Structural and thermodynamical insights into the binding and inhibition of FIH-1 by the N-terminal disordered region of Mint3

Received for publication, May 8, 2021, and in revised form, October 9, 2021. Published, Papers in Press, October 14, 2021, https://doi.org/10.1016/j.jbc.2021.101304

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Edited by Wolfgang Peti

Mint3 is known to enhance aerobic ATP production, known as the Warburg effect, by binding to FIH-1. Since this effect is considered to be beneficial for cancer cells, the interaction is a promising target for cancer therapy. However, previous research has suggested that the interacting region of Mint3 with FIH-1 is intrinsically disordered, which makes investigation of this interaction challenging. Therefore, we adopted thermodynamic and structural studies in solution to clarify the structural and thermodynamical changes of Mint3 binding to FIH-1. First, using a combination of circular dichroism, nuclear magnetic resonance, and hydrogen/deuterium exchange–mass spectrometry (HDX-MS), we confirmed that the N-terminal half, which is the interacting part of Mint3, is mostly disordered. Next, we revealed a large enthalpy and entropy change in the interaction of Mint3 using isothermal titration calorimetry (ITC). The profile is consistent with the model that the flexibility of disordered Mint3 is drastically reduced upon binding to FIH-1. Moreover, we performed a series of ITC experiments with several types of truncated Mint3s, an effective approach since the interacting part of Mint3 is disordered, and identified amino acids 78 to 88 as a novel core site for binding to FIH-1. The truncation study of Mint3 also revealed the thermodynamic contribution of each part of Mint3 to the interaction with FIH-1, where the core sites contribute to the affinity (ΔG), while other sites only affect enthalpy (ΔH), by forming noncovalent bonds. This insight can serve as a foothold for further investigation of intrinsically disordered regions (IDRs) and drug development for cancer therapy.

Cancer is a major cause of death worldwide. Thus, the development of its treatment is of great interest in various research areas, including molecular targeted therapies. Although the types of cancer vary widely and a large part of the complex mechanism remains unclear, some common factors or phenomena could serve as promising targets for drug development. The Warburg effect is one of the phenomena that are considered to play a pivotal role in the survival and proliferation of tumor cells. It is characterized by increased aerobic glycolysis, even in the presence of abundant oxygen, which is frequently observed in malignant tumor cells. Since the glycolytic approach of energy production is less efficient in terms of ATP production than oxidative phosphorylation, which is the primary pathway of ATP production in most normal cells, the preference for aerobic glycolysis is assumed to have some implications on the differentiation between cancer cells and normal cells. Although the benefits of the Warburg effect for cancer cells are still unclear, numerous proposals have been made in the past decades, based on intensive research (1, 2). The possible explanations are an adaptation to hypoxia, which is the condition that most cancer cells confront, and/or acceleration of production of building blocks required for cell proliferation, which are provided as by-products of glycolysis. The mechanisms of the Warburg effect are also being gradually revealed along with the investigation of these functions. Hypoxia-inducible factor-1 (HIF-1) is a key transcription factor that regulates the expression of genes necessary for cellular responses to hypoxic conditions. The major examples of the proteins expressed from these genes include vascular endothelial growth factor, erythropoietin, glucose transporters, and several glycolytic enzymes such as phosphoglycerate kinase 1.

HIF-1 is composed of α and β subunits, and its transcriptional activity is regulated by factor inhibiting HIF-1 (FIH-1), an asparaginyl hydroxylase that prevents HIF-1α binding to p300/CPB through modification of the Asn803 residue. Since the enzymatic activity of FIH-1 requires Fe (II) and α-ketoglutaric acid (αKG), the regulation of HIF-1 is known to be oxygen-dependent. However, Sakamoto and Seiki (3) identified another factor, Mint3, a member of the X11 protein family, which controls the suppression activity of HIF-1 by FIH-1.

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Previous research by Sakamoto and Seiki (3) showed that Mint3 binds to FIH-1 through its N-terminal portion and inhibits the regulatory activity of FIH-1 by competing with HIF-1α. Accordingly, the protein–protein interaction (PPI) between Mint3 and FIH-1 is one of the critical pathways that enhances the activity of HIF-1, which is followed by a metabolic effect that is beneficial for cancer cells. Therefore, Mint3-FIH-1 PPI is a promising target for the development of cancer therapies.

Previous research has implied that the N-terminal region of Mint3 is the binding site of FIH-1 (3). It is noteworthy that the comprehensive protein sequence and functional database (UniProt) predicts that there are no secondary structures in the N-terminal region of Mint3 (UniProt: O96018). Although it is traditionally assumed that the function of a protein originates in its three-dimensional structure, many intrinsically disordered proteins (IDPs) and IDRs have recently been identified to be biologically functional (4–7). Moreover, most of these biological functions are activated by PPIs between IDPs/IDRs and their binding partners. Here, we hypothesized that the intrinsically disordered part of Mint3 has some core sites generating its inhibitory function for FIH-1. However, owing to the difficulties in analyzing the structural dynamics of IDPs/IDRs, clarification of the molecular mechanisms of their PPIs is still challenging. To test our hypothesis, we adopted a truncation approach, which is presumed to be effective for the IDP and performed a series of physicochemical analyses to clarify how Mint3 binds FIH-1 at a molecular level.

Results

Intrinsically disordered N-terminal region of Mint3

Mint3, also known as X11γ/APBA3, belongs to the X11 protein family. Although the remaining two members, Mint1/X11α/APBA1 and Mint2/X11β/APBA2, are neuronal proteins, Mint3 is ubiquitously expressed (8). All three X11s have a conserved C-terminal half that is composed of a phosphotyrosine-binding (PTB) domain, which is known to be the binding site of amyloid precursor protein (APP), and two PDZ domains (Fig. 1A). However, their N-terminal halves are variable. Although both Mint1 and Mint2 have an N-terminal binding site to munc18-1, which is a neural protein involved in synaptic vesicle exocytosis, Mint3 does not have one (3, 9). Instead, the N-terminal region (1–214) of Mint3 is reported to mediate binding to FIH-1 (Fig. 1A), although it has been predicted to be intrinsically disordered according to its primary structure in the UniProt database (3). To characterize the structure of the involved region, we expressed the N-terminal fragment (1–214) of Mint3 (hereafter called Mint3NT) and performed a series of structural analyses in solution.

