion through S-G2 phase into mitosis and for the stable association of cohesin with chromatin (Fig 6A). The loss of spliceosome components could compromise the splicing of sororin pre-mRNA and, as a consequence, the abundance of mature sororin protein. Consistent with this hypothesis, analysis of the presence of exon-intron junctions in sororin RNA by real-time PCR indicated the increased retention of intron 1 and 2 in cells depleted of splicing factors (Fig 6B). In contrast, depletion of spliceosome subunits only had minor effects on the retention of introns in the RNAs encoding core cohesin subunits (Fig 6B). Retention of intron 1 in sororin’s mRNA would cause a translational frame shift after the coding exon 1 generating a stop codon in exon 3 or intron 2 if the latter is not removed. The resulting truncated protein would be non-functional.

Defects in pre-mRNA splicing could reduce protein levels acutely especially when coupled to a reduced half-life of the corresponding mRNA and/or protein. To assess the stability of sororin mRNA, we treated cells with actinomycin D to inhibit transcription. Sororin mRNA displayed reduced stability compared to the mRNAs encoding core cohesin subunits and to the mRNA encoding glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (Supplementary Fig S7A). In contrast to core cohesin subunits, sororin protein abundance dropped to about 25% within 2 h after releasing cells from mitosis into the G1 phase of the cell cycle (Fig 6C, left panel; for fluorescent quantification see Supplementary Fig S7B). The reduction in sororin protein level was further enhanced by treatment of cells with the protein synthesis inhibitor cycloheximide (Fig 6C, right panel; Supplementary Fig S7B). Thus, consistent with sororin being a substrate of the anaphase-promoting complex/cyclosome (APC/C) (Rankin et al., 2005), the protein is highly unstable in G1 cells.

Altogether, our observations raise the possibility that the loss of splicing machinery components impacts on sister chromatid cohesion by compromising the splicing of sororin pre-mRNA and, as a consequence, perturbing the cellular level of mature sororin protein. Consistent with this hypothesis, the loss of sororin and the depletion of splicing factors have similar phenotypic consequences: the loss of sister chromatid cohesion during interphase and the failure of cohesin to stably associate with chromatin.

**Expression of an intronless version of sororin restores sister chromatid cohesion in cells lacking splicing factors**

To test the hypothesis that failure of sororin pre-mRNA processing is causally linked to the loss of cohesion in cells depleted of spliceosome subunits, we generated a cell line stably expressing an intronless version of sororin (AcFL-Sororin-r) was also rendered resistant to sororin siRNA. Expression of AcFL-Sororin-r fully suppressed the loss of sister chromatid cohesion observed upon depletion of endogenous sororin suggesting that the transgene was functional (Fig 7A). Importantly, the intronless sororin transgene also potently restored sister chromatid cohesion in cells depleted of the splicing factors MFA1, NHP2L1, SART1 and CDC5L (Fig 7B and C). In contrast, expression of AcFL-Sororin-r only had a minor effect in cells depleted of the core cohesin subunit SCC1 or the centromeric protector protein SGOL1 indicating that the transgene cannot efficiently compensate for loss of core cohesin functions (Fig 7C). These results suggest that the defective synthesis of mature sororin protein makes a major contribution to the loss of sister chromatid cohesion in cells lacking spliceosome subunits (Fig 8B). Expression of the intronless sororin transgene did not, however, restore mitotic progression in cells transfected with siRNAs targeting spliceosome subunits indicating that the lack of as yet unidentified additional factors also contributes to the mitotic arrest in those cells (Supplementary Fig S8A).

**The loss of sister chromatid cohesion upon depletion of spliceosome subunits requires the anti-establishment factor WAPL**

Sororin ensures the stable association of cohesin with chromatin in interphase by antagonizing the activity of the cohesin release protein WAPL (Nishiyama et al., 2010). If defective splicing disrupts cohesin’s binding to chromatin and thereby sister chromatid cohesion through compromising sororin function, loss of WAPL should restore the links between sister chromatids in cells lacking spliceosome components. Indeed, we found that depletion of WAPL strongly suppressed the sister chromatid cohesion defect not only in cells depleted of sororin but also in cells transfected with siRNA duplexes targeting the splicing factors MFA1, NHP2L1, SART1 and CDC5L (Fig 8A; for protein depletion data see Supplementary Fig S8B). This observation supports the conclusion that spliceosome factors contribute to sister chromatid cohesion largely by maintaining sororin function, which counteracts the cohesin releasing activity of WAPL (Fig 8B).
Discussion

Cohesin and sister chromatid cohesion lie at the heart of chromosome biology. They are required for chromosome segregation, DNA repair, chromosome structure and gene regulation. Using a functional genomic approach, we have defined a requirement of pre-mRNA splicing factors for sister chromatid cohesion in human cells. Depletion of a multitude of splicing factors by RNAi abrogated the stable association of cohesin with chromatin after DNA replication and culminated in a dramatic loss of sister chromatid cohesion upon mitotic entry. The fact that we detected a severe loss of sister chromatid cohesion upon depletion of more than 26 spliceosome subunits suggests that defects in pre-mRNA splicing itself or in the processing of one or multiple transcripts are causally linked to the...
observed chromosomal phenotype. At this point, we have no
evidence that would support a direct involvement of spliceosome
components in cohesion.

