A genetic interaction map centered on cohesin reveals auxiliary factors in sister chromatid cohesion

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Abstract

Eukaryotic chromosomes are replicated in interphase and the two newly duplicated sister chromatids are held together by the cohesin complex and several cohesin auxiliary factors. Sister chromatid cohesion is essential for accurate chromosome segregation during mitosis, yet has also been implicated in other processes, including DNA damage repair, transcription and DNA replication. To assess how cohesin and associated factors functionally interconnect and coordinate with other cellular processes, we systematically mapped genetic interactions of 17 cohesin genes centered on quantitative growth measurements of >52,000 gene pairs in budding yeast. Integration of synthetic genetic interactions unveiled a cohesin functional map that constitutes 373 genetic interactions, revealing novel functional connections with post-replication repair, microtubule organization and protein folding. Accordingly, we show that the microtubule-associated protein Irc15 and the prefoldin complex members Gim3, Gim4 and Yke2 are new factors involved in sister chromatid cohesion. Our genetic interaction map thus provides a unique resource for further identification and functional interrogation of cohesin proteins. Since mutations in cohesin proteins have been associated with cohesinopathies and cancer, it may also identify cohesin interactions relevant in disease etiology.
Introduction

Sister chromatid cohesion ensures close proximity of the two sister chromatids from the time of replication until their separation to opposite spindle poles during mitosis. Sister chromatid cohesion is mediated in all eukaryotic cells by a multiprotein complex called cohesin (Michaelis et al., 1997). In budding yeast, Smc1, Smc3, Scc1 and Scc3 make up the core of the cohesin complex, which is loaded onto chromatin during G1-phase. It forms a ring-like structure that encircles sister chromatids generated during DNA replication in S-phase in a manner dependent on Smc3 acetylation by Eco1. Subsequently the cohesive status is sustained throughout G2- and M-phase by several maintenance factors, including Rad61, Pds5 and Sgo1. Several accessory proteins have also been implicated in promoting sister chromatid cohesion, including Elg1, Ctf18, the alternative RFC complexes, the replisome component Ctf4, the Chl1 helicase-like protein, the chromatin remodeler Chd1 and the S-phase checkpoint proteins Mrcl and Tof1 (Petronczki et al., 2004, Parnas et al., 2009, Hanna et al., 2001, Skibbens, 2004, Xu et al., 2004, Boginya et al., 2019). Finally, sister chromatid cohesion is dissolved at the metaphase to anaphase transition by proteolytic activity of Esp1 towards Scc1 (Uhlmann et al., 1999, Cohen-Fix et al., 1996, Xiong and Gerton, 2010).

Besides ensuring proper chromosome segregation, cohesin has been shown to impact the repair of DNA double-strand breaks (DSBs) (Unal et al., 2004, Unal et al., 2007, Strom et al., 2004, Heidinger-Pauli et al., 2009, Gelot et al., 2016, Wu et al., 2012, Kong et al., 2014), gene expression (Gullerova and Proudfoot, 2008, Dorsett, 2011, Lengronne et al., 2004) and nuclear organization (Harris et al., 2014, Yamin et al., 2020). In addition, several developmental disorders have been causally linked to germline mutations in cohesin genes and are collectively referred to as cohesinopathies. These include Cornelia de Lange Syndrome (Deardorff et al., 2012, Liu and Baynam, 2010), Roberts Syndrome (Vega et al., 2005) and Warsaw Breakage Syndrome (van der Lelij et al., 2010). Somatic mutations in cohesin genes, on the other hand, have been found with high frequency in various types of cancer (Thol et al., 2014, Bailey et al., 2014, Repo et al., 2016, Deb et al., 2014), underscoring the importance of cohesin genes in the development of pathogenesis. However, despite the important role that cohesin genes play in various cellular processes, including those relevant to disease manifestation, our understanding of
how the cohesin complex functionally interconnects with these processes is still rather limited.

Genetic interaction screens have highlighted the connectivity between genes and their corresponding pathways, thus providing insight into the biological role(s) of individual genes (Mani et al., 2008). In yeast, such screens have led to the identification of new genes that contribute to efficient sister chromatid cohesion (Mayer et al., 2004, Chen et al., 2012) and provided valuable insight into the connectivity between cohesin genes and genes involved in DNA repair and DNA replication (McLellan et al., 2012, Warren et al., 2004). However, these studies were focused on a rather limited number of cohesin genes. Here, we examined genetic interactions between 17 different cohesin genes and more than 1400 genes involved in various biological processes in a quantitative manner. The resulting genetic interaction map describes novel connections for cohesin genes in various cellular processes, including post-replication repair, microtubule organization and protein folding, and reveals that the microtubule-associated protein Irc15 and prefoldin complex members Gim3, Gim4 and Yke2 are novel regulators of sister chromatid cohesion. Thus, we provide a unique and powerful resource for the identification and functional interrogation of cohesin proteins.