Circular dichroism (CD) experiments were performed to measure the composition of the secondary structure of Mint3NT. Consequently, a negative peak at 200 nm, which is the hallmark of the random coil, was observed, and significant peaks that originate in secondary structures, especially α-helices and β-sheets, could not be observed (Fig. 1B). The result is suggesting that the isolated N-terminus of Mint3 is mostly composed of a random coil (10). Following that, differential scanning calorimetry (DSC) was performed to analyze the protein folding of Mint3NT. No significant peak caused by the collapse of the higher-order structure was observed in the thermal denaturation experiment (Fig. S1).

To determine the structural molecular details of the N-terminal regions of Mint3, hydrogen/deuterium exchange–mass spectrometry (HDX-MS) and nuclear magnetic resonance (NMR) spectroscopy were conducted. In the HDX-MS analysis, the exchange events are mediated by the conformational fluctuation of the protein. The exchange rate of amide hydrogen is influenced by the formation of a hydrogen bond, pH, and to a minor degree the temperature (11, 12). In this experiment, the deuteration processes were performed under the condition that the pH and the temperature are well controlled, so that we can focus on the influence of the formation of a hydrogen bond. Therefore, the amide hydrogen of an extended/flexible will be replaced by solvent deuterium at a higher rate, as compared with that of a folded/rigid region. In fact, the exchange rates of the whole Mint3NT were observed as an immediate saturation in the deuteration level, suggesting an intrinsically disordered state in solution (Figs. 1C and S2A, Table S1A). This is also supported by the NMR analysis. We measured the 1H-15N HSQC spectrum of Mint3NT and found that the dispersion of resonance peaks was very poor particularly along the 1H-axis, suggesting the absence of strong hydrogen-bonding interactions (Fig. 1D). Furthermore, the superposition of two HSQC spectra separately recorded for the fragment proteins of Mint3NT (Mint3(1–117) and Mint3(101–214)), Figure 1E reproduced the spectrum of whole protein (Mint3NT) very well, suggesting the absence of significant interaction between the N- and C-terminal regions of Mint3NT. However, a careful examination of the spectra revealed a slight change in the chemical shift values between the full length of Mint3NT (Mint3(1–214)) and fragment proteins (Mint3(1–117) or Mint3(101–214)) for several peaks. Based on the resonance assignment of Mint3(1–117), these exceptional residues were located at the terminal region of the fragment protein (residues 110–117), suggesting that the effect of cleavage of peptide bond was highly localized.

It should also be worth to mention here that the linewidth of each peak was very sharp even though the spectra were recorded at low temperature (4 °C), as is the case often observed for highly disordered proteins. In addition, the resonances derived from the side chains of three tryptophan residues were well resolved at (10 ppm, 130 ppm). All these results suggested that Mint3NT is mostly unstructured in solution. Also, we considered that truncation of Mint3NT is an effective methodology for the following detailed interaction analysis.

We also performed a titration experiment of 15N-labeled Mint3NT by unlabeled FIH-1 and found that the peak intensities for many residues were remarkably decreased in a concentration-dependent manner (Fig. S3). At the same time,
the opposite cases were also found, in which neither chemical shift value nor peak intensity was changed significantly by the addition of unlabeled FIH-1. These observations suggest that the interaction between $^{15}$N-Mint3NT and FIH-1 is site-specific. The assignment of the NMR resonances and the residue-specific identification of the interaction site will be published elsewhere (Maeda et al., in preparation).

**Physicochemical properties of the interaction**

Sakamoto and Seiki (3) identified Mint3NT as a construct that is sufficient for binding to FIH-1 in immunological assays. We performed a couple of thermodynamic analyses on the interaction between Mint3 and FIH-1 using isothermal titration calorimetry (ITC), to confirm the interaction quantitatively. The Mint3 constructs used in these experiments were Mint3PTB (1–361), which contains the PTB domain within the C-terminal half, and Mint3NT. Exothermic interactions were observed for both the Mint3s. Moreover, the thermodynamic parameters [the binding free energy ($\Delta G$), binding enthalpy ($\Delta H$), and binding entropy ($-T\Delta S$)] were the same for Mint3PTB and Mint3NT, within the error range (Figs. 1A and 2, A and B). The PTB domain did not contribute significantly to Mint3NT binding. Thus, we regarded that the Mint3NT

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**Figure 1. Molecular information of Mint3.** A, Mint3 constructs used in this study. B, CD spectrum of Mint3NT. C, HDX-MS chart of Mint3NT. D and E, comparison of the 1H-15N HSQC spectra of Mint3NT (D) and the superposition (E) of the fragment proteins of Mint3NT, corresponding to the N-terminal half (Mint3(1–117) in blue) and the C-terminal half (Mint3(101–214) in red). All spectra were recorded under nearly physiological conditions at pH 7.3 and 4 °C.
construct was of sufficient length in the context of binding to FIH-1. Of note, the binding of Mint3NT to FIH-1 produced a large change in enthalpy ($\Delta H = -32.2 \pm 0.7$ kcal mol$^{-1}$) and entropy ($-T\Delta S = -23.6 \pm 0.8$ kcal mol$^{-1}$) (Fig. 2B). This phenomenon can be interpreted by entropy–enthalpy compensation, namely (i) the large negative $\Delta H$ suggests the formation of a large number of intra- and intermolecular bonds, and (ii) the large positive $-T\Delta S$ reflects the loss of conformational flexibility of Mint3NT through the interaction. Additionally, the binding stoichiometry ($N$ value) for Mint3NT was estimated as $0.48 \pm 0.02$. Since in the ITC experiment, Mint3 and FIH-1 were loaded into the syringe and cell, respectively, this number suggests that the binding molar ratio between Mint3 and FIH-1 is 1:2. To confirm the oligomerization state of FIH-1, size-exclusion chromatography (SEC) and SEC–multangle laser scattering (SEC-MALS) measurements were conducted. In the SEC measurement, the chromatogram showed a single peak at the elution volume of the molar mass of the FIH-1 dimer. The SEC-MALS measurement also provided a chromatogram with a single peak, and the measured molar mass was approximately 80 kDa, which is consistent with that of the FIH-1 dimer (Fig. S4). Notably, although the concentration of FIH-1 used in SEC-MALS ($\sim70$ μM) was approximately ten times greater than that of SEC ($\sim7$ μM), the results suggested that most FIH-1 forms a dimer under both the conditions. Considering that the concentration of FIH-1 used in the ITC measurement (30 μM) was between these two concentrations, it was clear that FIH-1 presented as a dimer uniformly. Therefore, the stoichiometry can be interpreted to indicate that one Mint3 binds to a dimer of FIH-1.

