Global Analysis of SUMO-Binding Proteins Identifies SUMOylation as a Key Regulator of the INO80 Chromatin Remodeling Complex

Eric Cox\textsuperscript{a,h,c}, Woochang Hwang\textsuperscript{d}, Ijeoma Uzoma\textsuperscript{c}, Jianfei Hu\textsuperscript{d}, Catherine M. Guzzo\textsuperscript{e}, Junseop Jeong\textsuperscript{d}, Michael J. Matunis\textsuperscript{e}, Jiang Qian\textsuperscript{d}, Heng Zhu\textsuperscript{c,f,*} and Seth Blackshaw\textsuperscript{b,f,g,h,*}

From the \textsuperscript{a}Biochemistry, Cellular and Molecular Biology Graduate Program, \textsuperscript{b}Solomon H. Snyder Department of Neuroscience, \textsuperscript{c}Department of Pharmacology and Molecular Sciences, \textsuperscript{d}Wilmer Eye Institute, \textsuperscript{f}Center for Human Systems Biology, \textsuperscript{g}Institute for Cell Engineering, \textsuperscript{h}Department of Ophthalmology, Johns Hopkins University School of Medicine, Baltimore, Maryland, \textsuperscript{e}Department of Biochemistry and Molecular Biology, Bloomberg School of Public Health, Johns Hopkins University, Baltimore, Maryland

*To whom correspondence should be addressed at sblack@jhmi.edu (S.B), hzhu4@jhmi.edu (H.Z.), and jiang.qian@jhmi.edu (J.Q.)

ABSTRACT
SUMOylation is a critical regulator of a broad range of cellular processes, and is thought to do so in part by modulation of protein interaction. To comprehensively identify human proteins whose interaction is modulated by SUMOylation, we developed an \textit{in vitro} binding assay using human proteome microarrays to identify targets of SUMO1 and SUMO2. We then integrated these results with protein SUMOylation and protein-protein interaction data to perform network motif analysis. We focused on a single network motif we termed a SUMOmodPPI (SUMO-modulated Protein-Protein Interaction) that included the INO80 chromatin remodeling complex subunits TFPT and INO80E. We validated the SUMO-binding activity of INO80E, and showed that TFPT is a SUMO substrate both \textit{in vitro} and \textit{in vivo}. We then demonstrated a key role for SUMOylation in mediating the interaction between these two proteins, both \textit{in vitro} and \textit{in vivo}. By demonstrating a key role for SUMOylation in regulating the INO80 chromatin remodeling complex, this work illustrates the power of bioinformatics analysis of large datasets in predicting novel biological phenomena.

INTRODUCTION
The functional protein microarray is a powerful and versatile systems biology and proteomics tool that allows the profiling of the activity of thousands of proteins in parallel. Applications of functional protein microarrays range from the profiling of protein interactions, construction of enzyme-substrate networks, and identification of novel enzymatic activities. Since the development of the yeast proteome microarray over 15 years ago (1), more recent work has seen the development of complete and near-complete proteome arrays representing viruses, bacteria and plants (2-4). However, most existing human protein microarrays are comprised of only a minority of the human proteome (5-9). We have recently developed a human proteome microarray, the HuProt array, which includes >17,000 full-length human proteins (10). The proteins used to generate this microarray were expressed in yeast and purified under native conditions.
Expressing recombinant eukaryotic proteins in yeast improves the likelihood that proteins will retain their biological activity relative to prokaryotic and in vitro expression systems. Numerous studies have harnessed the power of the HuProt array to profile a wide range of protein activities, including protein-protein interactions (11, 12), protein-RNA interactions (13, 14), analysis of monoclonal antibody specificity (10), biomarker discovery (15), identification of substrates of protein kinases (16) and S-nitrosylation (17).

Recently, we have conducted SUMOylation assays using the HuProt microarray to identify numerous previously uncharacterized SUMO E3 ligase-dependent substrates using a subset of human SUMO E3 ligases (18). Protein SUMOylation is an essential posttranslational modification in most organisms, including yeast (19), C. elegans (20), Arabidopsis (21), and mice (22, 23). The reversible SUMO-modification of target proteins involves an enzymatic cascade chemically similar to ubiquitylation, involving E1 activating, E2 conjugating, and E3 ligating enzymes, and SUMO proteases. As the only classes of SUMOylation enzymes for which multiple members have been identified, the SUMO E3 ligases and the SUMO proteases have been proposed to be the major factors determining substrate specificity.

Although SUMOylation can regulate a diverse array of protein activities, at the molecular level SUMOylation alters protein surfaces and their ability to interact with other proteins (23). It has been proposed that SUMO acts like a “molecular glue” that mediates non-covalent interactions between modified substrates and SUMO-binding proteins (24). This is illustrated by the interaction between the promyelocytic leukemia (PML) protein and thymine DNA glycosylase (TDG), in which SUMO-modification as well as SUMO-interaction motifs (SIMs) on both proteins contribute to binding (24). Similarly, SUMO1 modification of RanGAP1 promotes its interaction with RanBP2 (25). The canonical SUMO-interaction motif (SIM) contains a hydrophobic core consisting of four hydrophobic amino acids that can be either preceded or followed by multiple negatively charged amino acids. Recently, other protein motifs have been shown to mediate SUMO-binding, including MYM-type and ZZ-type zinc finger domains (26, 27).

