The non-detergent sulfobetaine-201 acts as a pharmacological chaperone to promote folding and crystallization of the type II TGF- β receptor extracellular domain.

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Abstract

The roles of the extracellular domain of type II TGF- β receptor (TBRII-ECD) in physiological processes ranging from development to cancer to wound healing render it an attractive target for exploration with chemical tools. For such applications, large amounts of active soluble protein are needed, but the yields of TBRII-ECD we obtained with current folding protocols were variable. To expedite the identification of alternative folding conditions, we developed an on-plate screen. This assay indicated that effective folding additives included the non-detergent sulfobetaine-201 (NDSB-201). Although NDSB-201 can facilitate protein folding, the mode by which it does so is poorly understood. We postulated that specific interactions between NDSB-201 and TBRII-ECD might be responsible. Analysis by X-ray crystallography indicates that the TBRII-ECD possesses a binding pocket for NDSB-201. The pyridinium group of the additive stacks with a phenylalanine side chain in the binding site. The ability of NDSB-201 to occupy a pocket on the protein provides a molecular mechanism for the additive's ability to minimize TBRII-ECD aggregation and stabilize the folded state. NDSB-201 also accelerates TBRII-ECD crystallization, suggesting it may serve as a useful crystallization additive for proteins refolded with it. Our results also suggest there is a site on TBRII-ECD that could be targeted by small-molecule modulators.

Keywords:

Non-detergent sulfobetaine 201, transforming growth factor β , type II transforming growth factor β receptor, protein folding, pharmacological chaperone

Abbreviations:

TBRII-ECD, extracellular domain of type II TGF- β receptor; TBRII-ECD-PR, protease resistant variant of TBRII-ECD; NDSB, non-detergent sulfobetaine; TGF- β , Transforming growth factor β ; TBRII, type II TGF- β receptor; TBRI, type I TGF- β receptor; Trx, thioredoxin; GSH, reduced glutathione; GSSG, oxidized glutathione

Graphical abstract:

See Graphical_abstract.jpg

Highlights:

- An on-plate assay for TBRII-ECD folding identifies NDSB-201 as an effective additive.
- NDSB-201 can be used in large-scale folding experiments to reproducibly improve TBRII-ECD yield.
- X-ray crystallography reveals a specific binding site for NDSB-201 on TBRII-ECD.
- NDSB-201 can function as a pharmacological chaperone.

Introduction

Transforming growth factor beta (TGF- β) signaling controls cell proliferation, differentiation, and tissue remodeling [1]. There are three main TGF- β members—TGF- β 1, TGF- β 2, and TGF- β 3. TGF- β 1 and TGF- β 3—each of which are 25 kDa homodimeric proteins that form a multimeric assembly with the type II TGF- β receptors (TBR2) [2]. The resulting complexes bind to type I receptors (TBR1) and promote their phosphorylation. This activated TGF- β /TBR1/TBR2 kinase complex then mediates SMAD protein phosphorylation, which leads to the translocation of phospho-SMAD proteins into the nucleus where they regulate the expression of numerous genes [3]. The outcome of TGF- β responses depends upon the concentration of TGF- β , cell age, state of differentiation, presence of other growth factors, and the cellular environment [4]. Therefore, spatial and temporal control of TGF- β signaling is crucial for directing cell differentiation or to aid wound healing. Alternatively, dysregulation of TGF- β signaling has been implicated in several diseases, including proliferation of cancer cells and vascularization of tumors [5].

Because of the myriad roles of TGF- β signaling, chemical tools that modulate TGF- β signaling are valuable. An inhibitor of the kinase activity of the TBR1, SB-431542, blocks migration and vascular endothelial growth factor secretion, processes which are crucial for cancer growth and metastasis [6]. Exogenous application of TGF- β 3 to wounds reduces scarring [7]. These diverse signaling outcomes led us to seek synthetic ligands for the extracellular domains of TBR that are orthogonal to TGF- β . Our goal was to immobilize such ligands to control TGF- β signaling, such that the pathway could be activated locally, as would be desirable for promoting wound healing.