The abovementioned structural analysis depicted the intrinsically disordered state of isolated Mint3NT, and the thermodynamic parameters of the interaction suggested a loss of conformational flexibility for Mint3NT in a complex with FIH-1. CD analysis was further performed to determine whether Mint3NT forms secondary structures through this interaction. We compared the spectrum obtained from the Mint3NT-FIH-1 complex with a theoretical spectrum generated by merging the spectra of isolated Mint3NT and FIH-1 (Fig. S5). As both the spectra were similar, Mint3NT did not seem to form secondary structures through binding to FIH-1, suggesting that a significant conformational change was not observed, at least at the level of the secondary structures.

**Thermodynamic/kinetic contribution of each part of Mint3NT to the interaction**

In a previous study, the 101 to 110 region within Mint3NT was identified as a core region responsible for its binding to FIH-1 (13). Accordingly, it is hypothesized that Mint3NT can be divided into the binding and nonbinding sites. To clarify the contribution of each part of Mint3NT, especially in quantitative terms, we performed a series of thermodynamic analyses for several truncated and mutated versions of Mint3NT using ITC (Fig. 2, C and D). Some constructs (1–117, 1–92, 93–117, Mint3NTmut) were observed in the exothermic reactions. The

![Figure 2. Thermodynamic profiles of the interaction between truncated Mint3s and FIH-1. A, ITC profiles of Mint3PTB and Mint3NT binding to FIH-1. B, thermodynamic parameters of the interactions between FIH-1 and Mint3. C, deletion mutants of Mint3NT. D, thermodynamic parameters of the interactions between FIH-1 and deletion mutants of Mint3NT.](image-url)
N values of these truncated mutants were also around 0.5, suggesting that they bind to a dimer of FIH-1, similarly to Mint3NT.

A short peptide of Mint3 (93–117) containing the predicted core region (101–110) was observed in the ITC measurement, and its ΔG was calculated to be −7.5 ± 0.1 kcal mol⁻¹, based on the measured equilibrium dissociation constant (K_D = 3.0 ± 0.6 µM). Next, alanine mutations were introduced into the peptide 93 to 117 (GILSAEAGRD → AAAAAAGRD, 93–117mut), since this Alanine mutant is the sequence that was designed in the past and had successfully lost the activity (13). The interaction was found to be completely disrupted. Notably, the introduction of the same mutations into the whole Mint3NT (Mint3NTmut) weakened the binding (K_D = 1.31 µM, ΔH = −13.9 kcal mol⁻¹, −TΔS = 5.9 kcal mol⁻¹), but did not disrupt the interaction completely (Fig. S6). Thus, we considered that the predicted region (101–110) was just one of the binding sites.

After confirming that Mint3 has a binding site at 101 to 110, further truncation analyses were conducted. First, the N-terminal half (1–117) of Mint3NT, which contains the binding site (101–110), also exhibited significant exothermic interaction with FIH-1. The measured K_D was 0.65 ± 0.07 µM, and the calculated ΔG was the same (within error) with Mint3NT. Notably, the measured ΔH for the N-terminal half of Mint3NT was approximately half of that for Mint3NT (Fig. 2, C and D). Next, a Mint3 fragment (1–92) that lacked the binding site (101–110) was constructed. Remarkably, although the fragment (1–92) displayed a significant decrease in exothermic interaction, it retained its binding ability to FIH-1 (Fig. 2, C and D, and Table S2). Therefore, we hypothesized that the N-terminal half (1–117) of Mint3NT is sufficient to generate affinity (ΔG), and that the C-terminal half (111–214) does not contribute to ΔG because its contribution to ΔH is canceled by −TΔS. Furthermore, there is at least one more binding site in the 1 to 92 region of Mint3. Thus, we found the following characteristics of the binding process from the ITC experiments: (i) One dimer of FIH-1 binds to one Mint3NT monomer; and (ii) there are other interaction sites in Mint3NT, apart from Mint3 (93–117).

**Role of the dimer interface of FIH-1**

According to the binding stoichiometry measured with ITC, Mint3NT binds to a dimer of FIH-1. Thus, we hypothesized that the pivotal binding site of FIH-1 is located around the dimer interface, from the standpoint of symmetry. Although FIH-1 monomers strongly interact with each other by meshing their hydrophobic C-terminal helices, the substitution of Leu340 for arginine is known to disrupt the dimerization (14). Hence, a single-point mutant FIH-1L340R was constructed to
examine the role of the dimer interface (Fig. S7). As expected, FIH-1L340R was mostly eluted in the monomeric fraction in SEC (Fig. 3A).

To examine the conformational identity between the wild-type and monomeric mutant, CD measurements and differential scanning fluorometry (DSF) were conducted. In CD measurements, a recession of α-helicity was observed for the monomeric mutant. In quantitative terms, α-helix decreased by 5.4% and random-coil increased by 5.8%, whereas β-sheet and turn were practically unchanged. These results suggest that the mutation may disturb the structure of FIH-1, especially the α-helices (Fig. 3B). In DSF, the monomeric mutant showed lower thermal stability than the wild-type, and notably, the baseline level of the monomeric mutant was higher than that of the wild-type (Fig. S8). Because SYPRO Orange, which is a protein dye used in the DSF experiment, binds nonspecifically to the hydrophobic surface and emits fluorescence, the elevated baseline is likely to represent the exposure of the hydrophobic dimer interface of FIH-1L340R. Additionally, HDX-MS was applied to depict the structural differences between the wild-type and monomeric mutant in detail. The difference in the deuteration level was observed only around the C-terminal α-helices, namely the dimer interface (Figs. 3C and S2B, Table S1B). According to the results of CD, DSF, and HDX-MS, the substitution of Leu340 for arginine can interrupt the dimerization, by loosening the interfacial α-helices, and the effect on the structure is limited to the periphery of the dimer interface.