Pre-mRNA splicing is an essential process for gene expression in
eukaryotes (Braunschweig et al., 2013). Accordingly, we found that
depletion of spliceosome subunits causes cell lethality as a terminal
phenotype (data not shown). However, our analysis suggests that
an enhanced turnover of cohesin on interphase chromatin followed
by a cell cycle arrest in mitosis with split sister chromatids is an early
cellular consequence of compromising splicing. This suggests that
sister chromatid cohesion is highly sensitive to pre-mRNA splicing
defects. Furthermore, it provides a potential explanation for the prom-
inent identification of spliceosome components as mitosis-defective
hits in reverse genetic screens (Kittler et al., 2007; Hofmann et al.,
2010; Neumann et al., 2010).

Our experiments indicate that compromised splicing leads to
defects in sister chromatid cohesion largely through the loss of sorro-
rin function, thus providing a molecular explanation for the essential

Figure 7. Expression of an intronless construct of sororin suppresses the loss of cohesion caused by depletion of splicing factors.

A Immunoblot analysis of sororin protein levels in whole-cell extracts prepared from cells stably expressing either the AcFL tag or an AcFL-tagged intronless and siRNA-
resistant transgene of sororin (AcFL-sororin-r). Cells were transfected with sororin (+) or control (−) siRNA duplexes 72 h prior to analysis (left). The endogenous
protein (filled arrowheads) and the transgenic AcFL-sororin-r counterpart (open arrowheads) are indicated. Chromosome spreads (right) from the indicated cell lines
were prepared 28 h after transfection with sororin siRNA (n > 100 cells, mean from 3 experiments). Scale bar, 10 μm.

B Representative images of chromosome spreads from the indicated cell lines transfected with SGOL1, NHP2L1 (34 h), control, MFAP1, SART1 (40 h) or CDC5L (56 h)
siRNA duplexes. Scale bar, 10 μm.

C Quantification of sister chromatid cohesion status corresponding to the experiment described in (B) (n > 100 cells, mean of 3 experiments). SCC1 siRNA was
transfected 56 h prior to analysis.

Source data are available online for this figure.
role of splicing factors in mitosis (Fig 8B). Several lines of experimental evidence support this conclusion. Loss of sororin and loss of splicing factors share a phenotypic signature (Schmitz et al., 2007): failure of cohesin to stably associate with chromatin and defects in sister chromatid cohesion in post-replicative interphase cells despite cohesin loading and SMC3 acetylation. Furthermore, we detected aberrant processing of sororin pre-mRNA and reduced sororin protein levels in cells depleted of splicing factors. Lastly, expression of an intronless version of sororin and depletion of the cohesin release protein WAPL, whose activity is antagonized by sororin, were both able to suppress the sister chromatid cohesion defect observed in cells lacking spliceosome components. These rescue experiments also indicate that the unrestrained WAPL-induced dissociation of cohesin from chromatin in interphase might be responsible for the loss of cohesion observed in mitotic cells upon depletion of splicing factors. While we describe 26 splicing factors whose depletion severely compromises sister chromatid cohesion, we analysed the effects of depletion of four proteins, MFAP1, NHP2L1, SART1 and CDC5L, in detail. Based on the phenotypic similarity caused by depletion of these 4 components, it is tempting to speculate that compromised sororin function may underlie the loss of sister chromatid cohesion elicited by depletion of most if not all splicing factors described here. This extrapolation is supported by the analysis and global RNA profiling of cells lacking the splicing factor SNW1 (Zhou...
et al., 2002; see van der Lelij et al., 2014), a protein also identified in our screen for spliceosome components that are required for cohesion. Based on our results, it is conceivable that compromised processing of sororin mRNA and the ensuing loss of cohesion might also contribute to the mitotic defects previously reported in human cells depleted of factors linked to RNA metabolism, such as the TREX helicase UAP56, the RNA-binding protein RBMX, the splicing protein SON and the Prp19 splicing complex (Huen et al., 2010; Yamazaki et al., 2010; Matsunaga et al., 2012; Hofmann et al., 2013).

Our results support the notion that sister chromatid cohesion in human cells is exquisitely sensitive to the dosage of sororin protein. This highlights the delicate balance that cells have to strike between the activity of the cohesin release factor WAPL and the sororin-dependent mechanism that counteracts WAPL and promotes the establishment of long-lived connections between sister chromatids. This balancing act is presumably necessitated by the important roles of cohesin dissociation from chromatin. WAPL-dependent cohesin release from mitotic chromosome arms drives the formation of the iconic X shape of chromosomes and facilitates the correct partitioning of sister genomes during anaphase (Kuang et al., 2006; Haarhuis et al., 2013; Tedeschi et al., 2013). It also spares a soluble pool of the cohesin complex from cleavage by separase at the metaphase-to-anaphase transition providing a ready supply of intact cohesin complexes that can be loaded onto chromatid during telophase/G1 to control chromatin structure and gene regulation (Wendt et al., 2008; Hadjur et al., 2009; Nativio et al., 2009; Sofueva et al., 2013; Tedeschi et al., 2013). Lastly, WAPL-mediated release of cohesin from DNA prevents chromatin condensation in interphase cells (Tedeschi et al., 2013).