We identified two peptide ligands (LTGKNFPMFHRN and MHRMPSFLPTTL) that can bind both the extracellular domains (ECDs) of TBRI and TBRII to pre-organize the receptors on the cell surface [8]. This preorganization increases the sensitivity of peptide-bound cells to minute concentrations (pM) of soluble TGF- β and allows cell differentiation in a spatially-controlled manner [9] [10]. To exploit the potential of these peptides and design next-generation probes of TBR function, a molecular level knowledge of their binding mode is needed. To pursue the required structural and biophysical studies, large quantities of TBRs are required. In our hands, the previously reported expression and purification protocols resulted in variable yields of functional TBRII-ECD [11-13]. The variability resulted from the folding step; therefore, we sought to develop an on-plate screen to optimize folding.

We employed thioredoxin-fused TBRII-ECD (Trx-TBRII-ECD) in a folding screen that surveyed a variety of conditions. Non-detergent sulfobetaine 201 (NDSB-201) was found to be an effective additive, with a yield of active TBRII-ECD up to three-fold higher than previously observed. NDSB-201 also facilitates the folding of urea-denatured protease-resistant untagged TBRII-ECD (TBRII-ECD-PR) in solution. To understand the molecular basis for the ability of NDSB-201 to promote folding, we used X-ray crystallography. The data reveal that TBRII has an NDSB-201 binding site. These findings suggest that NDSB-201 acts as a pharmacological chaperone [14-16] not only by preventing aggregation of folding intermediates but also by binding and stabilizing the folded state [14-18] .

Material and Methods

Expression of Thioredoxin-Fused TBR II (20-136) (Trx-TBR II-ECD)

The sequence encoding the extracellular domain of TBR II (20-136) was amplified from a TBR II construct (encoding amino acids 15-136) [12] and cloned into the KpnI and NcoI sites of pET32b. Upstream of the multiple cloning sites, is a thioredoxin coding sequence, so the resulting construct encodes a fusion protein. The plasmid was transformed into the Tuner (DE3) strain (Novagen), a BL21 (DE3) derivative with a lactose permease deletion allowing uniform induction by isopropyl β -thiogalactopyranose (IPTG). Cells were grown in Terrific Broth (Research Products International Corp.) at 37 °C to OD₆₀₀ of 0.6. Gene expression was induced by addition of 0.1 mM IPTG and the cells were incubated for 14 hours at 15 °C. Bacteria were harvested by centrifugation and lysed (25 g wet pellet) via sonication in 50 mL of 50 mM sodium phosphate (pH 7.4), 300 mM NaCl, 0.1 mg/mL lysozyme, 2.5 mM benzamidine, 1 mM phenylmethylsulfonyl fluoride (PMSF), and 1 tablet/100 mL SIGMAFAST™ Protease Inhibitor Cocktail Tablet (EDTA-Free) (Sigma-Aldrich). Lysate was clarified by centrifugation at 100,000 x g for 1 hour and filtration through a 0.45 μ m syringe filter.

Folding Assay

An on-plate screen for protein folding was implemented. Clarified lysate (200 μ L) containing Trx-TBR II-ECD was added to each well of a Ni-NTA HisSorb plate (Qiagen). After 2 hours at room temperature, the plate was washed 3 times for 5 minutes each, with 200 μ L of 50 mM sodium phosphate (pH 7.4), 300 mM NaCl, and 25 mM imidazole. Buffered solutions were screened for their ability to support folding (200 μ L) by adding them to each well and holding the samples at 4 °C for 40 hours. The plate was