Results

Mapping genetic interactions of cohesin

To gain more insight into the relationship between sister chromatid cohesion and other cellular processes, a comprehensive genetic interaction map centered on cohesin was generated. To this end, query strains carrying gene deletion or temperature-sensitive alleles of 17 different cohesin genes and 18 DNA damage response (DDR) genes (Table S1) were crossed by using the synthetic genetic array (SGA) methodology (Tong and Boone, 2006) against a panel of 1494 array strains (Table S2) carrying gene deletion or Decreased Abundance of mRNA Perturbation (DAmP) alleles of genes that represent various biological processes (Fig. 1A). We previously used the 18 DDR mutants to map interactions of the DDR network and included these in the current study to warrant quality control and quality assurance (Guenole et al., 2013, Srivas et al., 2013). Genetic interactions were scored by quantifying colony sizes of the double mutants, which were normalized and
statistically analyzed to provide each mutant with a quantitative S-score (Fig. 1A). S-scores ≤ -2.5 represent negative or synthetic sick/lethal interactions, whereas S-scores ≥ 2 represent positive or alleviating/repressive interactions (Costanzo et al., 2019, St Onge et al., 2007, Hartman et al., 2001). In total, the profile map contains S-scores for 52,290 gene pairs (Fig. 1A and Table S3). Several routine quality control metrics were employed to ensure a high-quality map (Fig. S1). We observed a correlation of at least 50% between the genetic interactions identified in our screen and previously published genetic interaction maps (Fig. S1A-B) (Guenole et al., 2013; Collins et al., 2010, Costanzo et al., 2010). In addition, genetic interactions with the highest S-scores showed a high enrichment of interactions present in the Biogrid database (Fig. S1C).

Our genetic interaction map revealed in total 678 interactions, including 55 positive and 632 negative interactions (Fig. 1B). Validation of ~70 interactions resulted in an overall false discovery rate (FDR) of 31% (Fig. S1D-G). In particular, we identified 348 negative and 25 positive interactions for the cohesin-related genes along with 342 negative and 33 positive interactions for the DDR genes (Fig. 1B). As expected, interactions found in the cohesin-associated group were highly enriched for GO terms “sister chromatid cohesion” and “chromosome segregation”, whereas interactions for the DDR-associated genes were enriched for DNA repair-related GO terms (Fig. 1C, Tables S4, S5 and S6). In conclusion, a high-quality genetic interaction map centered on cohesin was generated, providing a useful resource to mine for crosstalk between sister chromatid cohesion and other cellular processes.

Cohesin genes interconnect with genes involved in various biological processes
To better understand the complexity of the interplay between sister chromatid cohesion and other biological processes, we generated a genetic interaction network comprising interactions with S-scores ≤ -2.5 and ≥ 2 for the cohesin-related query genes (Fig. 2). This interaction network may be relevant for other species as the vast majority of genes are orthologous to both fission yeast and human genes (Table S7). As expected, we observed a strong relationship between sister chromatid cohesion factors and genes involved in cell cycle control (e.g. SIC1, CTF19, BUB1, BUB3), as well as in DNA replication (e.g. RTT101, MMS22, POL2), which is in agreement with the required coordination of these three processes to guarantee faithful chromosome
duplication and segregation (Lengronne and Schwob, 2002, Fernius and Marston, 2009, Alexandru et al., 1999, Zhang et al., 2017, Edwards et al., 2003). Our network also revealed several known interactions between cohesin factors, mainly the non-essential cohesin accessory factors such as ELG1, TOF1 and RMI1, and genes involved in DSB repair (e.g. RAD51, RAD52, SRS2) (Ben-Aroya et al., 2003, Chang et al., 2005, Kanellis et al., 2003). Moreover, several interactions between cohesin factors and chromatin remodeling or histone-modifying complexes, such as ASF1, IES1, HTZ1, SWR1, HDA1 and HST3, strengthen the link between sister chromatin cohesion and chromatin architecture (Huang et al., 2004, Huang and Laurent, 2004, Munoz et al., 2019, Sharma et al., 2013, Thaminy et al., 2007). Finally, we found a strong interplay between both essential and non-essential cohesin genes and genes encoding for ribosomal subunits such as RPL15B, RPBL41B and RPBL19B. This is consistent with recent findings showing that defects in cohesin genes lead to defects in the production of ribosomal RNA and translation efficacy in both budding yeast and patient cells (Sun et al., 2015, Bose et al., 2012, Xu et al., 2014, Lu et al., 2014).