Although we cannot rule out that defective processing of other transcripts also impairs cohesion in splicing factor-depleted cells, defective sororin function appears to be the major contributing factor. In contrast to other cohesin proteins, sororin is degraded at the end of the cell cycle through APC/C-mediated proteasomal degradation (Rankin et al., 2005) thereby necessitating renewed sororin translation from processed mRNA for every ensuing S phase. We hypothesize that this could render sister chromatid cohesion vulnerable to defects in pre-mRNA splicing. The reduced half-life of sororin mRNA and protein is likely to accelerate the conversion of splicing defects into a reduction of mature protein. Our data suggest that the removal of introns from sororin pre-mRNA may have a particularly strict requirement for intact spliceosomes. While this strict requirement may not be unique to sororin pre-mRNA on a transcriptome-wide basis (see van der Lelij et al., 2014), sororin appears to be the most sensitive pre-mRNA to depletion of spliceosome components amongst essential cohesin factors. This sensitivity may act synergistically with the high turnover of sororin gene products to cause a reduction of sororin protein level and a resulting defect in sister chromatid cohesion. The molecular basis for a potentially selective dependency of sororin pre-mRNA splicing (and that of some other pre-mRNAs) on intact spliceosomes remains to be determined. Its elucidation will be facilitated by the transcriptome-wide mapping of spliceosome assembly and branch site usage in conjunction with the global analysis of intron retention following loss of particular spliceosome components.

There is precedence for a specific mitotic defect caused by compromised splicing of a transcript in budding yeast. Removal of the intron from the gene encoding α-tubulin alleviated the mitotic arrest in cells that harboured spliceosome mutations (Burns et al., 2002). Expression of an intronless allele of sororin restored sister chromatid cohesion but did not alleviate the mitotic block in human cells depleted of splicing components. This suggests that the defective processing of additional transcripts may contribute to the spindle assembly checkpoint-mediated arrest that we observed in cells lacking splicing factors.

We noted that the efficiency of cohesion rescue by expression of an intronless sororin transgene was reduced if cells lacking spliceosome components were analysed at later time points. This suggests that cohesion fatigue, the gradual loss of cohesion in mitotically arrested cells (Daum et al., 2011; Stevens et al., 2011), contributes to the observed sister chromatid cohesion in cells lacking spliceosome components. However, our results indicate that a defect in sororin function rather than cohesion fatigue caused by a protracted mitotic arrest is the major cause for the disruption of cohesion in cells depleted of splicing factors. Firstly, splicing factors are required for the stable association of cohesin with chromatin during interphase. Second, depletion of splicing factors abrogates cohesion prior to mitotic entry. Lastly, expression of an intronless sororin transgene and the depletion of WAPL efficiently suppresses the loss of sister chromatid cohesion elicited by depletion of spliceosome subunits.

The phenotypic link between splicing and sister chromatid cohesion that we describe here may have biomedical implications. One of the splicing factors whose depletion by siRNA causes a dramatic loss of cohesion is SF3B1. SF3B1 has recently emerged as one of the most frequently mutated genes in patients with chronic lymphocytic leukaemia (CLL) (Rossi et al., 2011; Wang et al., 2011; Quesada et al., 2012). Somatic SF3B1 mutations were also detected at high frequency in myelodysplastic syndrome (MDS) patients (Papaemmanuil et al., 2011; Yoshida et al., 2011). The recurrent nature of SF3B1 mutations in MDS and CLL suggests that they act as key drivers in hematopoietic proliferative disorders. SF3B1 is an essential component of the spliceosomal U2 snRNP (Wahl et al., 2009; Folco et al., 2011). SF3B1 mutations could drive dysplasia and malignancy by altering the splicing pattern of oncogenes or tumour suppressors (Gentien et al., 2014). Our findings raise the possibility that heterozygous SF3B1 mutations detected in CLL and MDS cells could contribute to pathology also by altering the turnover of cohesin on chromatin, and hence affecting chromosome stability, DNA repair and gene regulation.

Materials and Methods

Cells and growth conditions

HeLa Kyoto, HEK 293FT and HCT116 cells used in this study were grown in Dulbecco’s modified Eagle medium, DMEM (Invitrogen) supplemented with 10% FCS (Sigma) and 1% PenStrep (Invitrogen). In order to establish stable cell lines, media supplemented with either 0.35 μg/ml puromycin (Sigma) or 500 μg/ml G418 (Invitrogen) were used. Cells were grown in an incubator that was maintained at 37°C with 5% CO₂.

siRNA screen

To compile a list of candidate genes, we selected 851 genes that were either identified as causing mitotic failure upon depletion according
to the genome-wide or targeted functional genomic screens (Mukherji et al., 2006; Kittler et al., 2007; Conery & Harlow, 2010; Neumann et al., 2010) or gene products that were identified as midbody-associated by proteomic analysis (Skop, 2004). Subsequently, we eliminated genes that were either well-characterized regulators of cell division, pseudogenes, ribosomal proteins or annotation errors according to NCBI databases. Based on this winnowing process, we compiled a final candidate list of 718 genes that were targeted in the primary screen. Dharmacon siGENOME smartpool siRNAs (0.1 nmol) were obtained for the 718 genes in 96-well plate format. Each well contained a pool of 4 siRNAs targeting a single gene. Positive controls (known mitotic regulators such as ECT2, MgcRacGAP, AURKB) and non-targeting negative controls (scrambled siRNA and RISC free siRNA) were included as well. Prior to cell seeding, siRNA pools (final concentration 37.5 nM) and the transfection reagent [Lipofectamine RNAiMAX (Invitrogen)] at 0.167 μl/well at a final siRNA to RNAiMAX ratio of 1:600 were deposited in the wells [clear-bottomed 96-well plates (Falcon, Beckton Dickinson)] and mixed. Subsequently, 2,500 HeLa Kyoto cells were added to each well. Cells were grown for 52 h at 37 °C and then processed for fixation with ethanol at 0% overnight. Following the fixation, the cells were stained with Cellomics Whole Cell Stain (1:75 dilution, Thermo Scientific), and the DNA was visualized by DAPI (4′,6-diamidino-2-phenylindole dihydrochloride, Invitrogen) at a final concentration of 1 μg/ml. The screen was performed in triplicate. Automated image acquisition was performed using an ArrayScan ATi microscope (Thermo Scientific) equipped with a 10× objective. Twenty images per well were captured and subsequently processed using ImageJ software (http://rsweb.nih.gov/ij/). The processed images were scored for the percentage of cells containing abnormally shaped nuclei (fragmented, multinucleated nuclei) or multiple nuclei. The three biological plates were scored independently, and the median value (the percentage of cells with abnormal nuclei) was calculated and plotted.