then washed with phosphate-buffered saline (PBS) with 0.05% Tween 20 (PBST). The relative amount of correctly folded TBRII-ECD was measured directly on the plate using a previously described protocol that monitors TGF- β 1 interaction with the receptor [13]. Briefly, the plate was blocked with 200 μ L of 5% BSA in PBST at 37 °C for 1.5 hours. TGF- β 1 (5 ng/mL, 100 μ L, Cell Signaling Technology) was added. After 2 hours at room temperature (rt), a solution of 100 μ L (100 ng/mL) of biotinylated anti-TGF- β 1 (BAF240, R&D Systems) was added, and the mixture was allowed to stand at rt for 1 hour. After washing, 100 μ L (200 ng/mL) of streptavidin-HRP (Jackson ImmunoResearch) was added. After 30 minutes, functional complex formation was monitored via HRP activity by addition of the colorimetric substrate 1-Step Ultra TMB (100 μ L, Pierce). The reaction was quenched with addition of 100 μ L of 2 M sulfuric acid. Absorbance at 450 nm was measured to quantify the amount of correctly folded TBRII-ECD.

Production of TBRII (26-136) Q26A K97T (TBRII-ECD-PR)

The gene encoding untagged TBRII (26-136) containing two substitutions, Q26A and K97T (derived from Trx-TBRII-ECD by site-directed mutagenesis) that should confer protease resistance, was ligated into a pET24a plasmid between the NdeI and BamHI sites. The resulting construct was transformed into Tuner (DE3) cells. The bacteria were grown in Terrific Broth supplemented with 50 μ g/mL kanamycin and 0.2% glucose until OD₆₀₀ reached 0.6. Gene expression was induced with 1 mM IPTG, and the mixture was cultured for 6-8 hours. Cells were collected by centrifugation and kept frozen at -80 °C until use.

Folding and purification of TBR11 (26-136) Q26A K97T (T β RI-ECD-PR)

Production of TBR11-ECD-PR afforded inclusion bodies, which were isolated using a procedure similar to a protocol for production of TGF- β 3 described previously [19]. Briefly, bacterial pellets were resuspended in 100 mM Tris (pH 8.0), 10 mM EDTA, 0.1 mg/mL lysozyme, 2.5 mM benzamidine, and 1 mM PMSF. The mixture was sonicated, and the insoluble material was isolated by centrifugation. The pellet was resuspended in 100 mM Tris (pH 8.0), 10 mM EDTA, and 1% Triton X-100 and sonicated again. After centrifugation, the pellet that contained the inclusion bodies was resuspended by sonication in 100 mM Tris (pH 8.0), 10 mM EDTA, and 1 M NaCl. The resulting suspension of washed inclusion bodies was pelleted. This pellet was solubilized in 20 mM Tris (pH 8.0), 20 mM Bis-Tris, 100 mM DTT and 8 M urea at room temperature using a homogenizer. Insoluble materials were removed by centrifugation and the pH of the supernatant was adjusted to 6.0. After passage through a 0.22 μ m filter, the solution was loaded onto an anion exchange column (HiTrap Q FF, GE Healthcare) equilibrated with 20 mM Bis-Tris pH 6.0 and 8 M urea. TBR11-ECD-PR was eluted using a linear gradient of 0-300 mM NaCl. Fractions containing the desired protein were identified after analysis by SDS-PAGE. The protein concentration of the eluents was estimated by UV absorbance at 280 nm using an extinction coefficient of 8480 M⁻¹ cm⁻¹ [20]. The protein was dispensed as 50 mg aliquots and kept at -80 °C until use.

To initiate folding, the protein solution was added dropwise to a chilled solution of 75 mM Tris (pH 8.0), 1 M NDSB-201, 2 mM reduced glutathione (GSH), and 0.5 mM oxidized glutathione (GSSG). The protein concentration was kept at approximately 0.1 mg/mL. After stirring for 2 days at 4 °C, the solution was concentrated using an Amicon stirred cell (3 kDa cut-off) under nitrogen pressure, and the concentrate was dialyzed