Previous efforts to identify SUMO-binding proteins were limited either by their focus on a small number of cell or tissue types, or else were restricted to identifying nuclear proteins that bound to only a single SUMO isoform (24, 28, 29). One such study relied on the use of an MS-coupled pulldown (30), while others used yeast two-hybrid analysis, which typically correlates only moderately well with biochemical protein-protein interaction assays (24, 28). These methods, while powerful, often do not distinguish between direct and indirect interactions.

For our studies, we utilized the HuProt microarray containing >17,000 individually purified human proteins, thus avoiding bias in favor of a particular cell or tissue type or subcellular compartment, and allowing identification of direct and/or low-abundance target proteins. We also looked at binding to both SUMO1 and SUMO2 monomers, as well as SUMO1 and SUMO2 trimers, which were used as model SUMO chains. Our dataset represents a nearly two-fold increase in the number of known SUMO-
interacting proteins. The vast majority of these proteins had not been previously reported as having SUMO binding activity.

To use our data set to make novel predictions about SUMOylation function, we integrated our SUMO-binding and SUMOylation data with protein-protein interaction data from publicly available databases to perform network motif analysis. We identified 21 three-component network motifs. Nine of these network motifs were significantly enriched in this data set, while four network motifs were significantly depleted. We focused on a single network motif containing a SUMO-binding protein and a SUMO-modified protein that were previously reported to interact, along with SUMO itself (31, 32). This specific network motif suggests a potential role for SUMO in modulating the protein-protein interaction between the other 2 nodes; and we thus termed this motif a SUMOmodPPI (SUMO-modulated Protein-Protein Interaction). We then validated an example of this network motif that comprised the INO80 chromatin remodeling complex subunits INO80E and TFPT, along with SUMO2. We found that TFPT could be SUMOylated in transfected cells and identified the relevant SUMOylation site on the protein. We also validated the SUMO-binding activity of INO80E and identified the region of INO80E important for this interaction. Finally, we demonstrated that SUMO helps facilitate the interaction between INO80E and TFPT. This work demonstrates the power of integrative analysis of large datasets in predicting novel biological phenomena.

RESULTS

Identification of SUMO-binding partners using HuProt arrays

Previous work reported that the first alpha helix and second beta strand of both SUMO1 and SUMO2 can act as a SUMO paralog recognition region that interacts with the SUMO interaction motifs (SIMs) of SUMO paralog-specific binding proteins (26, 33, 34). Although only a handful of SUMO paralog-specific binding proteins have been identified to date, this suggests that more paralog-specific binding proteins remain to be discovered. In order to identify novel SUMO paralog-specific binding proteins, we purified SUMO1 and SUMO2 monomers, as well as SUMO1 and SUMO2 trimers, to be used as polySUMO chain mimics (Figure 1A). We then used these protein probes in a novel human proteome microarray-based SUMO-binding assay (Figure 1B). For detection of SUMO binding proteins on the HuProt microarrays, we prepared fluorescently labeled SUMO isoform-specific monoclonal antibodies. We then performed SUMO-binding assays in triplicate for each of the four probes. Among 457 total SUMO binding proteins that were identified by our assay, only 13 had been previously identified among the 270 previously known SUMO-binding proteins (Dataset S1 and Supplemental Fig. S1). 11 of these 13 previously identified SUMO-binding proteins were found by two SUMO2-focused affinity purification and mass spectrometry based screens (29, 35). The four SUMO protein probes that we tested showed wide variation in their binding specificity (Figure 2A). The SUMO2 monomer probe bound to the largest number of proteins (306), as well as the largest number of unique proteins (128). The SUMO1 trimer bound 197, the SUMO1 monomer bound 183, and the SUMO2 trimer bound 139 proteins in total. The pairwise probe combinations of SUMO1 trimer and SUMO2 monomer, and SUMO1 monomer and SUMO2 monomer, each bound the largest number
of proteins (108). On the other hand, the SUMO1 trimer and SUMO2 trimer bound to the smallest number of shared proteins (66). 39 proteins bound to all four SUMO probes.

**GO analysis reveals potential new function of SUMOylation**

We next conducted gene ontology (GO) analysis on the novel SUMO binding proteins (Figure 2B). This analysis revealed several notable properties of these proteins. With regard to cellular component, we observed significant enrichment for cytoplasmic localization among proteins that were bound by all four probes. Considering that many previously identified SUMO-modified proteins are localized to the nucleus, including many transcription factors, this enrichment of cytoplasmic proteins suggests that the importance of SUMOylation in the cytoplasm may be underappreciated.

We also found that some GO categories that were selectively enriched among proteins that bound to all 4 probes, different combinations of 2 or 3 probes, or else a single probe. For example, proteins associated with the GO categories of “actin binding”, “catalytic activity” and “oxidoreductase activity” were selectively enriched among targets bound by all 4 probes. A particularly surprising result of this analysis was the finding that of all pairwise combinations of probes tested, proteins that bound the SUMO1 trimer and SUMO2 monomer, which are two of the most dissimilar probes, shared the largest number of molecular function GO terms.