siRNA sequences and transfection protocol

The following siRNA duplexes were used at a final concentration of 37.5 nM: control siRNA (Thermo Scientific siGENOME Non-Targeting siRNA #1 D-001210-01 and #4 D-001210-04, RISC free siRNA D-001210-01) and Thermo Scientific siGENOME smartpool siRNAs targeting Mad2 (M-003271-01), ESCO1 (M-023413-01), ESCO2 (M-025788-01), MFAP1 (M-020071-01), SART1 (M-017283-01), NHP2L1 (M-019900-01), CDC5L (M-011237-00), SGOL1 (M-015475-01) and SCC1 (M-006832-01). siGENOME individual siRNAs were used against MFAP1 (D-020071-03) (Figs 1B–E, 2, 4–8, and Supplementary Figs S4, S6 and S8), SART1 (D-017283-01) (Figs 4–8 and Supplementary Figs S4, S6 and S8), NHP2L1 (D-019900-01) (Figs 4–8 and Supplementary Figs S4, S6 and S8), CDC5L (D-011237-03), SGOL1 (M-015475-01) and SCC1 (M-006832-01). siGENOME individual siRNAs were used against MFAP1 (D-020071-03) (Figs 1B–E, 2, 4–8, and Supplementary Figs S4, S6 and S8), SART1 (D-017283-01) (Figs 4–8 and Supplementary Figs S4, S6 and S8), NHP2L1 (D-019900-01) (Figs 4–8 and Supplementary Figs S4, S6 and S8), CDC5L (D-011237-03), SGOL1 (M-015475-01) and SCC1 (M-006832-01). siGENOME individual siRNAs were used against MFAP1 (D-020071-03) (Figs 1B–E, 2, 4–8, and Supplementary Figs S4, S6 and S8), SART1 (D-017283-01) (Figs 4–8 and Supplementary Figs S4, S6 and S8), NHP2L1 (D-019900-01) (Figs 4–8 and Supplementary Figs S4, S6 and S8), CDC5L (D-011237-03), SGOL1 (M-015475-01) and SCC1 (M-006832-01). siRNA sequences and transfection protocol

To create tagged versions of the genes used in the study (MFAP1, NHP2L1, SART1) for expression in human cells, AcGFP (Aequorea coeruleus G F P) was amplified from pAcGFP-N1 (Clontech) and inserted into pIRESpuro3 (Clontech). During amplification, a Kozak sequence (CGCCACC) and a FLAG epitope (DYKDDDDK) were added to AcGFP before the start codon and after the last amino acid, respectively, creating pIRESpuro3-AcFL (Su et al., 2011; Lekomtsev et al., 2012). The coding sequences of MFAP1, NHP2L1 and SART1 were amplified from cDNA (Source Biosciences) and introduced into pIRESpuro3-AcFL using the restriction enzymes Agel and EcoRI. To create siRNA-resistant variant of the genes, the following sequences were mutated using the Quikchange II site directed mutagenesis kit (Stratagene) (underlined are the silent nucleotide substitutions). For MFAP1, the sequence aagtgaaggtaaagcgtta was mutated to agtggtggagctgctg; for NHP2L1, the sequence tggagctgctg was mutated to tggagctgctg; and for SART1, the sequence ggacagctgctg was mutated to gcaagactgctg. The plasmids were transfected into HeLa Kyoto cells using FuGENE 6 transfection reagent (Roche). Clonal cell lines were isolated after 2 weeks of antibiotic selection and characterized by immunofluorescence microscopy and immunoblotting. HeLa Kyoto cells were transfected with a plasmid encoding nuclear-targeted RNase H1-EGFP [a generous gift from Robert Crouch (Cerritelli & Crouch, 2009)] following which GFP-positive cells were enriched by using fluorescence-activated cell sorting (FACS). Cells expressing H2B-mCherry were kindly provided by Su et al. (2011) and were grown in DMEM media supplemented with 500 μg/ml G418 (Invitrogen). A cell line stably expressing H2B-mCherry and α-tubulin-GFP was kindly provided by Laurent Sansregret. Cells stably expressing SMC1-EGFP were used for photo-bleaching experiments and were a kind gift from Schmitz et al. (2007).