extensively against 20 mM Bis-Tris (pH 6.0). Refolded protein was captured with an anion exchange column (HiTrap Q HP, GE Healthcare) equilibrated with 20 mM Bis-Tris (pH 6.0). Protein was eluted with a linear gradient of 0-300 mM NaCl (Figure 1). Fractions containing properly folded TBR II-ECD-PR were identified using the ELISA protocol similar to that employed in the aforementioned folding assay. For this ELISA, TBR II-ECD-PR was absorbed directly onto a Nunc MaxiSorp ELISA plate (Thermo Scientific). We found that this functional binding assay was effective in this format as well. Active fractions (i.e., those with functional protein) were pooled and dialyzed against 50 mM sodium phosphate (pH 7.0). Ammonium sulfate was added to 1 M, and the resulting solution was subjected to hydrophobic interaction chromatography (HiTrap Butyl HP, GE Healthcare) to capture minor impurities. Active TBR II-ECD-PR did not bind the column under these conditions. The TBR II-ECD-PR-containing solution (flow through) was purified further on a Superdex 75 16/60 column using 20 mM Tris (pH 7.4), 300 mM NaCl as a running buffer.

Crystallization and structural analysis of TBR II (26-136) Q26A K97T (TBR II-ECD-PR)

Purified TBR II-ECD-PR was dialyzed extensively against water and concentrated to 20 mg/mL. The protein concentration of the final solution was determined by UV absorbance at 280 nm using an extinction coefficient of 9230 M⁻¹cm⁻¹ [20]. The resulting sample was subjected to previously reported crystallization conditions (100 mM sodium citrate pH 5.0, 30% PEG 2000) supplemented with 50 mM NDSB-201 [21]. Crystals appeared and grew to full size in 2-3 days. This time scale is accelerated compared with that reported previously (and consistent with our experience) of 7-14 days for conditions in which NDSB-201 is absent [21]. The crystals were cryoprotected

using paratone, then vitrified and stored in liquid nitrogen. Single crystal diffraction data were collected at beamline 21-ID-D (LS-CAT) at the Advanced Photon Source, Argonne National Laboratory. Data reduction and scaling were performed with HKL2000 [22] (Table 1). Molecular replacement was performed with PHENIX AutoMR [23, 24] using a previously reported TBRII structure (PDB ID 1M9Z) [25] as a search model. NDSB-201 coordinates and restraints were generated using eLBOW [26]. Coot [27] and PHENIX.refine [28], respectively, were used for model fitting and refinement (Table 1). The model was validated using MolProbity [29]. Figures depicting the protein structure were generated with PyMOL [30].

Results

Folding screen for the extracellular domain of TBRII

Because TBRII engagement by TGF- β initiates signaling, the receptor is an attractive target for chemical tools and therapeutics that modulate TGF- β activity. An efficient protocol to produce active TBRII-ECD is therefore an enabling technology. Our attempts to follow reported protocols for folding TBRII-ECD from inclusion bodies [11, 12, 31] afforded highly variable yields of active protein; often less than 1 mg of protein was recovered from 50 mg of solubilized inclusion bodies. Although a thioredoxin fusion enhanced the solubility of chicken Trx-TBRII-ECD [32, 33], production of the corresponding human Trx-TBRII-ECD fusion resulted in soluble but nonfunctional protein that failed to bind TGF- β 1. A protocol for folding Trx-TBRII-ECD on Ni-NTA agarose was previously reported [13]; however, it employs a high concentration of arginine at pH 8.0, which interferes with protein binding to Ni-NTA agarose [34]. Still, we were inspired by this heterogenous folding strategy [13] to examine on-plate folding

as a means of rapidly identifying improved conditions. For this assay, soluble Trx-TBR II-ECD was immobilized onto a Ni-NTA-coated 96-well plate (Figure 2), and different conditions were tested. Each well was treated with an amalgam of the background buffer, one additive, and one redox couple; this mixture constituted a “folding buffer”. In total, 81 conditions were screened. Because TBR II-ECD contains multiple disulfide bonds, solutions that varied in redox potential were tested. To facilitate disulfide exchange, the pH was held constant at 8.0, which is similar to the pKa of cysteine thiol. The additives examined were selected on the basis of literature precedents [35-38] (Table 2). To quantify the efficiency of folding, a functional assay for receptor binding was employed by monitoring the amount of TGF- β 1 bound as detected by a TGF- β 1-specific antibody (Figure 3).