Our network also revealed several unanticipated interactions (Fig. 2). For example, several interactions between cohesin factors and genes involved in nucleotide excision repair, such as RAD16 and RAD1 with SMC1 and RAD10 with RAD61, in mismatch repair, such as MSH2 with MDC1 and RAD61, or in template switching, such as RAD5 with DCC1 and RMI1, might indicate a novel role for cohesin in post-replication repair. Supporting this notion, the separase complex is required for cohesin dissociation during post-replicative DNA repair (Nagao et al., 2004, McAleenan et al., 2013). Moreover, Smc1 is phosphorylated in an ATR-dependent manner after exposure to ultra violet (UV)-induced DNA damage and smc1-259 mutant shows a high sensitivity to UV (Garg et al., 2004, Kim et al., 2002). Finally, several other unanticipated interactions were found between cohesin factors and genes involved in microtubule organization and protein folding, highlighting potential novel functional connections. Taken together, our genetic interaction map provides a resource of known as well as novel interactions between cohesin and genes involved in various biological processes, which may serve as a starting point for unraveling cohesin functions in these processes.
Irc15 promotes the loading of centromeric cohesin

The cohesin interaction network may not only reveal new connections between cohesin genes and distinct biological processes, but may also uncover new factors involved in sister chromatid cohesion. Since genes acting in the same pathway tend to have similar genetic interaction profiles, we employed unsupervised hierarchical clustering of genetic interactions involving both cohesin and DDR related query genes (Fig. 3A, left panel). Strikingly, a cluster of array genes interacted specifically with the cohesin query genes, which clustered separately from the DDR query genes (Fig. 3A, right panel). Interestingly, within this cluster, genes implicated in the establishment of pericentromeric cohesion, namely CTF19, IML3 and CHL4, clustered together but did not interact with the three non-essential cohesin factors MRC1, TOF1 and ELG1. While this cluster furthermore included genes implicated in chromosome segregation (e.g. BIM1, MAD2 and BUB1), it was mostly dominated by genes involved in sister chromatid cohesion. Interestingly, among these genes were four genes GIM4, GIM3 and YKE2, all members of the prefoldin complex, and IRC15, a microtubule binding protein, whose role in this process was unknown. We confirmed the negative genetic interactions of gim3Δ, yke2Δ and irc15Δ with smc3-1, and of gim4Δ and yke2Δ with smc1-249 at semi-permissive temperature (Fig. S2).

To assess their role in sister chromatid cohesion, we first examined whether GIM4, GIM3, YKE2 and IRC15 affect the loading of cohesin onto chromosomes. PAC10, which encodes another member of the prefoldin complex, did not display any significant negative interaction with cohesin genes and was therefore included as a negative control. Scc1 loading was assessed by chromatin immunoprecipitation at known cohesin-binding sites in G2 cells (Fig. 3B,C). A region on chromosome III devoid of Scc1 was used as a negative control (Pal et al., 2018). Scc1 loading was comparable in wild type (WT) cells and cells lacking GIM3, GIM4, YKE2 or PAC10, suggesting that the prefoldin complex is not involved in cohesin loading. However, Scc1 levels were decreased at centromeric regions in the absence of IRC15, while they were increased on chromosome arms, indicating that Irc15 regulates the distribution of cohesin on chromosomes. The defect in centromeric cohesin loading in irc15Δ may stem from a translocation of cohesin from the centromeres to the chromosome arms. However, we could not detect any such translocation of Scc1 by ChIP when cells proceeded from G1-phase to G2/M-phase (Fig. S3A-F). Thus, we identify Irc15 as a new factor involved in the loading of centromeric cohesin.
Interestingly, *irc15Δ* cells present a delayed pre-anaphase mitotic entry due to defective kinetochore-microtubule attachments (Keyes and Burke, 2009). Potentially, reduced cohesin loading and consequently impaired sister chromatid cohesion may have affected the maintenance of kinetochore-microtubule attachments during mitosis. To address this, we examined whether overexpression of Scc1 could rescue the kinetochore assembly defects observed in the absence of *IRC15* (Keyes and Burke, 2009). To this end, we monitored binding of the kinetochore-associated Ndc80 complex, which is involved in kinetochore assembly (McCleland et al., 2003), by ChIP of GFP-tagged Ndc80 at four different centromeres (CEN2, CEN3, CEN4 and CEN8) and a negative control locus (Neg1p2) (Lefrancois et al., 2013) in WT and *irc15Δ* strains carrying a galactose-inducible allele of *SCC1* (Fig. S3G). We found that Ndc80 binding was increased around 4-fold in the absence of *IRC15* (Fig. S3H), indicative of a kinetochore assembly problem and agreeing with a previous observation (Keyes and Burke, 2009). Importantly, Ndc80 binding was not affected by Scc1 overexpression (Fig. S3H), suggesting that reduced cohesin loading in the absence of *IRC15* may not affect the maintenance of kinetochore-microtubule attachments.