With regard to biological process, enriched GO terms common to proteins that bound all 4 probes included 3 terms: “metabolic process”, “muscle contraction” and “oxidation reduction”. Among the proteins that we identified that are associated with muscle contraction, including myoglobin, myosin light chain 6B, sarcospan and alpha-1-syntrophin, each of these contained at least one predicted canonical SIM. Sarcospan and alpha-1-syntrophin are members of the dystrophin-glycoprotein complex (DGC), which plays a role in membrane integrity and signal transduction in heart and skeletal muscle and links the actin cytoskeleton to various receptors. Although a role for SUMO in muscle function has been largely unexplored, skeletal muscle actin is SUMOylated and altered cellular localization of SUMO1 has been observed in skeletal muscle in response to physical exercise (36, 37). As with molecular function, the SUMO1 trimer and SUMO2 monomer binding proteins likewise shared the largest number of GO categories compared to any other pairwise probe combination.

**Analysis of integrated SUMO networks**

We recently conducted SUMOylation assays using the HuProt microarray to identify numerous previously uncharacterized SUMO E3 ligase-dependent substrates using a subset of human SUMO E3 ligases (Data Set S2). We hypothesized that by integrating our SUMO-binding data with our in vitro SUMOylation data (18), as well as protein-protein interaction data from publicly available databases, we could make novel predictions about SUMOylation function. This combined dataset includes a total of 2910 high-confidence SUMOylation substrates, 489 high-confidence SUMO binding proteins, and 6121 reported protein interactions for proteins that are either SUMOylation substrates or SUMO binding proteins, or both. The integration of these three types of datasets resulted in a network of 2503 nodes, connected by 9520 edges. To gain biological insights from this complex network, we searched for tripartite network motifs that were statistically overrepresented. These network motifs can be considered to be the
basic building blocks for the network and provide the information underlying the design principle of SUMO-dependent networks. We can describe each network motif using two parameters, \( n \), the number of occurrences of the motif, and \( Z \), an enrichment score (Figure 3, supplemental fig. 4 and Data Set S3). This enrichment score reflects the difference between the observed occurrence of this network motif in the integrated SUMOylation network and its expected occurrence in a set of random networks (38). In these network motifs, at least one node represents SUMO and at least one node represents a protein from either the SUMO-binding or SUMOylation datasets. These 3 proteins are then connected by edges that represent one of three types of protein-protein interaction, including 1) noncovalent SUMO-binding identified in our assay or 2) covalent SUMOylation, or 3) a protein-protein interaction from one of several publicly available databases. Among the 21 classes of tripartite network motifs, we found nine enriched (\( Z>0 \)) and four depleted (\( Z<0 \)) motifs (Figure 3). One representative example is shown for each class of the tripartite motif. For example, network motif class 10 represents two interacting proteins that are both SUMOylated. One such instance of this network motif includes the proteins TWF2 and CAPZA2, which interact with each other and are both SUMOylated by SUMO1. In total, we found 4,377 instances of this motif (\( n=4377 \)), and this number of occurrences is significantly more than what we would expect from a random network, as indicated by a positive enrichment score (\( Z=3.9 \)).

**INO80E, TFPT, and SUMO2 form the SUMOmodPPI network motif**

We then selected one network motif that included two proteins previously reported to interact and included a novel SUMO-binding protein (this dataset) and a SUMOylated protein (38). This network motif immediately suggests a potential role for SUMO in modulating this protein-protein interaction and thus, we termed this network motif a “SUMO-modulated protein-protein interaction” (SUMOmodPPI). One such SUMOmodPPI consists of the INO80 chromatin remodeling complex subunits INO80E and TFPT, and SUMO2 (Figure 3). Our proteome microarray assay revealed that INO80E binds specifically to the SUMO2 monomer and the SUMO2 trimer. The other protein in this triad, TFPT, was previously found to be specifically modified by SUMO2 in the presence of the SUMO E3 ligases PIAS1 and PIAS3 (38). As a candidate SUMOmodPPI, we hypothesized that SUMO may play a role in modulating the interaction between INO80E and TFPT. Before we could proceed with testing our hypothesis, we first set out to validate their SUMO-binding and SUMOylation properties, using conventional assays.

**INO80E preferentially interacts with SUMO2 via its SIM domain**

First, we evaluated the binding of INO80E, translated \textit{in vitro} in rabbit reticulocyte lysate in the presence of \([^{35}S]\)-methionine, to immobilized GST-tagged SUMO1 or SUMO2 polymers. We observed a strong interaction between INO80E and the SUMO2 polymer in contrast to a very weak interaction between INO80E and the SUMO1 polymer (Figure 4A), in close agreement with our human proteome microarray data.

Visual inspection of the amino acid sequence of INO80E revealed a canonical SUMO-interaction motif (SIM) at the protein C-terminus containing a hydrophobic core
“VIDI” preceded by 3 negatively charged aspartate residues (Figure 4B). In order to test whether this predicted SIM was important for SUMO binding, we made two mutant forms of the INO80E protein. In the first mutant, we mutated the hydrophobic residues in the hydrophobic core of the SIM to alanines (mtSIM). In the second mutant, we made a truncation mutant that lacked the ten C-terminal residues, including the hydrophobic core sequence and the preceding negatively charged residues (ΔC10). The mtSIM mutant showed a 70% reduction in binding of INO80E to GST-SUMO2, while binding to GST-SUMO1 was unchanged (Figure 4C). However, while the INO80E truncation mutant (ΔC10) lacking the ten C-terminal amino acids showed a 20% reduction in binding to GST-SUMO1, it showed an 80% reduction in binding to GST-SUMO2, approaching the lower limit of detection in our assay. This stands in contrast to previous work reporting the importance of acidic residues specifically in SUMO1 binding (28). This result demonstrates the importance of both the charged and hydrophobic residues in mediating interaction between INO80E and SUMO, both for SUMO1 and, particularly, for SUMO2.