Preparation of stable cells lines using lentiviral infection

To create tagged alleles of sororin for expression in human cells, the coding sequence of sororin was introduced into pIRESpuro3-AcFL using the restriction enzymes Agel and EcoRI. To create an siRNA-resistant variant of sororin, the sequence cgaggggaggaggaggaggaggaggaggaggggaggaggaggagggggaggagggaggaggggaggaggaggaggaggaggaggaggaggaggggaggaggaggagggould be excised from pIRESpuro3 using the restriction enzymes Clal and EcoRI and cloned into pLVX-puro (Clontech). HEK 293FT cells were pre-seeded and grown to about 50% confluency before being transfected with pLVX-puro-Sororin (WT or siRNA resistant) and a second-generation packaging system [pAX2 and pMD2.G (Addgene)] using Lipofectamine 2000 (Invitrogen). Viral particles were collected 48 h after transfection by harvesting the supernatant and filtering it through a 0.45-μm PVDF filter unit (Milllex HV). HeLa Kyoto cells were infected with different titres of viral particles in presence of 8 μg/ml polybrene (Sigma). The cells were grown for 48–60 h in the presence of viral particles before the medium was supplemented with 0.35–0.4 μg/ml puromycin (Sigma) for transgene selection. After passing the cells for 3 generations, clonal cell lines were generated after 2 weeks of antibiotic titrations of viral particles in presence of 8 μg/ml polybrene (Sigma). The cells were then incubated for the indicated durations and processed for experiments.

Generation of stable cell lines using plasmid transfection

To create tagged versions of the genes used in the study (MFAP1, NHP2L1, SART1) for expression in human cells, AcGFP (Aequorea coeruleus GFP) was amplified from pAcGFP-N1 (Clontech) and inserted into pIRESpuro3 (Clontech). During amplification, a Kozak sequence (CGCCACC) and a FLAG epitope (DYKDDDDK) were added to AcGFP before the start codon and after the last amino acid, respectively, creating pIRESpuro3-AcFL (Su et al., 2011; Lekomtsev et al., 2012). The coding sequences of MFAP1, NHP2L1 and SART1 were amplified from cDNA (Source Biosciences) and introduced into pIRESpuro3-AcFL using the restriction enzymes Agel and EcoRI. To create siRNA-resistant variant of the genes, the following sequences were mutated using the Quikchange II site directed mutagenesis kit (Stratagene) (underlined are the silent nucleotide substitutions). For MFAP1, the sequence aagtgaaggtaaagcgtta was mutated to agtggtggagctgctg; for NHP2L1, the sequence tggagctgctg was mutated to tggagctgctg; and for SART1, the sequence ggacagctgctg was mutated to gcaagactgctg. The plasmids were transfected into HeLa Kyoto cells using FuGENE 6 transfection reagent (Roche). Clonal cell lines were isolated after 2 weeks of antibiotic selection and characterized by immunofluorescence microscopy and immunoblotting. HeLa Kyoto cells were transfected with a plasmid encoding nuclear-targeted RNase H1-EGFP [a generous gift from Robert Crouch (Cerritelli & Crouch, 2009)] following which GFP-positive cells were enriched by using fluorescence-activated cell sorting (FACS). Cells expressing H2B-mCherry were kindly provided by Su et al. (2011) and were grown in DMEM media supplemented with 500 μg/ml G418 (Invitrogen). A cell line stably expressing H2B-mCherry and α-tubulin-GFP was kindly provided by Laurent Sansregret. Cells stably expressing SMC1-EGFP were used for photo-bleaching experiments and were a kind gift from Schmitz et al. (2007).
selection. The cell lines were characterized by immunofluorescence microscopy and immunoblotting.

Other cell lines

Cells expressing H2B-mCherry were kindly provided by Su et al (2011) and were grown in DMEM media supplemented with 500 μg/ml G418 (Invitrogen). A cell line stably expressing H2B-mCherry and α-tubulin-GFP was kindly provided by Laurent Sansregret. Cells stably expressing SMC1-EGFP were used for photobleaching experiments and were a kind gift from Schmitz et al (2007).

Cell synchronization and drug treatments

To synchronize cells, 2.5 mM thymidine (Sigma) was added to cells either at the time of transfection or 24 h later. At the time of release, cells were rinsed twice with DMEM and fresh medium was added. The cells were allowed to proceed through the cell cycle either for 3 h to enrich for cells in S phase (Fig 4C) or for 5 h to enrich cells in G2 phase (Figs 4A, 5, 6A–C, and 7A left panel). To synchronize cells for performing chromosome spreads, asynchronous HeLa Kyoto cells that were grown for 24, 30, 36 or 52 h after siRNA transfection were treated with 330 nM nocodazole (Sigma) for 4 h. To study mRNA half-life (Supplementary Fig S7A), asynchronous HeLa Kyoto cells were treated with 5 μg/ml actinomycin D (Sigma) and cells were harvested at the indicated time points after drug treatment for RNA analysis. To study protein half-life (Fig 6C and Supplementary Fig S7B), HeLa Kyoto cells were synchronized with 2.5 mM thymidine for 24 h following which they were released for 8 h and then treated with 30 ng/ml nocodazole for 5 h after which the cells were harvested by mitotic shake-off and allowed to re-enter the cell cycle. Two hours after release from nocodazole, cells were treated with either DMSO or 100 μg/ml cycloheximide (Sigma) and harvested at the indicated time points for protein analysis. To score for the ability of the cells to form a metaphase plate (Fig 1E), HeLa Kyoto cells were synchronized in metaphase by treatment for 3 h with the proteasome inhibitor MG132 at a final concentration of 10 μM.