Subsequently, binding analysis was performed using ITC, and significant disruption of the binding to Mint3NT was observed for FIH-1L340R. Notably, the binding free energy (ΔG) decreased significantly (Figs. 2, A and B and 3D). Since the L340R mutation only affects the dimer interface according to the HDX experiment, we concluded that the disruption of interaction was caused by the structural change in the dimer interface. Therefore, it is clear that the dimer interface of FIH-1 plays a critical role, namely in terms of contribution to the ΔG in the interaction with Mint3.

Regions of FIH-1 involved in the interaction

Oxygen-dependent regulation of HIF-1 activity is known to be mediated by the hydroxylation of Asn803 at its HIF-1α subunit. This modification is accomplished by the catalytic activity of FIH-1 as an asparaginyl hydroxylase. FIH-1 has a catalytic center, Fe(II), in its β-strand jellyroll core with its cosubstrate αKG. Hence, FIH-1 is categorized into the Fe(II) and αKG-dependent dioxygenases (15). Since previous research suggested that Mint3 inhibits the regulatory activity of FIH-1 by competing with HIF-1α in the PPI, we hypothesized that the catalytic pocket, an obvious binding site of HIF-1α, may be involved in the interaction with Mint3 (3). First, ITC measurements were performed to examine the effect of the cofactors on the interaction between FIH-1 and Mint3NT. The absence of Fe(II) and αKG reduced the ΔG by 0.5 and 0.7 kcal mol⁻¹, respectively, and the decrease in ΔG caused by the elimination of both cofactors (1.3 kcal mol⁻¹) was approximately the sum of each value (Fig. 4, A and B). According to the values, each cofactor is not critical but is certainly involved in the interaction. In addition, as the effect on ΔG is additive, it is likely that each cofactor independently affects the binding affinity. Moreover, αKG influenced the ΔH, whereas Fe (II) did not (Fig. 4, A and B). Therefore, we assumed that the cofactors, especially αKG, may stabilize the conformation of FIH-1 that binds to Mint3.

Figure 4. Effects of FIH-1 cofactors on binding to Mint3NT and FIH-1 structure. A, ITC profiles and B, thermodynamic parameters of the interaction between FIH-1 and Mint3NT, in the presence/absence of αKG and/or EDTA. C, differences between HDX-MS charts for FIH-1, in the absence and presence of αKG. D, structural image of FIH-1, with the differences in HDX-MS data shown in blue (PDB ID: 1MZF).
Next, HDX-MS was performed against FIH-1 ± αKG, to further investigate its effect on the structure of FIH-1. Upon comparing the deuteration levels of the apo- and holo-forms of FIH-1, a significant difference was observed in several regions (Figs. 4C and S2C, Table S1C). Merging the results of HDX-MS and the crystal structure revealed that some of the varied regions were around the binding site of αKG, namely the prespecified catalytic pocket. However, it was also discovered that αKG affects not only its binding site, but also its adjacent helical region located at the outer edge of the FIH-1 dimer (Fig. 4D). Subsequently, HDX-MS was conducted against FIH-1 ± Mint3NT, to detect the binding site. As a result, the region in FIH-1, the deuteration level of which was shown to be affected by the presence of Mint3NT, corresponded to the αKG-related site (Figs. 5, A and B and S2D, Table S1D). Accordingly, the corresponding region, namely the outer helices (hereafter called the HDX region), is likely to be involved in the interaction with Mint3NT. Meanwhile, it should be noted that there was no information about the dimer interface because it is buried and intrinsically hard to deuterate.

To validate the role of the HDX region, mutation analysis was performed using ITC. Six amino acids within the HDX region (Tyr71, Lys75, Glu79, Asp152, Lys157, and Asn171) were selected based on their orientation and polarity and replaced with Ala. In addition, three amino acids (Glu29, Asp248, and Glu250) apart from the HDX region were replaced with Ala, as negative controls (Fig. 5B). The thermodynamic parameters of binding to Mint3NT were measured three times for each FIH-1 mutant, and the standard deviation was calculated as the experimental error. In terms of the ΔG, there was no significant difference between the mutants, including the negative controls. On the other hand, the ΔH commonly decreased by approximately 3 kcal mol⁻¹ for most of the mutants. According to the result of the negative control, the ~3 kcal mol⁻¹ decrease in ΔH was probably an experimental error caused by the mutation. However, D152A and K157A showed a significant decrease of approximately 8 kcal mol⁻¹, which was larger than 3 kcal mol⁻¹ (Fig. 5, B and C). Notably, the CD spectra of each mutant suggested that even a single mutation in FIH-1 may affect its structure, but the magnitude of the structural change did not correlate with the decrease in ΔH (Fig. S9). Therefore, a common decrease in ΔH can be expressed by structural fluctuations. However, the ΔH decrease caused by D152A and K157A is possibly caused by their interaction with Mint3NT. In conclusion, there is a possibility of the HDX region of FIH-1 being involved in the interaction with Mint3 without contribution to ΔG.