**The prefoldin complex is involved in sister chromatid cohesion**

While Irc15 promotes the loading of centromeric cohesin, its contribution to sister chromatid cohesion is unclear. Also unclear is whether the prefoldin complex affects this process. To examine this, we employed a strain in which a tandem LacO array was integrated 10 kb away from the *CEN4* locus and LacR-GFP protein is stably expressed (Fig. 4A). An increased number of G2/M cells with more than one GFP focus indicates a defect in sister chromatid cohesion in this strain (Fig. 4A,B). In our assays, a *kre1Δ* mutant defective in beta-glucan assembly was included as negative control, while *chl1Δ*, *bub1Δ* and *rts1Δ* mutants served as positive controls (Kitajima et al., 2005, Kitajima et al., 2006). As expected, two GFP foci were evident in ~10% of the *kre1Δ* cells in G2/M-phase, which was comparable to that in WT cells (Fig. 4C, top). In contrast, at least ~20% of the *chl1Δ*, *bub1Δ* and *rts1Δ* cells displayed two GPF foci, indicative a cohesion defect. Importantly, at least 20% of the *gim3Δ, gim4Δ, yke2Δ, pac10Δ* and *irc15Δ* cells showed more than two GFP foci, suggesting a defect in sister chromatid cohesion. Noteworthy, an increased number of the
prefoldin mutant cells also harbored two GFP spots in G1-phase. This may result from chromosome mis-segregation during the previous mitosis, which might be a consequence of defective cohesion (Hoque and Ishikawa, 2002, Sonoda et al., 2001), although we could not detect any aneuploidy in these mutants (Fig. 4C, bottom), likely due to the low frequency of these events (<10%). To determine whether the prefoldin holocomplex is involved in cohesion establishment, we compared sister chromatid cohesion in gim4Δ and yke2Δ single and double mutants (Fig. 4D). gim4Δ and yke2Δ were epistatic with regard to their cohesion defect, suggesting that the prefoldin complex as a whole functions in the same pathway for cohesion establishment. In addition, we also evaluated if Irc15 functions in one of the two parallel non-essential cohesion pathways or defines a new cohesion pathway (Xu et al., 2007). To this end, we generated double mutants of IRC15 with CHL1 or MRC1, which encode components of the cohesion pathways involving Csm3 and Ctf18-RFC, respectively (Xu et al., 2007). While irc15Δ was epistatic with mrc1Δ, it displayed additive cohesion defects with chl1Δ. These results suggest that Irc15 functions with Mrc1 in the cohesion pathway involving Ctf18-RFC. Finally, we compared the resumption of cell cycle progression of irc15Δ and the prefoldin mutants following a G2/M arrest. Although WT cells progressed through mitosis and started to enter G1 by 60 minutes, the majority of the irc15Δ and prefoldin mutant cells were still in mitosis at that time, showing a clear delay in cell cycle progression (Fig. 4F), consistent with a sister chromatid cohesion defect (Sonoda et al., 2001). Thus, we reveal that Irc15 and the prefoldin complex promote efficient sister chromatid cohesion. While Irc15 promotes this process likely by facilitating the loading of centromeric cohesin, unclear is how the prefoldin complex would affect this process. Given that prefoldin delivers unfolded proteins to cytosolic chaperonins (Vainberg et al., 1998), we checked whether it may affect the stability of the cohesin core subunits. However, Smc1, Smc3, Scc1 and Scc3 stability remained unaffected in gim3Δ cells (Fig. S4).

**Discussion**

Here, we generated a comprehensive genetic interaction network centered on cohesin comprising 373 genetic interactions specific for cohesin factors. The network uncovered novel connections for cohesin genes in various cellular processes. Moreover, it also revealed new factors involved in sister chromatid cohesion, namely
the microtubule-associated protein Irc15 and the prefoldin complex members Gim3, Gim4 and Yke2. Thus, our genetic interaction map provides a unique resource for further identification and functional interrogation of cohesin proteins.

Irc15 was initially identified in different screens that were designed to identify factors involved in chromosome segregation and DNA repair (Alvaro et al., 2007, Measday et al., 2005, Daniel et al., 2006, Jordan et al., 2007). It was also shown that Irc15 associates with microtubules, regulating their dynamics and mediating tension between kinetochores (Keyes and Burke, 2009). Here, we identified a novel role for Irc15 in centromeric cohesin loading and cohesion establishment. Proper centromeric cohesion is a prerequisite to generate a dynamic tension between microtubules and sister chromatids in yeast (Goshima and Yanagida, 2000, He et al., 2000, Tanaka et al., 2000). This tension is also required for the establishment of stable microtubule-kinetochore attachments (Ault and Nicklas, 1989, Nicklas and Ward, 1994, Koshland et al., 1988, Skibbens et al., 1995). Indeed, loss of Scc1 impairs both sister chromatid cohesion and kinetochore function in higher eukaryotes (Sonoda et al., 2001). However, in the case of irc15Δ our results suggest that the kinetochore defect did not result from the cohesin loading defect observed in this mutant background. Conversely, several inner and central kinetochore proteins play a role in the recruitment of pericentromeric cohesin (Eckert et al., 2007, Hinshaw et al., 2017). However, cells with defective microtubule-kinetochore attachments exhibit high levels of Scc1 loading at centromeres (Eckert et al., 2007). Given that Irc15 controls tension between kinetochores and microtubules (Keyes and Burke, 2009) and that we observed a decrease in centromeric cohesin loading in the absence of IRC15, it is thus unlikely that the cohesion defect in irc15Δ cells stems from a kinetochore defect. Rather, Irc15 may play independent roles in cohesin loading and microtubule-kinetochore attachment at centromeres.

