

Mass spectrometry reveals modularity and a complete subunit interaction map of the eukaryotic translation factor eIF3

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The eukaryotic initiation factor 3 (eIF3) plays an important role in translation initiation, acting as a docking site for several eIFs that assemble on the 40S ribosomal subunit. Here, we use mass spectrometry to probe the subunit interactions within the human eIF3 complex. Our results show that the 13-subunit complex can be maintained intact in the gas phase, enabling us to establish unambiguously its stoichiometry and its overall subunit architecture via tandem mass spectrometry and solution disruption experiments. Dissociation takes place as a function of ionic strength to form three stable modules eIF3(c:d:e:l:k), eIF3(f:h:m), and eIF3(a:b:i:g). These modules are linked by interactions between subunits eIF3b:c and eIF3c:h. We confirmed our interaction map with the homologous yeast eIF3 complex that contains the five core subunits found in the human eIF3 and supplemented our data with results from immunoprecipitation. These results, together with the 27 subcomplexes identified with increasing ionic strength, enable us to define a comprehensive interaction map for this 800-kDa species. Our interaction map allows comparison of free eIF3 with that bound to the hepatitis C virus internal ribosome entry site (HCV-IRES) RNA. We also compare our eIF3 interaction map with related complexes, containing evolutionarily conserved protein domains, and reveal the location of subunits containing RNA recognition motifs proximal to the decoding center of the 40S subunit of the ribosome.

hepatitis C virus internal ribosome entry site | subunit organization
model3 | top-down analysis of macromolecular complexes |
translation regulation | in-solution disruption

Since its identification in the 1970s (1–3), the translation initiation factor eIF3 has been subjected to intense scrutiny. Despite considerable interest, knowledge of many aspects of its structure and function remain elusive because of its overall structural complexity and the lack of facile genetic approaches. It is established, however, that eIF3 is involved in both ribosome biogenesis and protein synthesis in eukaryotes (4). Concerted binding of initiation factors is required to initiate protein synthesis and recruit transfer and messenger RNAs to the 40S subunit before assembly of active ribosomes (5). eIF3 binding may take place initially during this process, together with eIF1 and eIF1A to the 40S, followed by binding of the Met-tRNA_i-eIF2-GTP complex. Then mRNA binding, scanning, and AUG recognition occur, enabling the 60S subunit to join to form elongation-competent 80S ribosomes (6). An alternative pathway of initiating protein synthesis, often used by viruses, involves a structured sequence in the 5' untranslated region of mRNA known as the internal ribosome entry site (IRES). These sequences promote translation initiation without requiring the full complement of eukaryotic initiation factors (5–7). The hepatitis C virus (HCV) IRES is recognized specifically by the small ribosomal subunit and eIF3 before viral translation initiation, forming stable complexes amenable to numerous structural studies (7, 8).

Much of the structural information on eIF3 is derived from electron microscopy images, where eIF3 is revealed as a five-lobed particle that binds to the 40S subunit of the ribosome on the side opposite the 60S subunit. HCV IRES RNA was found to bind diagonally across eIF3, and it also binds to the solvent side of the 40S subunit, inducing conformational changes in the 40S upon binding (7, 8). eIF4G binds to the same eIF3 “arm” as that occupied by part of the IRES, consistent with its role in mRNA recruitment to the 43S preinitiation complex. It is also established, from directed hydroxyl radical probing, that subunit eIF3j binds to the decoding center of the human 40S subunit (9).

Human eIF3 shares an intriguing degree of homology with two other complexes whose functions appear unrelated: the COP9 signalosome and the 19S proteasome lid. All three complexes consist of subunits with either PCI (Proteasome, COP9, eIF3) or MPN (Mpr1-Pad1 N-terminal) signature domains (10), and share a common 6PCI + 2MPN domain stoichiometry. The mammalian eIF3 also has an additional five non-PCI/MPN subunits. Two of these additional subunits (b and d) have been shown to cross-link with mRNA within both the 48S preinitiation complex (11) and the IRES-eIF3 binary complex. eIF3b, together with eIF3g that is also a non-PCI/MPN subunit, contain RNA recognition motifs (RRM), indicating their potential role in RNA binding. The roles of the MPN and PCI domains are not well understood, but current opinion is that these domains serve as a structural scaffold, interacting with each other and to other binding partners. It has also been proposed that MPN subunits have a regulatory role in coordinating overall activity (12).