A single experiment folding screen indicated that urea, NDSB-201, or NDSB-256 each can serve as an effective additive (Figure 3). Compared to the efficiency of folding obtained using PBS, the use of a redox buffer at pH 8.0 improves the yield of folded Trx-TBR II-ECD. We therefore held the redox couple constant (2 mM GSH + 0.5 mM GSSG) and repeated the refolding experiment in triplicate (Figure 4). Using the level of folding obtained in redox buffer alone as a baseline, the influence of additives was compared. When either urea, NDSB-201, or NDSB-256 was added, a high level of active Trx-TBR II-ECD was obtained.

Although urea was effective, we focused on the NDSBs. NDSBs can promote protein folding but their mechanism of action is poorly understood. We reasoned that their effects on TBR II-ECD might be leveraged to provide insight into the mechanism of NDSB-aided folding. Moreover, we postulated that these additives could bind TBR II-ECD. This hypothesis arose because all NDSBs are zwitterionic but not all NDSBs

improved folding in our assay, and these two observations suggest that, the effects of the NDSBs on folding are unlikely to arise solely from bulk solution effects. As NDSB-201 was the most cost-effective of the useful additives (~\$0.4/g vs ~\$13/g), it was used in larger scale solution phase folding experiments of the untagged TBRII-ECD-PR construct that had been used previously for structure determination [25]. Our optimized conditions consisted of 75 mM Tris (pH 8.0), 1 M NDSB-201, 2 mM reduced glutathione (GSH), and 0.5 mM oxidized glutathione (GSSG) at 4 °C for 40 hours. This protocol yields 8-13 mg of purified TBRII-ECD-PR from 50 mg of urea-solubilized protein. This protocol was successful for either Trx-TBRII-ECD or TBRII-ECD-PR, suggesting that the conditions identified in the on-plate folding assay were also effective for folding in solution. Because the protease-resistant variants of TBRII-ECD-PR undergo folding, we conclude these mutations do not impact folding significantly.

Crystallographic analysis of TBRII-NDSB-201 complex

We used X-ray crystallography to examine whether NDSB-201 binds to the TBRII fusion protein. Purified TBRII-ECD-PR could be crystallized using previously reported conditions [21]. When NDSB-201 was added, protein crystals grew and reached full size in 2-3 days as opposed to 1-2 weeks. Analysis by X-ray crystallography revealed a molecule of NDSB-201 bound to TBRII-ECD-PR, as the electron density of the pyridinium ring and the sulfonate group were apparent in the difference density map (Figure 5). The NDSB-201 interaction site is a hydrophobic pocket formed by W65, F111, F126, and a disulfide bridge involving cysteine residues at positions 38 and 44. The cationic pyridinium ring of NDSB-201 stacks onto the phenyl ring π system of F126 in a staggered conformation. The pyridinium ring also interacts with W65 in an edge-on

manner. There are additional contacts to several adjacent hydrophilic residues (D39, N40, K67, E102, and T109); however, functional group atoms of these side chains are more than 4 Å away from the sulfonate group of NDSB-201 and may not contribute substantially to the NDSB-201 binding. The total buried surface area of the complex is 375 Å² (PISA) [39]. These results indicate that NDSB-201 occupies a binding pocket and engages in favorable interactions with TBR II-ECD.

Discussion and Conclusions

An on-plate assay was developed to rapidly identify conditions that promote folding of the Trx-TBR II-ECD. The most effective conditions from the plate assay were also useful in folding untagged TBR II (TBR II-ECD-PR) in solution, indicating that the results from the higher throughput assay translated well to solution refolding. The assay provided valuable information about chemical sensitivity of TBR II-ECD during the folding process. Specifically, a low concentration of urea is highly effective at folding TBR II-ECD, but the efficiency of that process rapidly dropped with increasing urea concentration (Figure 4A). These results highlight the importance of controlling urea concentration when applying the previously published protocol [12]. Because our screen covers significant chemical space of known refolding reagents, it could readily be applied to other monomeric receptors or enzymes for which an appropriate detection system for functionally folded protein is available. Small amounts of candidate folding buffers can readily be generated by mixing stock solutions in 96-well blocks, followed by transfer directly to the folding plate; accordingly, this assay and screening format are readily adaptable for a high-throughput platform using liquid handling robots.