Real-time PCR for measurement of RNA levels

The following primers were used for the real-time quantitative PCR (RT-qPCR) experiments (Fig 6B and Supplementary Fig S7A); sororin Exon1-Intron1 (Fwd: ACAGGTTCATAGAGAAGGCGAG, Rev: AGTTATGTCTGGGAGGGCAA), sororin Exon2-Intron2 (Fwd: AGAGGGATGAACTGAGGCTC, Rev: TGGGGCGTCTGCTCTTAA), sororin Exon5 (Fwd: AAGTTGAGGCTTCTACAGC, Rev: TCGAAGCCAAGGCAAG), SMC3 Exon1-Intron1 (Fwd: AGCAATCTGGTCATTGTTTGTCTTTTTTAT, Rev: TGGCAGTTTTTTTTCTTATGCTTT), SMC3 Intron1 (Fwd: AAAGTGTGATGAAAGAAATTGGCGAAC, Rev: GTTT GAGGGGCAGGATGTTG), SMC1A Exon9-Intron9 (Fwd: AGATCATCAGCGCCCTTTACC, Rev: GGATTGCAAACCTTGGCTCTTAA), SMC1A Exon9 (Fwd: GACAGAGAGGAGGGATG, Rev: CAGGTTAAGGCGTGTAGT), SCCI 1 Intron1-Exon2 (Fwd: AAGAQAGACTATGAAAGCGCCGAGGA, Rev: AGGCTTGCAAGAGATTGCG), SCCI 1 Intron1 (Fwd: CCTAATCAGATTTGTTGTGGTCC, Rev: TACACTC GAACCATGGGCC), SA2 Exon4-Intron4 (Fwd: AAGCATGAGGAGGATATAGACT, Rev: TGTGACTATTTAGATTGAGC), SCA2 Exon5 (Fwd: AGATTATCACTTACCATGCTG, Rev: CAGGGTGCTTGTA TGTCGA) and GAPDH (Fwd: CCTCCCGCTTCGCTCTCT, Rev: CTGGCGAGGCAAAAAGAAGA). G2 synchronized HeLa Kyoto cells transfected with the indicated siRNAs (Fig 6B) or asynchronous HeLa Kyoto cells treated with DMSO or 5 μg/ml actinomycin D (Supplementary Fig S7A) were harvested and lysed in RLT plus buffer (Qiagen RNeasy Plus kit) + β-mercaptoethanol (Sigma). RNA was extracted from the cells using the RNeasy plus MiniKit (Qiagen) according to the manufacturer’s instructions. After measuring the RNA concentration in a NanoDrop spectrophotometer, 1.5 μg of total RNA was reverse transcribed with random hexamer primers using the TaqMan reverse transcription kit (Applied Biosystems). Pre-mRNA and mature mRNA levels were assessed by real-time quantitative PCR (RT-qPCR) performed using the indicated primer pairs using IQ-SYBR Green Supermix and CFX96 Real-Time System (Bio-Rad) using the manufacturer’s instructions. The relative amount of RNA was calculated using the ΔΔCt method as described earlier (Winer et al, 1999). GAPDH mRNA was chosen for normalization of the mRNA measurements (Fig 6B and Supplementary Fig S7A).

Immunofluorescence microscopy (IF)

Cells grown on coverslips with a diameter of 18 mm and thickness 1 (Assistent) were fixed overnight with methanol at −20°C or for 10 min at 37°C in 4% formaldehyde in PBS (Thermo Scientific, 16% PFA stock diluted with PBS). After fixation, samples were washed three times in 0.01% Triton X-100 in PBS for 5 min each and then permeabilized with 0.2% Triton X-100 in PBS for 10 min. Following three washes in 0.01% Triton X-100 in PBS, the coverslips were incubated with blocking solution (3% BSA in PBS containing 0.01% Triton X-100) for 1 h. Samples were incubated with the primary antibody in the blocking solution at 4°C overnight. Samples were then washed with three times with 0.01% Triton X-100 in PBS being incubated with secondary antibody diluted in blocking solution for 1 h along with 1 μg/ml DAPI. Following three more washes with 0.01% Triton X-100 in PBS, the coverslips were mounted onto microscopic slides with ProLong Gold (Molecular Probes) and left to dry overnight. Images were acquired on Zeiss Axio Imager M1 or M2 microscopes using a Plan Neofluor 40×/1.3 oil objective lens or 63×/1.4 Apochromat oil objective lens (Zeiss) equipped with an ORCA-ER camera (Hamamatsu) and controlled by Velocity 6.1, software (Improvision). Images were deconvolved using Velocity’s iterative restoration function. The images were processed using Imagej and Adobe Photoshop software.