Many of the interactions established in eIF3 are thought to be conserved between human and yeast (6) suggesting that they share a common core, although yeast eIF3 contains only a subset of the subunits identified in human. A comprehensive subunit interaction map for human eIF3 has not yet been reported, although models have been proposed for the yeast complex (6). We have shown that mass spectrometry (MS) can be used to uncover complete interaction maps for large multiprotein complexes, including the yeast exosome (13) and 19S proteasome lid (14). These interaction maps were achieved by generating large numbers of subcomplexes as “building blocks” of the entire complex.

Our previous MS studies of human eIF3 have shown that we can determine close to 30 phosphorylation sites, primarily on subunits

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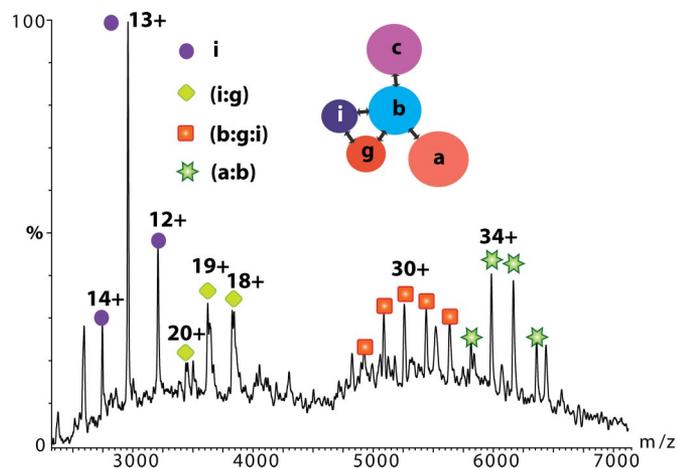


Fig. 3. MS spectrum of the yeast eIF3 isolated by tagging subunit eIF3b. Charge state series are assigned on the basis of masses to subcomplexes eIF3i:g, eIF3b:g:i, and eIF3a:b. eIF3i is observed dissociating from the yeast complex at $\approx m/z$ 3,000. (Inset) The interaction network for yeast eIF3 derived from seven subcomplexes observed by MS.

human eIF3 in our spectra suggests its resemblance to the yeast eIF3 core complex.

MS of Yeast eIF3 and IP Confirm Our Interaction Network. To provide the missing links within the subcomplex (a:b:c:i:g), we isolated eIF3 from yeast using a tandem affinity protocol (21). We were unable to record a spectrum of the intact complex despite confirmation from proteomics experiments that subunits a, b, c, g, and i, but not j, together with eIF5 were present in the complex (SI Text and Fig. S5). This implies that the complex is less stable under our conditions than that of its human counterpart. However, we were able to detect pairwise dimers eIF3g:i and eIF3a:b and the trimer eIF3b:g:i when the affinity tag is placed on subunit b (Fig. 3). By tagging subunit c, we observed additional subcomplexes assigned as eIF3(c:b:i), b:i, b:g, and (c:b:g) (data not shown). Together, MS data from yeast establish seven subcomplexes, enabling us to derive an interaction network for eIF3(a:b:c:i:g) with eIF3b serving as the scaffolding protein connecting the remaining subunits a,c,i and g (Fig. 3).

The results from our investigation of yeast eIF3 allow us to define analogous interactions within the human complex, which resembles closely one of the three relatively tightly associated modules generated by manipulating the ionic strength. These modules are denoted A(eIF3a:b:i:g), B(eIF3f:h:m), and C(eIF3k:l:e:d:c). The remaining subunit eIF3j, a labile subunit, attaches to the complex via eIF3b as previously defined (9, 22). The fact that all of the subcomplexes with masses <500 kDa are composed of members from only one or at most two of these modules supports our proposal of three tightly associated modules with weaker interactions occurring between them.

Having established interaction networks within these three modules, we then looked for connections between them. We noted that the effect of increasing ionic strength on the subcomplex (eIF3k:l:e:d:c:f:h:m) was formation of smaller subcomplexes l:k, e:l:k, and d:e:l:k while the intact (k:l:e:d:c:f:h:m) subcomplex persists, implying that it is not likely that the four proteins k, l, e, d interact strongly with the trimer f:h:m. Together with our earlier conclusion that the presence of subunit h is essential for binding the trimer f:h:m, it is reasonable to propose that c and h provide one of the electrostatic interactions between modules B and C. Similarly, establishment of the eIF3b:c interaction in yeast and the observation of subcomplex (a:b:c:i:g) strongly suggest the role of eIF3c in recruiting module C to A. It is, however, difficult to define unambiguously contacts between the module B and A based on MS

data alone. We therefore supplemented our data with results from IP. A number of binary interactions were found that support our subunit contacts for modules B and C (Fig. S3). Moreover, those between eIF3b:f and b:h are important for determining interactions between the modules A and B.