The additives we identified, urea and members of the NDSB series, are aggregation suppressors [40, 41], suggesting that a major bottleneck of TBRII-ECD folding is aggregation. Still, NDSB-201 and NDSB-256 outperformed other NDSBs in this screen, which led us to test for a binding site on the receptor. NDSB-201 and NDSB-256 contain pyridinium and benzyl groups, respectively, that can engage in arene–arene interactions with accessible aromatic amino acids [42]. The less effective zwitterionic additives, NDSB-195 and NDSB-221, possess quaternary ammonium groups, but lack an aromatic substituent that can participate in aromatic stacking (Figure 4B).

Addition of NDSB-201 to crystallization solutions affords a complex of NDSB-201 and TBRII-ECD-PR; electron density maps indicate NDSB-201 interacts with TBRII-ECD-PR in a shallow hydrophobic pocket composed of aromatic residues. The cationic pyridinium is oriented such that it stacks upon the aromatic phenylalanine in the binding site. Through this mode of interaction, NDSB-201 may stabilize the folded product in addition to preventing protein aggregation during folding, highlighting its potential to function as a pharmacological chaperone. Indeed, our structure reveals that the pyridinium group of NDSB-201 can stack with aromatic residues in a hydrophobic TBRII-ECD pocket. NDSB-201 binding should therefore stabilize the native disulfide bond in the pocket and mask hydrophobic surfaces within TBRII-ECD to stabilize the folded state and prevent protein aggregation.

To date, there are six structures of protein:NDSB-201 complexes in the Protein Data Bank [43–48], and in each of them, NDSB-201 interacts with aromatic residues. These interactions provide a rationale by which this compound promotes crystallization. In these previously reported structures, however, NDSB-201 was used primarily as a crystallization additive or a cryoprotectant after the correctly folded protein has been

obtained by other means. Our results illustrate that NDSB-201 can serve the dual roles of folding agent and crystallization additive. We postulate that for proteins that can be folded or refolded with NDSB-201, it will be advantageous to include NDSB-201 in crystallization trials.

The presence of a pocket for NDSB-201 on TBR11, suggests derivatives of this pyridinium-containing compound could be further optimized to generate probes of TGF- β signaling. Alignment of our NDSB-201-bound structure with that of the compounds of TBR11, TBR1 and TGF- β 3 (PDB ID 2PJY, Figure 6) reveals that the identified NDSB-201 binding site on TBR11 does not overlap with the regions bound by TGF- β 3 or TBR1-ECD. Thus, the NDSB-201 binding site is distinct. These data are intriguing in that they suggest NDSB-201 is unlikely to block TGF- β binding. Still, as demonstrated previously [8, 9], molecules that bind at a remote site can modulate TGF- β signaling. Our research not only suggests an alternative folding mechanism of TBR11-ECD by NDSB-201, but also reveals a potential small-molecule, binding pocket on TBR11-ECD that could be targeted for inhibitor or activator development.

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Table 1: Crystallographic statistics

Data collection statistics	
Wavelength (Å)	0.97919
Resolution range (Å)*	22.1 - 1.50 (1.55 - 1.50)
Space group	P 2 ₁ 2 ₁ 2 ₁
Unit cell (Å)	33.6, 40.5, 75.8
Total reflections	104,863 (10,083)
Unique reflections	17,064 (1,653)
Multiplicity	6.1 (6.1)
Completeness (%)	99.5 (98.5)
Mean I/σ(I)	18.7 (3.9)
Wilson B-factor (Å ²)	17.9
R-merge	0.054 (0.419)
R-meas	0.060 (0.458)
R-pim	0.024 (0.182)
Refinement Statistics	
Resolution range (Å)	21.4 - 1.50 (1.55 - 1.50)
R-factor	0.171 (0.194)
R-free (10%)	0.206 (0.242)
Number of atoms	1006
protein	890
ligand	26
water	90