Identification of the other binding site in Mint3NT

Although the 101 to 110 region was confirmed to be a pivotal binding site, there is presumably another binding site that exists within the N-terminal half (1–117) of Mint3NT. To identify the other predicted binding sites, we performed HDX-MS for isolated Mint3NT (apo-form) and Mint3NT-FIH-1.
complex (holo-form), since the HDX-MS experiment can provide structural information in peptide-scaled resolution. First, to examine whether the HDX-MS system is feasible for detecting the binding site of Mint3NT, the results of the peptides that contain the known binding site (101–110) were used as a positive control. For instance, as expected, the peptides IADAHGLLSAEAGRDDL (96–112) and AGRDDLLGL (107–115) exhibited a significant difference in the deuteration level between the apo- and holo-states. Moreover, the apo- and holo-states of the peptide PEPPAGAQSPE (184–195), which is located in the C-terminal half of Mint3NT, did not differ in terms of deuteration levels. This result is consistent with the speculation that the C-terminal half of Mint3NT does not contribute to the binding to FIH-1 (Figs. 6A and S2E, Table S1E). To confirm system reliability, other peptides within the N-terminal half of Mint3NT were also investigated. An obvious difference in the deuteration level between the apo- and holo-states was observed for the peptides lying immediately before the known binding site, for example, LVGPSGGAPCPLHIAHTGHGLASQE (71–95) (Fig. 6A). To determine the exact region that is important for binding, the amino acid sequence of human Mint3 was compared with that of mouse, rat, bovine, dog, and chicken. Consequently, a well-conserved region (78–88) was found within the peptide of concern. Following that, a new mutant of Mint3NT was constructed by altering five amino acids with a large side chain to alanine within the species-common region (hereafter called Mint3mut2) (Fig. 6B). In the binding analysis using ITC, a remarkable decrease in binding affinity was observed for Mint3mut2, as compared with Mint3NT (Fig. 6C). In addition, a short fragment (78–117) of Mint3NT containing both the detected binding sites was constructed. Compared with the predescribed peptides 93 to 117, there was an increase in the binding affinity of peptides 78 to 117, owing to the presence of a newly detected binding site (Fig. 6, D and E). To confirm the defect of interaction with FIH-1 in the full-length mutant Mint3 with the five alanine substitutions in the 78 to 88 amino acid region (Mint3mut2) within cells, FLAG-tagged FIH-1 was expressed in 293FT cells along with either V5-tagged wild-type Mint3 (WT-Mint3) or Mint3mut2 (Fig. 7A). Immunoprecipitation of FLAG-tagged FIH-1 coprecipitated V5-tagged Mint3mut2 much less than WT-Mint3. Thus, the conserved amino acids of Mint3 in the 78 to 88 amino acid region are essential for binding to FIH-1 even within cells. Mint3 binds to FIH-1

**Figure 6. Novel binding site of Mint3NT.** A, HDX-MS profiles of Mint3NT peptides, in the absence and presence of FIH-1. B, comparison of the amino acid sequence of Mint3NT among animal species, and amino acid sequence of Mint3mut2. C, ITC profiles of the interaction between FIH-1 and Mint3NT mutants. D, thermodynamic parameters of Mint3(93–117) and Mint3(78–117). E, amino acid sequences of Mint3(93–117) and Mint3(78–117).
and thereby activates HIF-1α transactivation activity (3, 16). Thus, to confirm the effect of the Mint3 mutation on the regulation of HIF-1α, we used a luciferase reporter assay that can specifically monitor FIH-1-mediated regulation of HIF-1α transactivation activity as described previously (3, 17). Expression of WT-Mint3 increased the HIF-1α transactivation activity by more than 3500% in 293FT cells, while that of Mint3mut2 increased it by only 500% even though Mint3mut2 was expressed at the comparable levels to WT-Mint3 in 293FT cells (Fig. 7B). Thus, the conserved amino acids of Mint3 in the 78 to 88 amino acid region are also important for activating HIF-1α transactivation activity in cells. Accordingly, we concluded that the 78 to 88 region is a major binding site in Mint3NT. Unfortunately, the peptides 78 to 88 were too poorly water-soluble to be synthesized and evaluated in vitro.

Description of the Mint3-FIH-1 complex

Through thermodynamic analysis using ITC and structural analysis using HDX-MS, we made four main observations: (i) Mint3 has two core regions (77–87 and 101–110), which contribute to its ΔG to FIH-1. (ii) The interfacial structure of the FIH-1 dimer plays a critical role in the interaction with Mint3. (iii) The regions of Mint3 other than the cores are not involved in the ΔG, while they do affect the ΔH. (iv) Some amino acids (Asp152 and Lys157) in the outer helices of the FIH-1 dimer do not contribute to the ΔG, but contribute to the ΔH. Based on these results, we generated a model of the Mint3-FIH-1 complex, composed of “binding” part and “touching” parts (Fig. 8). Here, the “binding” parts indicate the parts contributing to the ΔG, and “touching” parts represent the parts involved in the interaction by affecting the ΔH.

Figure 7. Confirmation of the role of novel binding in cells. A, immunoblotting using mouse anti-V5 antibody and rabbit anti-FLAG polyclonal antibody in 293FT cells. Left side is a sample of the whole cell lysate, while right side is the sample immunoprecipitated using an anti-FLAG polyclonal antibody. B, reporter assay expressed vector alone, wild-type Mint3, or Mint3mut2 in 293FT cells. Left side is the immunoblotting data of the whole cell lysate using an anti-V5 antibody, while right side is the data for the luciferase activities.

Figure 8. The interaction model between FIH-1 and Mint3NT, as inferred from this study. The three-dimensional structure is that of FIH-1 dimer, while the string-like illustration is a schematic for Mint3NT (PDB ID: 1MZF).
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rather than the $\Delta G$. In the “binding” part, the core regions (77–87 and 101–110) of Mint3 are interacting with the dimer interface of FIH-1, reflecting their critical roles in the $\Delta G$. On the other hand, the “touching” parts are composed of 1 to 76/111 to 214 of Mint3 and the outer helices of FIH-1 dimer, indicating that both of them contribute to producing balanced favorable enthalpy and unfavorable entropy through the interaction.

To confirm that the description was physically possible, we estimated the molecular size of the proteins. The scale of FIH-1 was obtained from its crystal structure, by measuring the distance between the outer helix (Lys157) and the center of the dimer interface (Leu340), which represents the semimajor axis of the FIH-1 dimer. As a result, the value was estimated at 57.0 Å (Fig. S10). Meanwhile, the radius of gyration $R_g$ of Mint3NT (1–214) can be estimated by $R_g = R_0^{*}N^*$ with $R_0 = 0.1927$ nm and $N^* = 0.598 \pm 0.028$, where $N$ represents the number of the amino acids (18). Thus, the approximated diameter of free Mint3NT, namely $2^*R_g$ is 95.4 Å, which suggests that free Mint3NT is a little smaller than FIH-1. As Supporting information, the fully extended lengths of 1 to 76 and 111 to 214 regions of Mint3NT were estimated at 273.6 Å and 374.4 Å, respectively, according to the average end-to-end distance of one amino acid (3.6 Å) (19–22). Comparing the physical size of the two proteins, Mint3NT is possible to “bind” to the dimer interface and “touch” at the outer helix of the FIH-1 simultaneously, while it needs to be extended a little through the interaction.