**TFPT is SUMOylated at K216**

Next, we turned to the other protein in this network, TFPT, and tested whether it could be SUMO-modified. Amino acid sequence analysis using GPS-SUMO software (39) revealed the presence of a consensus SUMOylation motif at K216 near the C-terminus of the protein. To test whether this lysine residue was important for SUMOylation, we created a lysine-to-arginine TFPT mutant, TFPTK216R. Then we conducted in vitro SUMOylation assays by expressing TFPT in rabbit reticulocyte lysate in the presence of [35S]-methionine, followed by incubation with or without the E1 and E2 SUMOylation enzymes with either SUMO1 or SUMO2 (Figure 5A). When wildtype TFPT was incubated with the SUMOylation enzymes and either SUMO1 or SUMO2, we observed a strong band representing a 20-kD molecular weight shift consistent with a SUMO-modified form of TFPT. However, when we incubated the in vitro translated TFPTK216R with the SUMOylation enzyme cocktails in the presence of either SUMO1 or SUMO2, no such shifted band was observed. This observation indicated that K216 is the major lysine residue for TFPT SUMO-modification by both SUMO isoforms SUMO1 and SUMO2.

To validate this result in vivo, we tested the ability of TFPT to be SUMOylated in transfected mammalian cells. We co-transfected V5-tagged TFPT with or without myc-tagged SUMO2 into HEK 293T cells, then immunoprecipitated V5-TFPT using an anti-V5 antibody, followed by immunoblotting using an anti-myc antibody to detect SUMOylated TFPT. In the absence of myc-SUMO2, no bands were observed in the anti-myc Western blot (Figure 5B). However, when we co-transfected SUMO2, we saw a strong band representing SUMO2-modified TFPT. Next, we tested the ability of TFPTK216R to be SUMOylated in mammalian cells. Although TFPTK216R was expressed (see input anti-V5 blot) at equal levels and immunoprecipitated (see IP:V5/IB:V5 blot) with equal efficiency, we failed to see any signals in the anti-myc blot, indicating that TFPTK216R was not SUMOylated. This shows that K216 is essential for SUMOylation of TFPT in mammalian cells.

**TFPT SUMOylation at K216 is enhanced by PIAS1 and PIAS3 in vivo**
Our group had shown significant enhancement of TFPT SUMOylation in the presence of the SUMO E3 ligases PIAS1 and PIAS3 using HuProt microarray based SUMOylation assays (18). In order to test whether PIAS1 and PIAS3 enhanced SUMOylation of TFPT in mammalian cells, we co-transfected TFPT constructs with either FLAG-tagged PIAS1 or untagged PIAS3 constructs (Figure 5B). In cells co-transfected with TFPT, SUMO2 and PIAS1 constructs, we observed a strong signal corresponding to mono-SUMOylated TFPT with several additional bands representing several higher molecular weight forms of SUMOylated TFPT, presumably due to either poly SUMOylation at multiple lysine residues or modification by polymeric SUMO2. However, in cells co-transfected with TFPT\textsuperscript{K216R}, SUMO2 and PIAS1 constructs, the anti-myc immunoblot showed only weak signal corresponding to the highest range of molecular weights. Consistent with our above observation, K216 is the major site in TFPT to be SUMOylated by PIAS1. Finally, we tested the ability of TFPT SUMOylation to be enhanced by the SUMO E3 ligase PIAS3. As with PIAS1, we saw a strong band representing mono-SUMOylated TFPT as well as a laddering band pattern representing several higher molecular weight forms of SUMOylated TFPT. This pattern was not observed when transfecting TFPT\textsuperscript{K216R}. Taken together, these results indicate that both SUMO E3 ligases PIAS1 and PIAS3 could effectively enhance TFPT’s SUMOylated on K216 in mammalian cells, and that PIAS1 and PIAS3 could further stimulate formation of higher molecular weight forms of SUMOylated TFPT.

**SUMOylation enhances interaction between TFPT and ION80E**

INO80E and TFPT are known subunits of the human INO80 chromatin-remodeling complex and interact with the N-terminal domain of the INO80 ATPase (31). The SUMOmodPPI network motif containing INO80E, TFPT and SUMO2 predicts that SUMOylation may enhance interaction between INO80E and TFPT. Considering that the INO80E C-terminal SIM is important for the ability of INO80E to interact with SUMO2, and that TFPT is modified by SUMO2, we hypothesized that the INO80E SIM may mediate the interaction between INO80E and SUMOylated TFPT. In order to test this hypothesis, we conducted an *in vitro* binding assay to allow us to measure binding between INO80E and SUMO-modified TFPT. We first generated recombinant radiolabeled TFPT using *in vitro* translation in the presence of \([^{35}S]\)-methionine, then incubated the reaction with or without SUMO1 or SUMO2 and the E1 and E2, to give us a mixture of unmodified and SUMO-modified TFPT (Figure 6A). We then used the resulting protein mixture to conduct binding assays with purified recombinant GST-INO80E bound to glutathione sepharose. Although the mixture of TFPT and SUMO2-modified TFPT forms contained a larger fraction of the unmodified form than the SUMOylated form, following binding to GST-INO80E, we only recovered the SUMO2-modified form of TFPT, suggesting that INO80E preferentially interacts with the SUMOylated form of TFPT.