Protein residues	102
RMSD (bonds, Å)	0.007
RMSD (angles, °)	1.18
Ramachandran favored (%)	98.0
Ramachandran outliers (%)	0.0
Average B-factor (Å ²)	22.4
protein	20.8
ligand	36.3
solvent	34.6

*Statistics for the highest-resolution shell are shown in parentheses. The coordinate and structure factors are deposited at the Protein Data Bank under accession code 4P7U.

$$R - merge = \frac{\sum_{hkl} |I_{hkl} - \langle I_{hkl} \rangle|}{\sum_{hkl} I_{hkl}}$$

$$R - meas = \frac{\sum_{hkl} \sqrt{\frac{n}{n-1}} |I_{hkl} - \langle I_{hkl} \rangle|}{\sum_{hkl} I_{hkl}}$$

$$R - pim = \frac{\sum_{hkl} \sqrt{\frac{1}{n-1}} |I_{hkl} - \langle I_{hkl} \rangle|}{\sum_{hkl} I_{hkl}}$$

$$Refinement R - factor = \frac{\sum_{hkl} |F_{hkl}^{obs} - F_{hkl}^{calc}|}{\sum_{hkl} F_{hkl}^{obs}}$$

Table 2: Composition of buffers used in the on-plate refolding screen.

Buffer	Additives	Redox buffer
75 mM Tris pH 8.0	Sorbitol (0.5, 1, 2 M)	2 mM GSH + 0.5 mM GSSG
	Urea (1, 2, 4 M)	2 mM GSH + 0.2 mM GSSG
	NaCl (100, 250 mM)	2 mM GSH + 0.05 mM
	PEG 3350 (0.05, 0.1, 0.5 %)	GSSG
	PEG 6000 (0.05, 0.1, 0.5 %)	
	MgCl ₂ +CaCl ₂ (1 mM each)	
	β-Cyclodextrin (5, 10 mM)	
	β-Cyclodextrin (5, 10 mM)	
	NDSB-195 (0.5, 1 M)	
	NDSB-201 (0.5, 1 M)	
	NDSB-221 (0.5, 1 M)	
	NDSB-256 (0.5, 1 M)	

Figure Legends

Figure 1: Representative ion-exchange chromatogram of the refolded TBRII-ECD-PR. The peak containing active TBRII-ECD-PR is marked with an asterisk. (1.5-column, see Figure_1.jpg)

Figure 2: Schematic depicting the on-plate refolding assay and ELISA scheme utilized to detect correctly folded Trx-TBRII-ECD (1-column, see Figure_2.jpg)

Figure 3: Preliminary screen for refolding conditions of Trx-TBRII-ECD. Relative amount of properly folded TBRII was measured by absorbance reading of the ELISA product. (2-column, see Figure_3.jpg)

Figure 4: A. Application of the Trx-TBRII-ECD refolding screen using a constant redox buffer: 2 mM GSH + 0.5 mM GSSG. Error bars represent standard deviations of triplicate measurements. Dashed line indicates the level of active Trx-TBRII-ECD when no additive is present, setting the baseline for effective additives.. B. Structures of the NDSB additives tested. (2-column, see Figure_4.jpg)

Figure 5: Crystallographic analysis of TBRII-ECD:NDSB-201 complex. A) Structure of NDSB-201 (orange) bound to TBRII-ECD-PR (blue). The $mF_o - DF_c$ difference map (black mesh, contoured 3.0σ above the mean) was calculated after the ligand was removed and the structure re-refined. B) Surface representation of the binding site. TBRII-ECD-PR surface atoms that are within 4 Å of the ligand are highlighted in yellow. Functional groups of hydrophilic side chains are not close enough to interact with NDSB-201. Most interactions are cationic π - π stacking and hydrophobic. (1-column, see Figure_5.jpg)

Figure 6: Alignment of the NDSB-201-TBR11-ECD-PR complex structure to a structure of the TGF- β signaling complex. The location of NDSB-201 is indicated by the boxes. (1-column, see Figure_6.jpg)

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