Discussion

Mint3 is a member of the X11 protein family that has conserved C-terminal domains, such as the binding site of APP and unique N-terminal domains (23, 24). Previous research has demonstrated that the N-terminal half of Mint3 binds to FIH-1 and acts as an activator of the HIF-1 pathway, which is involved in cellular responses to hypoxia and leads to the proliferation or metastasis of cancer cells (4, 13). However, the conformation and mechanism of the interaction between Mint3 and FIH-1 remain unclear. Thus, using CD, HDX-MS, and NMR analyses, we first demonstrated that the N-terminal portion, which is the functional part of Mint3, is highly disordered in its isolated form. This result is consistent with the emerging knowledge of IDRs. In particular, their biological roles include signal transfer, regulation, transcription, and replication (25, 26).

In general, many IDRs undergo transitions to more ordered or folded states upon binding to their targets, to execute their biological functions. The kinase-inducible transcriptional activation domain (KID) of cyclic AMP response element-binding protein (CREB) is one of the most characterized examples (25, 27, 28). Although KID is intrinsically disordered, it forms an orthogonal helix by binding to the CREB-binding protein (CBP) (29). However, although the unfavorable entropy suggested a loss of flexibility in binding to FIH-1, coupled folding and binding could not be observed upon CD measurement of Mint3NT. Therefore, we presumed that a truncation approach is feasible and effective for characterizing the PPI between Mint3NT and FIH-1.

To characterize the binding, we conducted thermodynamic and structural analyses using ITC and HDX-MS, respectively, since the classical approaches, particularly crystallographic methods, have little relevance for a mostly disordered protein such as Mint3NT. Although these methods cannot directly visualize the interaction, we succeeded in identifying the regions involved in the interaction, “binding” and “touching” sites, and revealing their different roles in thermodynamics, as mentioned above (Fig. 8). The “binding” sites of the dimer interface of FIH-1 for 78 to 88 and 101 to 110 regions of Mint3NT, which represent the regions that contribute to the $\Delta G$, are a usual interpretation. In particular, we proposed that the 78 to 88 amino acids in Mint3 are a critical binding region to FIH-1. On the other hand, the “touching” sites, which indicate the parts that do not contribute to the $\Delta G$ but certainly influence the $\Delta H$, are somewhat conceptual. Some previous reports can provide hints for interpreting the roles of the “touching” site. Tummino et al. (30) suggest that enhancing the enthalpic contributions of protein interactions would likely favor slower dissociation and, hence, a longer residence time. In addition, it has been reported IDPs/IDRs can regulate the binding kinetics (31). Therefore, the “touching” sites may affect the rate constant of the interaction, although further investigation is required in this regard.

Our model for the complex of Mint3NT and FIH-1 mentioned above is a description that can consistently explain all the results. Notably, the region between the “binding” site and the “touching” site of Mint3 is delineated to cover the binding pocket of aKg in FIH-1. This model is consistent with the fact that Mint3 abrogates the ability of FIH-1 to modify HIF-1α, by competing with HIF-1α to bind to the aKg binding site of FIH-1 (3). Although the detailed conformation of Mint3 needs to be determined directly using some experimental methods, this model can provide us insight into the suppression mechanism of Mint3 toward HIF-1 binding to FIH-1.

In addition, the detailed binding mechanism can be used as a guide for drug development in cancer therapy. According to the Mint3-FIH-1 complex model, there are two possible strategies for disturbing the suppression activity of Mint3. The first strategy is to aim at a critical site of binding affinity, which in this case is the “binding” site, as it seems to be more effective in inhibiting the interaction with FIH-1. The second strategy is to focus on the “touching” site, to prevent Mint3 from covering the aKg-binding site. The merit of the second scenario is that the inhibitor with lower affinity may still be effective in this case, as compared with the case with aiming at the “binding” site.

In conclusion, we found that the N-terminal region of Mint3 has properties corresponding to IDRs, and that this region interacts with FIH-1 in an enthalpy-driven manner. Using Mint3NT mutants, we suggest that the core region of Mint3 binds to FIH-1 in such a way that a single molecule rests on the FIH-1 dimer interface. We have elucidated the structural and thermodynamic features of how Mint3 suppresses FIH-1,
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and this insight can serve as a foothold for further investigation of IDRs and drug development for cancer therapy.

Experimental procedures

Recombinant protein expression and purification

The coding sequence for human Mint3NT (1–214) with a C-terminal hexa-histidine (His6) tag, and human FIH-1 (full-length) with an N-terminal His6 tag and cleavage site (ENLYFQ) of TEV protease were inserted into the pET28 vector. The plasmids were transformed into BL21 (DE3) cells and cultured at 28 °C in Luria broth medium containing 50 μg/ml kanamycin. M9 minimal medium containing 0.5 g/l $^{15}$NH$_4$Cl was used for the preparation of uniformly $^{15}$N-labeled protein. Protein synthesis was induced by the addition of 1 mM Isopropyl β-D-1-thiogalactopyranoside (IPTG) to the culture, when its optical density at 600 nm reached a value of ~0.6, and further culturing at 20 °C for 16 h. Cells were harvested by carrying out a round of centrifugation at 7000 g for 30 min, following which the supernatants were loaded onto a Ni-NTA Agarose (Qiagen) column that had been pre-equilibrated with the binding buffer. The column was washed with 20 column volumes of wash buffer composed of 20 mM Tris-HCl (pH 8.0), 500 mM NaCl, and 5 mM imidazole. The lysates were centrifuged at 40,000 g for 10 min and disrupted by sonication in a binding buffer composed of 20 mM Tris-HCl (pH 8.0), 500 mM NaCl, and 5 mM imidazole. The column was washed with 20 column volumes of wash buffer composed of 20 mM Tris-HCl (pH 8.0), 500 mM NaCl, and 20 mM imidazole. His$_6$-tagged Mint3NT and His$_6$-tagged FIH-1 were eluted with 10 ml of an elution buffer composed of 20 mM Tris-HCl (pH 8.0), 500 mM NaCl, and 200 mM imidazole. His$_6$-tagged Mint3NT and His$_6$-tagged FIH-1 were cleaved using His$_6$-tagged TEV protease. Mint3NT was purified using SEC with a HiLoad 26/60 Superdex 200 pg column (GE Healthcare) at 4 °C. The His$_6$-tag of FIH-1 was cleaved using His$_6$-tagged TEV protease. Mint3NT was purified using SEC with a HiLoad 26/60 Superdex 200 pg or 75 pg column (GE Healthcare) at 4 °C. The monomer fraction was collected from the purification. FIH-1 was purified using SEC with HiLoad 26/60 Superdex 200 pg at 4 °C. The elution profiles of both the proteins were monitored at a wavelength of 280 nm. The mutants/fragments of Mint3 and FIH-1 were prepared as described above.