When we conducted a binding reaction using the INO80E\textsuperscript{ΔC10} mutant lacking the C-terminal SIM, we saw a reduction in the recovery of SUMOylated TFPT. We then repeated this experiment in triplicate. We found a significant reduction in binding of the INO80E\textsuperscript{ΔC10} mutant to the SUMO2-modified form of TFPT, whereas binding to the SUMO1-modified form of TFPT was not significantly reduced (Figure 6B). Additionally,
the INO80EΔC10 mutant did not show any difference in its interaction with unmodified TFPT, relative to wildtype INO80E (data not shown). This suggests that the C-terminal SIM of INO80E is specifically required for interacting with the SUMO2-modified form of TFPT.

Next, we wanted to determine whether the INO80E C-terminal SIM was important for the interaction between INO80E and TFPT in mammalian cells. We conducted co-immunoprecipitation experiments in HEK293T cells to investigate the interaction between INO80E and TFPT. We transfected cells with myc-INO80E and V5-TFPT, then conducted immunoprecipitation using anti-myc, and Western blotting using anti-V5 (Figure 6C). This showed that wildtype INO80E could co-immunoprecipitate both wildtype TFPT and TFPTK216R (lanes 9 and 10). However, when we co-transfected the INO80EΔC10 mutant with TFPT, we observed a greatly reduced ability of mutant INO80E to co-immunoprecipitate both TFPT and TFPTK216R (lanes 11 and 12). We were unable to detect the SUMO-modified form of TFPT in the co-immunoprecipitated fraction, likely due to the low stoichiometry of this form in transfected cells.

Finally we co-transfected INO80E wildtype and INO80EΔC10 mutant with TFPT and SUMO2. We observed enhanced interaction between INO80E and TFPT under conditions of SUMO2 overexpression, and this was observed for both the wildtype INO80E and the INO80EΔC10 mutant (lanes 13 and 14). However, by calculating the ratio of co-immunoprecipitated TFPT relative to the amount of INO80E, we observed a greater than three-fold increase in the amount of TFPT bound to wildtype INO80E compared to the amount of TFPT bound to INO80EΔC10. These results suggest that SUMO2 promotes the INO80E-TFPT interaction, and that the INO80E C-terminal SIM is important for this interaction.

**DISCUSSION**

Harnessing the power of the human proteome microarray, we performed the most comprehensive and unbiased search to date for novel SUMO-binding partners. Furthermore, by integrating this SUMO-binding data with SUMO substrate data, as well as protein-protein interactions, we have generated an extensive list of SUMO-centered networks that suggest new functions of SUMO, particularly in mediating protein-protein interactions. This fits with previously proposed models of SUMO functioning as molecular glue that stabilizes protein-protein interaction (24, 40). We have found that a combination of SUMOylation and SUMO-binding mediate the interaction between INO80 complex subunits. SUMOylation of the INO80 complex subunit TFPT primes the protein for interaction with INO80E, via its SUMO interaction motif.

However, there are some important caveats to this model. For example, we observed that INO80E has the ability to interact with unmodified TFPT and that this interaction is enhanced when SUMOylated TFPT is absent from the reaction mixture. Under *in vitro* conditions in which SUMOylated TFPT is present, there is only limited binding of INO80E to unmodified TFPT. This suggests to us that both SUMO-modified and unmodified forms of TFPT may compete for binding to a single site on INO80E, and that TFPT interaction with this site may be enhanced by TFPT SUMOylation. In
mammalian cells, we found that the interaction between INO80E and TFPT may be dependent on the INO80E SIM, regardless of whether the SUMOylation site on TFPT that we identified has been mutated. This could indicate that the INO80E SIM may be required for interaction with unmodified TFPT. Another possibility is that TFPT could be SUMOylated on multiple additional sites that these minor SUMOylation sites may also contribute to the interaction with INO80E. Alternatively, the INO80E SIM may interact with other endogenous SUMO-modified proteins, which may in turn mediate the interaction between INO80E and TFPT.

Although the roles of the INO80E and TFPT subunits in INO80 complex function are unknown, and thus the significance of their interaction is unclear, these proteins have been proposed to serve a regulatory function that is likely specific to metazoans, as there are no known orthologs of these subunits in yeast (32). We propose that the SUMO-SIM interaction may be an important regulator of the INO80 complex.

Furthermore, we confirmed the results of a human proteome microarray-based screen to identify SUMO E3 ligase-dependent SUMOylation substrates that showed that SUMOylation of TFPT is enhanced by the SUMO E3 ligases PIAS1 and PIAS3 (18). Previously, PIAS1 and PIAS3 were identified as upstream regulators of chromatin remodeling through their role in stimulating SUMOylation of the protein MBD1, thus affecting its ability to interact with the histone methyltransferase SETDB1 (41). Thus, regulation of chromatin-associated protein-protein interactions may be a general role of the PIAS1 and PIAS3 SUMO E3 ligases.