We also identified the prefoldin complex as a new factor involved in sister chromatid cohesion. The prefoldin complex is a multi-subunit chaperone that assists in the proper folding of proteins in the cytosol (Vainberg et al., 1998). Even though it did not affect the stability of the cohesin core subunits, it is tempting to speculate that prefoldin targets one or more (other) factors involved in sister chromatid cohesion, thereby affecting this process. Alternatively, the involvement of the prefoldin complex in cohesion might also be related to its role in regulating chromatin structure during transcription elongation (Millan-Zambrano et al., 2013). To this end, it may either
influence the transcription of genes involved in cohesion or allow the loading of the cohesin complex by generating nucleosome-free region at transcribed genes (Millan-Zambrano et al., 2013). This hypothesis is supported by our genetic interaction network, which identified a strong relationship between cohesin factors and factors involved in gene expression and/or chromatin remodeling. To this end, it is interesting to note that the RSC remodeling complex facilitates the association of cohesin on chromosome arms by generating a nucleosome-free region (Huang et al., 2004, Huang and Laurent, 2004, Munoz et al., 2019). Moreover, the SWR1 complex deposits the histone variant H2A.Z, whose acetylation maintains sister chromatin cohesion levels (Sharma et al., 2013). Finally, it was also shown that the NAD+-dependent deacetylase Hst3, member of the sirtuin superfamily, is involved in sister chromatid cohesion through the acetylation of histone H3 at lysine K56 (Thaminy et al., 2007), and that strains harboring mutations in cohesin genes are sensitive to sirtuin inhibitors (Choy et al., 2015). These findings may enforce a potential link between prefoldin and chromatin remodeling in cohesion establishment.

Among the novel connections for cohesin genes, we identified several interactions linked to post-replication repair and nucleotide excision repair. Further studies may reveal the functional importance of the link between sister chromatid cohesion and these processes. Since defects in nucleotide excision repair are associated with Cockayne Syndrome or Xeroderma Pigmentosum, we anticipate that the link between cohesin factors and this repair process may be relevant for disease etiology. In line with this, it was recently shown that the nucleotide excision repair structure-specific endonuclease ERCC1–XPF complex interacts with the cohesin complex and other proteins at promoters to silence imprinted genes during development in mice (Chatzinikolaou et al., 2017). Moreover, since sister chromatid cohesion and the factors involved are well conserved from yeast to men (Xiong and Gerton, 2010), our network may also inform on genetic interactions of cohesin factors mutated in cohesinopathies or cancer.
Materials & Methods

Genetic interaction map analysis
The genetic interaction map was generated and analyzed as previously described (Srivas et al., 2013). Briefly, an array of 1494 genes (Table S2) was collected from the yeast deletion collection mat alpha and the DAmp library containing KANMX selection marker. To generate the query genes (Table S1), mutant strains carrying deletion mutations were generated by PCR gene targeting (Longtine et al., 1998), while mutants carrying point mutations were either generated using the MIRAGE method (Nair and Zhao, 2009) in a strain containing synthetic genetic array (SGA) anti-diploid selection markers and a NATMX selection marker, or obtained from Charles Boone and Philip Hieter lab. Primers used to generate these mutants are available upon request. Due to the presence of temperature sensitive mutants, the generation of double mutants was performed at permissive temperature (23°C) with use of the SGA procedure in quadruplicate using the ROTOR HDA (Singer Instruments) pinning robot (Tong and Boone, 2006). Genetic interactions were assessed at semi-permissive temperature (30°C). Pictures were taken with a Canon Powershot G3. Colonies sizes were quantified and normalized using Matlab Colony Analyzer. Quantitative S-scores were calculated using Matlab as previously described (Collins et al., 2010, Guenole et al., 2013). Network visualizations of genetic interactions were done using Cytoscape (Shannon et al., 2003). A Cytoscape plugin BiNGO was used for GO term enrichment analysis (Maere et al., 2005). Unsupervised clustering was performed using Cluster 3.0 using a selection of array genes that show magnitude of S-score > 2.0 in at least one of the query genes and with a variation with a standard deviation > 0.8 in the query genes. The clustering was visualized in a heatmap using Java TreeView.