Circular dichroism analysis

Far ultraviolet (UV) CD spectra of Mint3NT, FIH-1, and FIH-1 mutants were measured on a J-820 spectropolarimeter (Jasco) at 25 °C in 35 mM Na-phosphate (pH 7.5), 300 mM NaCl, and 500 μM αKG. The concentrations of the protein samples were maintained in the range of 5 to 10 μM, as appropriate to keep the HT voltage under 600 V. Each spectrum was generated upon accumulation of five measurements.

Hydrogen/deuterium exchange–mass spectrometry analysis

Mint3NT, FIH-1, and its mutants were prepared in Na-phosphate buffer composed of 35 mM Na-phosphate (pH 7.5) and 300 mM NaCl at a final concentration of 1.0 mg ml$^{-1}$. Each protein was diluted tenfold with Na-phosphate buffer (the solvents are as mentioned before) in heavy water (D$_2$O). The diluted solutions were then incubated separately at each time and at each low temperature as shown in Table S1. Deuterium-labeled samples were quenched by diluting about tenfold with quenching buffer at pH 2.3, composed of 2 M guanidine hydrochloride and 200 mM citric acid. All of the processes described above were performed automatically using HDX Workflow Solution (LEAP Technologies), and all the time points were determined in three independent labeling experiments. After quenching, the solutions were subjected to online pepsin digestion followed by LC/MS analysis using an UltiMate 3000 RSLCnano System (Thermo Fisher Scientific) connected to a Q Exactive Plus mass spectrometer (Thermo Fisher Scientific). Online pepsin digestion was performed with a Poroszyme Immobilized Pepsin Cartridge 2.1 × 30 mm (Waters Corporation) in formic acid solution (pH 2.5) for 3 min at 8 °C, at a flow rate of 50 μl min$^{-1}$. Desalting and analytical processes after pepsin digestion were performed using Acclaim PepMap 300 C18 (1.0 × 15 mm; Thermo Fisher Scientific) and Hypersil GOLD (1.0 × 50 mm; Thermo Fisher Scientific) columns. The mobile phase was 0.1% formic acid solution (buffer A) and 0.1% formic acid containing 90% acetonitrile (B buffer). The deuterated peptides were eluted at a flow rate of 45 μl min$^{-1}$, with a gradient of 10% to 90% of buffer B for 9 min. The conditions of the mass spectrometer were as follows: electrospray voltage, 3.8 kV; positive ion mode, sheath and auxiliary nitrogen flow rate at 20 and two arbitrary units; ion transfer tube temperature at 275 °C; auxiliary gas heater temperature at 100 °C; and a mass range of m/z 200 to 2000. Data-dependent acquisition was performed using a normalized collision energy of 27 arbitrary units. The MS and MS/MS spectra were subjected to a database search analysis using the Proteome Discoverer 2.2 (Thermo Fisher Scientific). Analysis of the deuteration levels of the peptide fragments was performed based on the MS raw files, by comparing the spectra of deuterated samples with those of nondeuterated samples, using the HDX Examiner software (Sierra Analytics). Data summary for each HDX-MS was shown in Table S1.

Nuclear magnetic resonance spectroscopy

$^1$H-$^{15}$N HSQC spectra were measured on an Avance I 600 spectrometer (Bruker BioSpin) equipped with a TXI-triple resonance triple gradient probe at 4 °C. The sample concentration was 50 μM $^{15}$N-labeled protein dissolved in 50 mM Na-phosphate buffer (pH 7.3) containing 10% D$_2$O. The spectra were processed using NMRPipe (32).

Differential scanning calorimetry analysis

Thermal denaturation experiments were performed using a Nano DSC (TA instruments). Mint3NT was dialyzed against Na-phosphate buffer composed of 35 mM Na-phosphate (pH 7.5), 300 mM NaCl at 4 °C for 16 h, and concentrated to 50 μM (∼1.2 mg ml$^{-1}$) using Amicon Ultra-15 Centrifugal Filter Units (EMD Millipore). The protein was loaded into the sample cell and heated from 10 to 100 °C, at 60 °C h$^{-1}$. 
Isothermal titration calorimetry

Calorimetric titration of Mint3NT with FIH-1 was performed using a MicroCal iTC200 system (GE Healthcare) at 25 °C. Both proteins were dialyzed against Na-phosphate buffer composed of 35 mM Na-phosphate (pH 7.5), 300 mM NaCl, and 500 μM aKG at 4 °C for 16 h and concentrated using Amicon Ultra-15 Centrifugal Filter Units. At each injection (total 25 times), 1.5 μl of 150 μM Mint3NT solution was added to a sample cell containing 30 μM FIH-1 solution. All titration data were analyzed by fitting to a single set of site models using Origin 7 (OriginLab). Calorimetric titration of Mint3NT mutants/fragments and FIH-1 mutants was performed as described above.