There is some evidence to suggest TFPT may function as a transcription factor. TFPT was reported to contain a b-ZIP domain (31), and it was found to possess sequence-specific DNA binding activity in a screen to identify DNA binding proteins using a transcription factor protein microarray (8). The transcription factor YY1, a known SUMO target, was shown to recruit the INO80 complex to specific gene promoters (42, 43). SUMOylated TFPT may play a similar role, by binding to both the genome and to the INO80 complex via the SUMO-binding subunit INO80E, in order to recruit the complex to specific genomic locations. This in turn should regulate chromatin conformation and gene transcription. This is a particularly attractive hypothesis considering the well-established role of SUMOylation in regulating transcription factors by directing the assembly of specific multiprotein complexes.

A recent study found that knockdown of the INO80 ATPase, as well as the other INO80 complex subunits INO80E and TFPT, resulted in embryonic stem cell differentiation (44). Additionally, knockdown of INO80E in mouse embryonic stem cells resulted in significant gene expression changes in >1200 genes as measured by microarray analysis (44). Interestingly, the INO80 complex was found to regulate the gene Sox2 and occupy multiple locations in the Sox2 gene in mouse embryonic stem cells. Interestingly, a YY1 binding site identified by the ENCODE project also overlaps with a TFPT DNA sequence motif identified previously (8) in the orthologous region of the human genome. This points to a potential role of both TFPT and YY1 working
redundantly to recruit the INO80 complex to regulatory elements of Sox2 to sustain embryonic stem cell pluripotency.

This work extends the list of known SUMO-binding proteins tenfold. Considering that many previously identified SUMO-modified proteins are localized to the nucleus, including many transcription factors, this enrichment of cytoplasmic proteins suggests that the importance of SUMOylation in the cytoplasm may be underappreciated. There are alternative explanations for the observed enrichment of cytoplasmic proteins among the SUMO-interacting proteins identified by our approach. It is possible that SUMO’s interaction with nuclear proteins could be dependent on the presence of other nuclear proteins or DNA that is not present in the reaction mixture incubated on the protein microarray, thus hindering our ability to identify nuclear SUMO-interacting proteins. Additionally, it is possible that in some cases, other proteins or biomolecules were co-purified with proteins that were spotted in the array. Although our analysis of randomly chosen purified proteins indicates that these proteins are generally of high quality and high purity, and we estimate that 98% of proteins have the correct band size (10), it is impractical to analyze every protein that we have purified and spotted on our protein arrays. Unidentified contaminants, including other interacting proteins, could interact strongly with our SUMO probes, thus giving us a false positive result. We believe that every experimental approach to identify SUMO-interacting proteins has their own inherent biases. Previous analysis of protein quantities spotted on our arrays versus protein subcellular localization and function showed that there are higher amounts of cytoplasmic proteins on our protein arrays compared to proteins localized to other parts of the cell. Additionally, transcription factors, which have frequently been associated with SUMOylation in the literature, are expressed at lower levels in our yeast expression system, which may contribute to the low number of transcription factors identified in our assay (10).

The in vivo role of the SUMO-binding activity of these proteins must be determined through systematic identification of the relevant SUMO interaction motifs, mutagenesis, and further study of the downstream consequences. We believe that our network motif analysis approach further contributes an important context in which to look at the role of the SUMO-binding activity of these proteins.

METHODS

SUMO binding assay on human proteome microarrays
SUMO binding experiments were conducted with 4 probes representing the SUMO1, SUMO2 and the SUMO1 and SUMO2 trimers, all in triplicate. Protein chips were incubated overnight in blocking buffer (20 mM HEPES-KOH pH 7.3, 100 mM potassium acetate, 2 mM magnesium acetate, 1 mM EGTA, 0.05% Tween-20, 2% BSA) at 4° C. Chips were then incubated with SUMO1 or SUMO2 monomer or trimer in assay buffer (20 mM HEPES-KOH pH 7.3, 100 mM potassium acetate, 2 mM magnesium acetate, 1 mM EGTA, 0.05% Tween-20) for 1 hr at room temperature. Following incubation with SUMO, chips were washed 3X with assay buffer and 3X with PBS, followed by incubation with DyLight 549-anti-SUMO1 (21C7) or DyLight 649-anti-SUMO2 (8A2).
Negative control experiments were done in parallel using antibodies only without SUMO. Chips were then washed 3X with 1X TBS with 0.05% Tween-20, 1X with water, then spun to dry and scanned.