Yeast strains and culture conditions
A strain expressing 18Myc-tagged Scc1 and HA-tagged Pds1 was used in flow cytometry and Scc1-based ChIP experiments. PCR gene targeting was used to generate the tagged alleles and gene deletions (Table S8). A strain carrying a LacO array integrated on chromosome IV 10 kb away from CEN4 and expressing a LacR-GFP fusion protein was used for sister chromatid cohesion assays (Shimada and Gasser, 2007). PCR gene targeting was used to generate gene deletions in this
background (Table S8). Primers used to generate yeast strains are available upon request. All yeast strains were cultured in rich YPAD medium or Synthetic Complete medium lacking methionine (SC-methionine).

**Chromatin immunoprecipitation**

Chromatin immunoprecipitation (ChIP) was performed as previously described with slight modifications (Cobb et al., 2003). Briefly, cells were grown to $5 \times 10^6$ cells/ml in YPAD and synchronized in G2/M by incubation with nocodazole (7.5 µg/ml) for 2 hours for Scc1 ChIP. Nocodazole (7.5 µg/ml) was added a second time after 1 hour of incubation. Alternatively, cells were synchronized in G1 with α-factor for 2 hours, washed and released in YPAD containing nocodazole for 0, 30, 60, 90 and 120 minutes. Samples were fixed with 1 % formaldehyde. For Ndc80-GFP ChIP, cells were grown overnight in SC-methionine containing 2% raffinose, diluted and grown in the presence of 2% glucose or 2% galactose for 4 hours, diluted to $5 \times 10^6$ cells/ml and fixed with 1% formaldehyde. Extracts were prepared in lysis buffer (50mM Hepes, pH=7.5, 140 mM NaCl, 1 mM Na EDTA, 1% trition x-100, 0.1 % Na deoxycholate) containing protease inhibitors. Extracts were subjected to immunoprecipitation with Dynabeads mouse or rabbit IgG (Invitrogen, M-280) coated with antibody against c-Myc (9B11, Cell Signaling) or GFP (ab290, Abcam). DNA was purified and enrichment at specific loci was measured using qPCR. Relative enrichment was determined by $2^{-\Delta\Delta C_T}$ method (Livak and Schmittgen, 2001, Cobb and van Attikum, 2010). Dynabeads alone were used to correct for background. An amplicon 11 kb downstream of ARS305, devoid of Scc1 binding, was used for Scc1 ChIP normalization (Tittel-Elmer et al., 2012). An amplicon devoid of Ndc80 binding (Neg1p1) was used for Ndc80 ChIP normalization (Lefrancois et al., 2013). Primers used are listed in Table S9.

**Sister chromatid cohesion assay**

Sister chromatid cohesion was assayed using a strain containing a LacO repeat integrated at chromosome 4 between *ARS1* and *CEN4* at 10 kb distance to *CEN4* and LacR-GFP integrated at *HIS3* locus (Shimada and Gasser, 2007). Cells were grown to midlog in YPAD, synchronized in G1 by incubation with α-factor for 1.5 hours, or in G2/M by incubation with nocodazole (15 µg/ml) for 1 hour. Cells were
fixed in 4% paraformaldehyde at room temperature for 15 minutes, washed and resuspended in KPO4/Sorbitol solution (10 mM KPO4, 1.2 M Sorbitol, pH=7.5). Images of cells were acquired on a Zeiss AxiolImager M2 widefield fluorescence microscope equipped with 100x PLAN APO (1.4 NA) oil-immersion objectives (Zeiss) and an HXP 120 metal-halide lamp used for excitation. Fluorescent signals were detected using the following filters: GFP/YFP 488 (excitation filter: 470/40 nm, dichroic mirror: 495 nm, emission filter: 525/50 nm). Images were recorded and analyzed using ZEN 2012 software.

**Flow cytometry**

Cells were grown to midlog in YPAD, synchronized in G1 by incubation with α-factor for 1.5 hours, or in G2/M by incubation with nocodazole (15 µg/ml) for 1 hour. Alternatively, cells were grown to midlog in YPAD, synchronized in G2/M by incubation with nocodazole (15 µg/ml) for 2 hours, washed and released in YPAD. Samples were prepared as previously described (Haase and Lew, 1997). Data were acquired on a BD FACSCalibur (BD Biosciences) or on a Novocyte (ACEA Biosciences, Inc) and analyzed with FlowJo or NovoExpress software, respectively.

**Spot dilution test**

Cells were grown overnight in YPAD and then plated in fivefold serial dilutions starting at a density of $6 \times 10^6$ cells/ml (OD600 nm= 0.5) on YPAD plates. Cells were grown for 3 days at semi-permissive temperature (30°C) before images were taken.