Antibodies and dyes

The following primary antibodies were used in this study for immunoblotting (IB) and immunofluorescence (IF) experiments: mouse monoclonal anti-AcGFP (JL8, Clontech, IF & IB 1:2,000, rabbit anti-AcGFP (Clontech, IF & IB 1:2,000), mouse monoclonal anti-α-tubulin (B512, Sigma, IF & IB 1:10,000), rabbit monoclonal anti-β-tubulin HRP Conjugate (9F3, Cell Signalling, IF 1:2,000), mouse monoclonal anti-SMC1 (6892, Cell Signalling, IB 1:1,000), rabbit polyclonal anti-SMC3 (ab9263, Abcam, IB 1:2,000), rabbit monoclonal anti-SA2 (5882, Cell Signalling, IB 1:1,000), rabbit polyclonal anti-MFAP1 (SAB2104903, Sigma, IB 1:2,000), rabbit polyclonal anti-NHP2L1 (95958, Abcam IB 1:5,000) and rabbit polyclonal anti-SART1 (SC376460, Santa Cruz, IB 1:1,000). Jan-Michael Peters (IMP,
Splicing factors are required for cohesion  
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Vienna) kindly provided the following antibodies: rabbit polyclonal anti-SGOL1 (g975, IB 1:500), rabbit polyclonal anti-sororin (g953, IB 1:2,000), rabbit polyclonal anti-SCC4 (g974, IB 1:1,000) and rabbit polyclonal anti-SCC1 (g890, IB 1:1,000). Katsuhiko Shirahige (University of Tokyo) kindly provided a monoclonal mouse antibody recognizing SMC3 acetylated at Lys105 and Lys106 (IB 1:1,000) (Nishiyama et al, 2010). Secondary antibodies conjugated to either Alexa Fluor 488 or 594 (Molecular Probes) at a final dilution of 1:1,000 were used for IF detection. DNA was stained with 1 mg/ml DAPI (Molecular Probes). HRP-conjugated secondary antibodies at a concentration of 1:5,000 were used to detect proteins on immunoblots using chemiluminescence (GE Healthcare). For fluorescent detection of Western blotting, anti-rabbit IgG conjugate (DyLight R 800) (Cell Signalling) at a final concentration of 1:1,000 were used for detection with Odyssey Imaging System (LI-COR Biosciences).

Time-lapse microscopy

To quantify mitotic duration and cell division status (Fig 1B and C), cells were seeded in 12-well plates (Nunc). An hour prior to imaging, cells were transfected with indicated siRNAs in 24-well ImageLock plates (Essen BioScience). 12 h after seeding, the plate was imaged using an Incucyte live cell imaging system (Essen BioScience). Images were processed using ImageJ software and analysed using the mitotic duration plugin. For high resolution imaging (Fig 1D), HeLa Kyoto cells stably expressing H2B-mCherry and x-tubulin-GFP were seeded in Labtek chambers (Nunc, Thermo Scientific) and imaged at intervals of 5 min using a Zeiss Axio Observer Z1 microscope controlled by SimplePCI software (Hamamatsu) equipped with an Orca 03G01 camera (Hamamatsu) and a Plan-Apochromat 10|/0.45 objective.

To quantify mitotic duration for Supplementary Fig 5A, cells were reverse transfected with the indicated siRNAs in 24-well ImageLock plates (Essen BioScience). 12 h after seeding, the plate was imaged using an Incucyte live cell imaging system (Essen BioScience). Images were processed using ImageJ software and analysed using the mitotic duration plugin. For high resolution imaging (Fig 1D), HeLa Kyoto cells stably expressing H2B-mCherry and x-tubulin-GFP were seeded in Labtek chambers (Nunc, Thermo Scientific) and imaged at intervals of 5 min using a Zeiss Axio Observer Z1 microscope controlled by SimplePCI software (Hamamatsu) with a 40|/1.3 DIC H oil objective.

Chromosome spreads

Chromosome spreads of HeLa Kyoto and HCT116 cells were performed as follows. Cells transfected with the indicated siRNAs at a final concentration of 37.5 nM were grown at 37°C for 24, 30, 36 or 52 h as indicated. To enrich for mitotic cells, the medium was supplemented with 330 nM nocodazole for 4 h. Cells were harvested by mitotic shake-off and centrifugation. Subsequently, cells were incubated in a hypotonic solution (DMEM: filtered deionized water at 1:2) for 6 min at room temperature (RT). Cells were subsequently fixed with freshly made Carnoy’s buffer (1:3 Glacial acetic acid: methanol) for 15 min at RT and pelleted. This fixation step was repeated three times. The suspension of cells in Carnoy’s buffer was dropped onto a clean slide from a distance of 2 feet and left to dry overnight at RT. The slides were washed in PBS solution containing 1 mg/ml DAPI and mounted using Prolong Gold mounting solution. Chromosome spreads from individual cells were classified and scored with regards to the status of sister chromatid cohesion based on the indicated morphological criteria (normal X-shaped/parallel/split) and represented in graphs. Within a single karyotype, the majority of chromatids and chromosomes typically conformed to one of the categories used and the spread was therefore classified accordingly. For sister chromatid cohesion percentage data associated with micrographs, the categories of parallel and fully split chromatids were combined.