**Immunoprecipitation and Immunoblotting**

HEK 293T cells were transfected with Fugene 6 (Promega) and HCT 116 cells were transfected with Fugene HD (Promega). Both cell types were plated in 6-well plates, and harvested 24-48 hours following transfection. In order to detect protein SUMOylation, cells were washed with 1X PBS, then lysed in SUMO IP lysis buffer (50 mM Tris-HCl pH 7.4, 150 mM NaCl, 2 mM EDTA, 1% Triton X-100, 0.5% sodium deoxycholate, 20 mM N-ethyl-maleimide, 1% SDS, and Roche protease inhibitor cocktail) for 10 minutes at 4° C. Then SUMO IP binding buffer ((50 mM Tris-HCl pH 7.4, 150 mM NaCl, 2 mM EDTA, 1% Triton X-100, 0.5% sodium deoxycholate) was then added to dilute the lysate ten-fold, and lysates were transferred to cold microcentrifuge tubes. Cells were then sonicated on ice and centrifuged at 15,000 g at 4° C for 10 minutes. Soluble cell lysates were incubated with Dynabeads (Life Technologies) pre-bound to either anti-Flag (F1804, Sigma), anti-myc (R95025, Life Technologies), or anti-V5 (R9605, Life Technologies) antibodies for 2 hours at 4° C and washed 3 times with SUMO IP wash buffer (50 mM Tris-HCl pH 7.4, 150 mM NaCl, 2mM EDTA, 1% Triton X-100, 0.5% sodium deoxycholate, and 0.1% SDS). Bound proteins were eluted with 1X LDS-sample buffer (NP0008, Life Technologies) with 5% beta-mercaptoethanol. In order to look at protein interactions between INO80E and TFPT by co-immunoprecipitation, cells were transfected and harvested as above, but lysed in INO80 lysis buffer (40 mM HEPES-NaOH pH 7.9, 0.3 M NaCl, 0.2% Triton X-100, and 10% glycerol), followed by centrifugation. Lysate supernatants were then bound to Dynabeads linked to anti-FLAG or anti-V5 for 2 hours at 4° C and then washed 3X with INO80 wash buffer (40 mM HEPES-NaOH 7.9, 0.25 M NaCl, 0.2% Triton X-100, and 10% glycerol). In both immunoprecipitation to detect SUMOylated proteins and co-immunoprecipitation experiments to detect interacting proteins, samples were resolved on 4-12% NuPage Bis-Tris gels (Life Technologies) in MES buffer and subject to immunoblotting using HRP-anti-V5 and HRP-anti-myc antibodies (Life Technologies).

**Recombinant proteins**

GST, GST-tagged SUMO1, SUMO2, and SUMO1 and SUMO2 trimers were expressed in bacteria and GST-IN080E was expressed in yeast. All GST-tagged proteins were purified by affinity chromatography on glutathione-Sepharose 4B beads (GE Healthcare) according to manufacturer’s instructions.

**In vitro binding assays**

Recombinant GST, GST-tagged SUMO1, SUMO2, or SUMO2 trimer were diluted into 100 µl of 1X PBS, 0.05% Tween 20 and incubated with either glutathione-sepharose or alternatively glutathione-coated 96-well plates (Pierce Biotechnology). Following overnight incubation at 4° C, wells were blocked for 1 hr at room temperature with 2% bovine serum albumin in assay buffer (20 mM HEPES-KOH pH 7.3, 100 mM potassium acetate, 2 mM magnesium acetate, 1 mM EGTA, 0.05% Tween-20). IN080E and TFPT were produced by in vitro transcription and translation in rabbit reticulocyte
lysate in the presence of [³⁵S] methionine according to manufacturer’s instructions (Promega). *In vitro* translated proteins (10 µl) were diluted into 100 µl of assay buffer, when appropriate, incubated with SUMO1 or SUMO2 and the SUMO E1 and E2 enzymes, then incubated with the immobilized GST-tagged proteins for 1 hr at room temperature. Unbound proteins were removed by washing, and bound proteins were eluted with SDS sample buffer and resolved by SDS-PAGE and autoradiography. Equal loading of immobilized proteins was verified by staining with SimplyBlue SafeStain (Life Technologies). Quantitative binding was determined by scanning densitometry measurements of radiolabeled protein relative to the amount of binding partner protein as indicated by Coomassie stain.

**In vitro SUMOylation assays**
TFPT was produced by *in vitro* transcription and translation in rabbit reticulocyte lysate in the presence of [³⁵S] methionine according to manufacturer’s instructions (Promega). Radiolabeled protein was then incubated either with or without SUMO1 or SUMO2 and the SUMO E1 and E2 enzymes for 1 hour at 37° C to yield a mixture of SUMOylated and unmodified TFPT for subsequent binding experiments.

**Protein Microarray Data Analysis**
To identify positive hits for each chip, uninformative spots, such as those in damaged or high-background regions of the chip, were first identified as having a coefficient of variation (CV) value >1.5, and excluded from analysis. A background correction was then applied to the remaining spots, defined as the signal intensity (SI) value of each spot as the odds ratio of the foreground median divided by the background median. If a spot has a weak signal, foreground median and background median signal intensity will be similar, and its SI will thus be ~1. To normalize the signal, we assumed that true hits are relatively rare and almost evenly dispersed in each block, thus we force each block on a chip to have a median SI of 1. To be considered a positive hit, the duplicate spots of each protein need to have both have signal intensities (foreground/background ratio) 5 SD above the mean value. Additionally, positive hits need to be detected in 2 out of 3 triplicate experiments to be considered positive. Positive hits that were detected in negative control chips were also excluded from analysis.

**Network Motif Analysis**
Protein microarray data from both SUMO-binding experiments and SUMOylation experiments (18) was integrated with protein-protein interaction data from publicly available databases, including BioGRID, DIP, MIPS, IntAct and HPRD and network motifs were identified and analyzed as previously described (38). The Z-value of a motif is calculated as the difference of its observed occurrence in a real network and its averaged occurrence in several hundred random networks, normalized with the standard deviation.