**Cycloheximide chase experiment**

Cells expressing Scc1-18Myc, Scc3-6FLAG, Smc1-6FLAG or Smc3-6FLAG were subjected to cycloheximide chase analysis as previously described (Buchanan et al., 2016). Samples were collected at 0, 30, 60 and 90 minutes after cycloheximide treatment. Whole cell extracts were prepared by post-alkaline protein extraction and analyzed by SDS-PAGE. Western blotting was performed using a c-Myc antibody (9E10, Santa Cruz Biotechnology) and FLAG antibody (clone M2, Sigma). Ponceau staining served as a loading control.
Curation of *S. cerevisiae* - *S. pombe* and *S. cerevisiae* - *H. sapiens* orthologs

Information about budding yeast-to-human and budding yeast-to-fission yeast orthologs was collected from two different sources, InParanoid (O’Brien et al., 2005) and PomBase (Lock et al., 2018), and is presented in Table S7. InParanoid inventories orthologs based on protein sequence similarity, whereas PomBase curates orthologs based on both function and sequence similarity.

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Competing interests

The authors declare no competing or financial interests.

Author contributions


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Figure 1. A genetic interaction map centered on cohesin

(A) Outline of the genetic interaction screen. Mutants in 17 cohesin and 18 DNA damage response (DDR) query genes were crossed against a panel of 1494 mutants in array genes involved in various biological processes. Genetic interactions were scored by quantification of colony sizes, providing each double mutant with a quantitative S-score.

(B) Total number of positive (S-score ≥ 2) and negative (S-score ≤ -2.5) interactions for all query (top), cohesin (bottom left) or DDR (bottom right) genes.

(C) GO term enrichment of interactions involving all (left), cohesin (middle) or DDR genes (right).
Figure 2. A genetic interaction network centered on cohesin

Visualization of significant genetic interactions of cohesin-related genes. Interacting genes were grouped based on GO annotation.
Figure 3. Identification of new cohesin factors with Irc15 as cohesin loader

(A) Heatmap displaying hierarchical clustering of genetic interactions scores (S-scores; left panel) identified a cluster of negative interactions involving cohesin factors and genes involved in chromosome segregation (right panel; blue = negative interaction, yellow = positive interaction, black = neutral interaction, grey = missing interaction). Potential new sister chromatid cohesion factors are highlighted in red.

(B) Schematic of chromosomal loci assayed for Scc1 loading. qPCR was performed at known cohesin binding sites either on centromeres (CEN9; CEN3) or genic (POA1; MRP10; MET10) and intergenic (Conv 32W-31C) regions on chromosome arms. ChrIII neg was a negative control.

(C) Enrichment of Scc1-Myc assessed by ChIP-qPCR at the indicated loci in nocodazole-arrested strains. Enrichment corresponds to the ratio of the Scc1-Myc signal over beads alone. Average enrichment with standard error of the mean of 3 (gim3Δ, gim4Δ, yke2Δ and pac10Δ) or 4 (WT; irc15Δ) independent experiments is shown. Asterisks indicate statistical differences using a Student t-test (* = p < 0.05; ** = p < 0.01).
Figure 4. The prefoldin complex and lrc15 affect cohesion establishment

(A) Schematic of the sister chromatid cohesion assay. A LacO array was integrated on chromosome IV 10 kb away from CEN4 in cells expressing LacR-GFP fusion protein. Upon synchronization of the cells in G1 with α-factor or in G2/M with...
nocodazole, cells with normal sister chromatid cohesion show one spot in G1 and G2/M in the majority of the cells. Cohesin mutants show a larger fraction of cells with two GFP spots.

(B) Representative images of the sister chromatid cohesion assay in nocodazole-arrested cells.

(C) Quantification of sister chromatid cohesion in cells from (B). Mean percentage of cells with more than one GFP spot (top) is shown. Around 400 cells were scored in at least 3 independent experiments for each strain. Error bars represent the standard error of the mean. Flow cytometry analysis of DNA content was used to monitor cell synchronization (bottom).

(D-E) Quantification of sister chromatid cohesion in the indicated cells as in (B).