SEC–multiangle laser scattering

MALS was detected using DAWN8+ (Wyatt Technology). FIH-1 was dialyzed against Na-phosphate buffer composed of 35 mM Na-phosphate (pH 7.5), 300 mM NaCl, and 500 μM aKG at 4 °C for 16 h, and concentrated to 70 μM using Amicron Ultra-15 Centrifugal Filter Units. The protein sample (50 μl) was loaded into a 10/300 Superdex 200 GL (GE Healthcare) column connected to the MALS detection unit for SEC. During the experiment, UV detection was performed using an SPD-10A UV/Vis Detector (Shimadzu). The obtained data were processed and analyzed using ASTRA (Wyatt Technology).

Differential scanning fluorometry analysis

The thermal shift assay was performed on a CFX Connect Real-Time System (Bio-Rad). FIH-1 wild-type and L340R were dialyzed against Na-phosphate buffer composed of 35 mM Na-phosphate (pH 7.5), 300 mM NaCl, and 500 μM aKG at 4 °C for 16 h, and concentrated to 10 μM using Amicon Ultra-15 Centrifugal Filter Units. One thousandth of the protein sample volume of SYPRO Orange Protein Gel Stain (5000× concentrated in Dimethyl sulfoxide (DMSO); Invitrogen) was added, and the proteins containing 5× SYPRO Orange were loaded onto Hard-Shell 96-well PCR Plates (Bio-Rad) for measurement. To denature each protein, an α-Mint3NT solution and μM FIH-1 solution were added to a sample cell containing 30 μl of 150 μM Mint3NT solution and 1.5 μl of 150 μM FIH-1 solution. The thermal shift assay was performed on a CFX Connect Real-Time System (Bio-Rad). FIH-1 wild-type and L340R were dialyzed against Na-phosphate buffer composed of 35 mM Na-phosphate (pH 7.5), 300 mM NaCl, and 500 μM aKG at 4 °C for 16 h, and concentrated to 10 μM using Amicon Ultra-15 Centrifugal Filter Units. The protein sample volume of SYPRO Orange Protein Gel Stain (5000× concentrated in Dimethyl sulfoxide (DMSO); Invitrogen) was added, and the proteins containing 5× SYPRO Orange were loaded onto Hard-Shell 96-well PCR Plates (Bio-Rad) for measurement. The denaturation of each protein was observed via fluorescence emitted from SYPRO Orange upon absorption to the exposed hydrophobic surface.

Cellular analysis using immunoprecipitation and reporter assay

293FT cells, which are derived from HEK293 human embryonic kidney cells and express the simian virus large T antigen, were purchased from Thermo Fisher Scientific. Cells were cultured at 37 °C in a humidified atmosphere containing 5% CO2 in high-glucose Dulbecco’s Modified Eagle Medium (DMEM) (Thermo Fisher Scientific) containing 10% fetal bovine serum, 100 units/ml penicillin, and 100 μg/ml streptomycin (Merck).

pcDNA3 (Thermo Fisher Scientific) expression vectors for V5-tagged Mint3 and FLAG-tagged FIH-1 were constructed as previously described (3). Expression vectors expressing mutant Mint3 (Mint3mut2) were generated using PCR-based methods.

Immunoprecipitation was performed as previously described (3, 33). Briefly, 293FT cells were cotransfected with expression plasmids encoding a V5-tagged Mint3 construct and a FLAG-tagged FIH-1 construct using Lipofectamine 2000 (Thermo Fisher Scientific). Cells were lysed in lysis buffer 24 h after transfection and centrifuged at 20,000g for 15 min at 4 °C. Supernatants were collected and incubated with beads conjugated to anti-FLAG M2 antibody (Merck). Beads were washed, and the proteins bound to them were eluted using FLAG peptide and analyzed using immunoblotting. Immunoblotting was performed as previously described (3, 33) using mouse anti-V5 antibody (R960-25, Thermo Fisher Scientific) and rabbit anti-FLAG polyclonal antibody (F7425, Merck).

Reporter assays were performed as previously described, with minor modifications (3, 34). A pGL4.35 reporter vector containing the firefly luciferase gene, under the control of a transcriptional regulatory unit comprising 9x Gal4-binding elements, was purchased from Promega. A pRL vector expressing Renilla luciferase (Promega) served as an internal control. 293FT cells were seeded into 24-well plates at a density of 2.5 × 10^4 cells/well and cotransfected with a reporter plasmid (100 ng), internal control vector (10 ng), pcDNA3 Gal4BD-HIF-1α CAD plasmid (3, 34) (50 ng), and other plasmids (200 ng) that expressed either vector alone, wild-type Mint3, or Mint3mut2. Transfection was performed using Lipofectamine 2000. Twenty-four hours after transfection, luciferase activity was measured using the Dual-Glo Luciferase Assay System (Promega), according to the manufacturer’s instructions. Luminescence was measured using a GloMax 20/20 luminometer (Promega).

Data availability

All data are available from the authors upon request. Please send request to Satoru Nagatoishi, ngtoishi@ims.u-tokyo.ac.jp.

Supporting information—This article contains supporting information.

Acknowledgments—We thank Thermo Fisher Scientific for the technical support in HDX-MS experiments.


Funding and additional information—This work was supported by JSPS KAKENHI under grant number JP18H02082, JP18H05425 (to S. N.) and JP16H02420, JP19H05766, JP20H02531 (to K. T.), by the
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Platform Project for Supporting Drug Discovery and Life Science Research [Basis for Supporting Innovative Drug Discovery and Life Science Research (BINDS)] from AMED of Japan under grant number JP20am0101094 (to K. T.), and by the P-CREATE (Project for Cancer Research and Therapeutic Evolution) from AMED of Japan under grant number JP21cm0106211 (to S. N., T. S., and K. T.).

Conflict of interest—The authors declare that they have no conflicts of interest with the contents of this article.

Abbreviations—The abbreviations used are: αKG, α-ketoglutaric acid; CD, circular dichroism; DSC, differential scanning calorimetry; DSF, differential scanning fluorimetry; FIH-1, factor inhibiting HIF-1; HDX-MS, hydrogen/deuterium-exchange–mass spectrometry; ITC, isothermal titration calorimetry; NMR, nuclear magnetic resonance; SEC, size-exclusion chromatography; SEC-MALS, SEC–multiance laser scattering.

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