**ACKNOWLEDGEMENTS**
The work was supported by NIH grant EY017015 (to S.B.), National Institutes of Health Common Fund Grant U54RR020839 to H.Z., and a predoctoral award from the American Heart Association 10PRE3040000 to E.C. We thank W. Yap for comments.
on the manuscript.
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18. Uzoma I., Hu J., Cox E. et al. (2017) Global Identification of SUMO Substrates Reveals that Crosstalk between SUMOylation and Tyrosine Phosphorylation Promotes Cell Migration. Submitted,
FIGURE LEGENDS

Fig. 1  Design of protein microarray-based identification of human SUMO binding proteins.
A, (Top) SUMO1 and SUMO2 protein surfaces generated using MacPyMOL with previously reported crystal structures (SUMO1 PDB ID 2PE6 and SUMO2 PDB ID 4BK6). The surfaces corresponding to the first alpha-helix and second beta-strand for both SUMO1 and SUMO2 are shown in color (nitrogens are shown in red, oxygens in blue and carbons in green). This surface has been reported to be important for interactions with other proteins. (Bottom) Cartoon showing 4 probes that were used in our experiments. SUMO1 and SUMO2 monomer probes were based on the proteolytically processed (mature) forms of SUMO1 and SUMO2. The SUMO1 and SUMO2 trimer probes, described previously (Zhu 2008), were expressed as trimeric fusion proteins. B, Schematic of the SUMO-binding protocol used with the human proteome microarray. C, Representative microarray images showing human proteome microarray visualized with an antibody to GST, showing all proteins (left, in green), and an antibody to SUMO2 following a SUMO-binding experiment with the SUMO2 trimer (right, in red).

Fig. 2  Identification of SUMO binding proteins using human proteome microarrays.
A, Venn diagram showing the number of SUMO-binding experiment hits for each probe tested. B, Gene ontology analysis on proteins identified in our screen for SUMO-binding proteins. Gene ontology categories that are significantly enriched among proteins that bound to one of our 4 probes are shown as a colored box, color-coded by probe.

Fig. 3  Network analysis of SUMO binding and SUMO-conjugated human proteins.
Schematic of network motif analysis approach showing the 3 data sets that were used and all the enriched and depleted network motifs. The “SUMOmod PPI” network motif that was the focus of subsequent experiments is boxed in orange.

Fig. 4  INO80E binds to SUMO1 and SUMO2 through a C-terminal SIM domain.
A, Cartoon showing INO80E, the location of the SUMO interaction motif (SIM) at the C-terminus of the protein, and the two SIM-deficient INO80E mutants generated for this study. B, Results from in vitro binding assay. INO80E, the INO80E SIM mutant and the INO80EΔC10 were generated by in vitro translation in the presence of [35S]-methionine and incubated with glutathione sepharose-bound GST-SUMO1 and GST-SUMO2. C, INO80E mtSIM shows reduced binding to GST-SUMO2, but not GST-SUMO1.

Fig. 5  PIAS1 and PIAS3 induce modification of TFPT by SUMO2.  A, Results from in vitro SUMOylation assay for TFPT. Wildtype TFPT and the K216R mutant were translated in vitro in the presence of [35S]-methionine and incubated either with or without the E1 and E2 SUMOylation enzymes, and with either SUMO1 or SUMO2. B, Results from the in vivo SUMOylation assay for TFPT. HEK 293T cells were co-transfected with TFPT and the K216R mutants both in the presence or absence of
SUMO2, and in the presence or absence of either PIAS1 or PIAS3. We immunoprecipitated V5-tagged TFPT and immunoblotted against myc-tagged SUMO2 to detect SUMOylated TFPT.

Fig. 6  NO80E selectively binds to SUMO-conjugated TFPT. A, Results from *in vitro* binding assay of [*35S*]-methionine-labeled TFPT to wildtype and SIM-deficient INO80E in the presence and absence of E1/E2 enzymes and SUMO. B, Quantitative analysis of these results by scanning densitometry. C, Results from *in vivo* binding assay of wildtype and mutant TFPT and INO80E the presence and absence of SUMO2. Myc-tagged INO80E was immunoprecipitated, then immunoblotted with anti-V5 to detect co-immunoprecipitated TFPT.
SUMO1

SUMO1 trimer

SUMO2

SUMO2 trimer

B

1) Block chip

2) Incubate with SUMO1 or SUMO2

3) Wash

4) Incubate with Dylight labeled-anti-SUMO1/SUMO2

5) Wash

6) Dry and scan

C

α-GST

TRUB1

INO80E

MAGEB1

SUMO1

SUMO2

α-SUMO2
Cox, et al. Figure 2
Cox, et al. Figure 3

SUMOylation data (Uzoma et al. (18))
SUMO-binding data (This study)
Protein-protein interaction data (databases)

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SUMO-modulated Protein-Protein Interaction “SUMOMod PPI”

SUMOylation
SUMO-binding
Protein-protein interaction

P1 = protein #1
P2 = protein #2
S = SUMO1 or SUMO2
n = number of examples of this motif type
Z = enrichment score
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Cox et al. Figure 4