(F) Flow cytometry analysis of M-phase progression of the indicated strains. Cells were arrested in G2/M by nocodazole treatment, released in YPAD and analyzed at the indicated timepoints.
Sun et al Figure S1

A

\[ r = 0.421, r^2 = 0.230, n = 29 \]

Query_B::NATMX x Array_A::KANMX

Query_A::NATMX x Array_B::KANMX

B

Correlation

C

Fold enrichment over random

Top % of Gene Pairs (Ranked by \( S \))

D

S-score

- Rank based on S-score
- Divide in 6 groups
- Generate top 10 double mutants de novo

Compare observed fitness to estimated fitness

E

Example of validation for \( ctf4 \)\( \Delta \) \( sir1 \)\( \Delta \) (cat. = 2)

F

<table>
<thead>
<tr>
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<td>Total</td>
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</tr>
</tbody>
</table>

G

\( P_{\text{adj}} < 0.05 \)

- FDR negative interactions: 25.0%
- FDR positive interactions: 53.3%
- FDR average: 31.0%
Figure S1. Quality control of the genetic interaction map

(A) Correlation of S-scores from genetic interactions in double mutants generated by reciprocal crossings of query and array strains: query gene A::NATMX x array gene B::KANMX and query gene A::KANMX, array gene B::NATMX.

(B) Correlation of genetic interaction scores of common interactions between this study and other studies.

(C) Fold enrichment for genetic interactions that are present in the Biogrid database (version 3.2; (Stark et al., 2006)) is shown. Fold enrichment is defined as n/r, where n is the number of highest scoring genetic interactions (x-axis) found in the Biogrid database, while r is the number of overlapping interactions expected at random.

(D) Outline of the validation of genetic interactions with S-scores ≥ 2 or S-scores ≤ -2.5. Negative interactions were ranked and divided in 6 groups. Double mutants corresponding to the top 10 interactions within each group were generated de novo in multitude using SGA technology (n>8). The fitness of double mutants was compared to the estimated fitness of the double mutants using t-test statistics and correction for multiple testing (Benjamini-Hochberg). Estimated fitness of the double mutant was calculated by combining fitness of the corresponding single mutants (Mani et al., 2008), which were also generated de novo by crossing them with WT dummy strains containing either a his3::KANMX or his3::NATMX allele. Positive interactions were validated in the same way, except that only the top 15 interactions were tested.

(E) Example of the outcome of the validation of the negative interaction between CTF4 and SIR1.

(F) Table showing the outcome of the validation of 71 genetic interactions.

(G) Table showing the false discovery rates based on the validation of 71 genetic interactions.
Figure S2. Validation of negative genetic interactions for IRC15, GIM3, GIM4 and YKE2.

Drop assay for the indicated strains. Ten-fold serial dilutions of exponentially growing cells were spotted on rich medium and incubated at semi-permissive temperature (30°C).
Sun et al_Figure S3

A) POA1

B) MRP10

C) MET10

D) Conv 32W-31C

E) CEN9

F) CEN3

G) Western blot analysis

H) Ndc80 ChIP

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Figure S3. Cohesion defects in irc15Δ do not stem from cohesin translocation from centromeres to chromosome arms and do not result in kinetochore defects.

(A-F) Enrichment of Scc1-Myc assessed by ChIP-qPCR at POA1 (A), MRP10 (B), MET10 (C), Conv 32W-31C (D), CEN9 (E) and CEN3 (F) in WT and irc15Δ cells. Strains were arrested in G1 and released in nocodazole. Enrichment corresponds to the ratio of the Scc1-Myc signal over beads alone. Average enrichment with standard error of the mean of 3 independent experiments is shown.

(G) Western blot analysis of Scc1-HA expression after incubation in medium with galactose (Gal; Scc1-HA overexpression) or glucose (Glu; Scc1-HA repression).

(H) Enrichment of Ndc80-GFP assessed by ChIP-qPCR at four centromeres in the indicated strains in presence of glucose (Scc1-HA repression) or galactose (Scc1-HA overexpression). Enrichment corresponds to the ratio of the Ndc80-GFP signal at CEN2, CEN3, CEN4, CEN8 and Neg1p2 over Neg1p1 in GFP IPs over IPs with beads alone. Average enrichment with standard error of the mean of 2 (glucose) or 3 (galactose) independent experiments is shown.
Figure S4. Cohesin core subunits stability is not affected in gim3Δ cells.

(A-D) Western blot analysis and quantification of Scc1-Myc (A), Scc3-Flag (B), Smc1-Flag (C) and Smc3-Flag (D) levels in WT and gim3Δ cells. The indicated strains were subjected to cycloheximide chase analysis. Ponceau staining served as loading control. Signal intensities were adjusted to Ponceau staining. The ratio was set to 100% for the first time point for each strain and subsequent time points were normalized to it.
Table S1: List of query genes
    Click here to Download Table S1

Table S2: List of array genes
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Table S3: S-scores of the cohesin/DDR interaction map
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Table S4: Gene Ontology analysis of the cohesin/DDR interaction map
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Table S5: Gene Ontology analysis of cohesin-related genes
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Table S6: Gene Ontology analysis of DDR-related genes
    Click here to Download Table S6

Table S7: Ortholog information for the cohesin interaction network
    Click here to Download Table S7

Table S8: Yeast strains used in this study
    Click here to Download Table S8

Table S9: List of primers used for qPCR
    Click here to Download Table S9