Fluorescence in situ hybridization (FISH) in mitotic cells

To analyse the status of sister chromatid cohesion in intact mitotic cells, we performed FISH experiments (Fig 2B). Cells were reverse transfected with indicated siRNAs onto 18-mm coverslips and grown for 48 h. Subsequently, the cells on the coverslips were fixed with Carnoy’s buffer for 15 min at RT. FISH probes for the centromeres of chromosomes 6 (LPE 06G) and chromosome 8 (LPE 08R) (Cytocell aquarius) were used to probe the status of sister chromatid cohesion. The FISH probes were diluted in hybridization buffer (Cytocell aquarius) 1:10 and subsequently added to the fixed coverslips. Denaturation was performed for 3 min at 75°C, and the slide was left to hybridize overnight at 37°C in a humidified and lightproof chamber. Following this, the coverslip was washed in 0.25x SSC for 3 min at 73°C and subsequently in 2x SSC with 0.05% Tween 20 for 30 s at RT. The coverslips were then incubated in PBS containing 1 mg/ml DAPI to counterstain DNA. Finally, the coverslips were mounted using Prolong Gold mounting solution. Images were then acquired on a Zeiss Axio Imager M2 microscope using a Plan Neofluor 40|/1.3 oil objective lens equipped with an ORCA-ER camera (Hamamatsu) and controlled by Volocity 6.1 software (Improvision). Distance measurements were performed in Volocity by locating the centres of each of the pair of centromids manually and measuring the distance along a straight line between the two centroids. If the distance between two closest signals for a chromosome was larger than 2 µm, they were considered to be split.

FISH in interphase cells

Interphase FISH experiments (Fig 4A) were performed as previously described (Schmitz et al, 2007). Briefly, cells were transfected with the indicated siRNA duplexes on coverslips with a diameter of 18 mm and thickness 1 (Assistent). 2.5 mM thymidine was added either at the time of transfection or 24 h later. Following 24 h of thymidine arrest, cells were released from the thymidine arrest by rinsing twice in DMEM and then allowed to proceed through the cell cycle. 5 h after release, they were fixed with Carnoy’s buffer (15 min at RT) and left to dry overnight. BAC clone RP11-113F1 corresponding to the human tff1 locus was kindly provided by Erwan Watrin (University of Rennes). The BAC was used to generate FISH probes labelled with Primeti II Random primer labelling kit (Stratagene) and Cy3-dCTP (Amersham Biosciences) as described earlier (Schmitz et al, 2007). Just before use, the probes were diluted in hybridization buffer (Cytocell Aquarius) 1:10. The probe was added to the fixed coverslips. The coverslip, along with the probes were denatured for 3 min at 75°C. Subsequently, they were incubated in a humidified chamber at 37°C overnight. After the incubation, the coverslips were washed briefly in 0.4x SSC made from a stock of 20x SSC (3M NaCl + 300 mM Na2C6H5O7) at 72°C for 3 min and then in 2x SSC with 0.05% Tween 20 at RT for 30 s.
Following this, DNA was counterstained with 1 μg/ml DAPI and the coverslips were mounted with ProLong Gold (Molecular Probes). Images were acquired on a Zeiss Axio Imager M1 microscope using a Plan Neofluor 40×/1.3 oil objective lens equipped with an ORCA-ER camera (Hamamatsu) and controlled by Volocity 6.1. software (Improvision). Images were deconvolved using Volocity’s iterative restoration function. Distance measurements were performed in Volocity by locating the centres of each of the pair of centroids manually and measuring the distance along a straight line between the two centroids. In control and SGOL1 depleted cells, since the distances were often difficult to resolve because of the proximity of the dots. As a consequence, only cells where at least 2 out of the 3 centroid pairs were resolvable were considered for the analysis. In sororin, MFAP1, NHP2L1 and SART1 depleted cells, the vast majority of paired signals were clearly resolved and could be measured.

Photobleaching experiments

Photobleaching experiments (Fig 5) were performed as described previously (Gerlich et al., 2006; Schmitz et al., 2007). Cells stably expressing SMC1-EGFP, obtained from Schmitz et al. (2007), were reverse transfected with the indicated siRNA duplexes at a final concentration of 37.5 nM in Labtek 2 chambered borosilicate coverglass chambers (Nunc, Thermo Scientific). 2.5 mM thymidine was added either at the time of seeding or 24 h post-seeding. Cells were released from thymidine arrest after 24 h (by rinsing twice in DMEM) and allowed to proceed through the cell cycle for 5 h. An hour before imaging, the medium was changed to CO2-independent medium without phenol red (Invitrogen) supplemented with 10% FCS, 0.2 mM L-glutamine, PenStrep and 1 mM Na-pyruvate (all Invitrogen). 1 μg/ml cycloheximide (Sigma) was added to avoid new synthesis of SMC1-EGFP. The imaging and bleaching was performed at 37°C using a Olympus FX1000D (InvertedMicroscopeX81) laser confocal scanning microscope equipped with a PlanApoN ×60/1.40 NA Oil Sc objective lens and controlled by FV10-ASW software (Olympus). One half of the nuclear region was bleached leaving the other half intact. Repeated bleaching was performed every 10 s for five iterations in order to remove the soluble pool of SMC1-EGFP. This resulted in a reduction in the fluorescence intensity of the unbleached nuclear area as well. The first post-bleach frame used for the downstream analysis was acquired 2 min after photobleaching to allow for complete equilibration of bleached soluble SMC1-EGFP across the nucleus. ImageJ software was used for the intensity measurements. EGFP intensities were measured by drawing a rectangular area of the same dimensions in the bleached and unbleached area followed by subtraction of the mean background signal outside of the cell. The difference between the background corrected mean fluorescence intensity of the unbleached and bleached area normalized to the difference in the first post-bleach frame was calculated and plotted over time for the different siRNA treatments.

Supplementary information for this article is available online: http://emboj.embopress.org

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Author contributions

MP and SS designed the study. SS performed and analysed most of the experiments with contributions from MDVN and SL. The siRNA screen was carried out in collaboration with MH. MP and SS wrote the manuscript.

Conflict of interest

The authors declare that they have no conflict of interest.

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