Taking this data together, we are able to propose a detailed interaction network for the stable interactions within the entire human eIF3 (Fig. 4). In this model, eIF3 subunits are represented as spheres scaled according to their mass. It was not possible to take into account subunits that may deviate from globular conformations because a complete atomic structure of only one subunit (eIF3k) has been reported to date (23). This simplistic interaction model is, however, consistent with the IP data, the interactions within yeast eIF3, and the 27 subcomplexes assigned for human eIF3 as a function of ionic strength.

Comparison of eIF3 with eIF3:IRES. If we now compare the mass spectra of free human eIF3 with that bound to HCV IRES, we observe an increase in m/z value consistent with the association of a 332-nucleotide RNA of 107 kDa with eIF3 (Fig. S4). MS/MS of the IRES:eIF3 peak isolated at m/z 13,800 yielded subunit k (Fig. S4c), indicating that gas-phase dissociation of eIF3k is not inhibited by IRES binding. Moreover, these high energy conditions give rise to marginally resolved charge states enabling the mass of the IRES:eIF3 complex to be measured as 907 kDa, confirming binary complex formation. If we compare the activated spectra of the intact eIF3 complex with those recorded for the complex bound to IRES, we note several differences: first, the disappearance of peaks corresponding to relatively large subcomplexes in the IRES:eIF3 spectrum, indicating that when the IRES is bound, subcomplex formation is diminished, implying a global stabilization. Particularly interesting is the observation that eIF3i and the l:k dimer can still dissociate in solution from the binary complex, suggesting that these three subunits are not in direct contact with the IRES. By contrast, two dimers, i:g and f:h, are not present in the IRES:eIF3 spectrum, suggesting that the subunits g, f, and h play a role in IRES binding (Fig. 5a).

Discussion

Here, we report a model of the subunit organization of human eIF3, the largest and most complex of the eukaryotic initiation factors. Our model, is based primarily on data from MS of intact subcomplexes generated in solution and is supported by a subset of interactions generated independently by IP and our analysis of the homologous yeast complex. Our results reveal facile dissociation of subunits eIF3i, j, k, and m as well as three well defined interaction modules generated by manipulating the ionic strength of the complex-containing solution. Furthermore, results from binding of human eIF3 to the IRES shows that although dissociation of larger subcomplexes is prevented, eIF3i and the eIF3l:k dimer are still able to dissociate, suggesting that these subunits are not directly involved in the interactions with the IRES.

If we compare our model with data for other protein complexes containing both MPN and PCI domains (10), we note that the two MPN domain-containing subunits, eIF3f and h, interact directly in our subunit organization map (Fig. 5a), a phenomenon also observed with another PCI-containing complex, the proteasome lid (14). Moreover, several directly interacting PCI domains are found in both complexes, supporting their role in providing a scaffold for active subunits. Our interaction map for human eIF3 also establishes direct interactions among eIF3b, g, i, and j, four of the five non-PCI/MPN subunits absent in the other two PCI complexes.

The high sensitivity of eIF3 to dissociation into subcomplexes as the ionic strength is raised suggests that electrostatic interactions may play a central role in maintaining the integrity of the complex and in regulating its function. The three largest subunits eIF3a, b, and c are extensively phosphorylated, occupy a central position in our model, and are conserved between human and yeast (16).

This is an intriguing observation because it implies a role of RNA recognition for these two subunits (eIF3b and g) in binding and presenting mRNA to the decoding centre of the ribosome.

The labile binding of eIF3j, as shown by its absence from all of our subcomplexes, implies its ready dissociation from the complex. Interestingly, eIF3j has been shown to dissociate from eIF3 in quiescent T lymphocytes, but upon activation, eIF3j binds eIF3 and 40S ribosomal subunits, implying that the interaction of subunit j with eIF3 may well be regulated (26). Moreover, eIF3j has been shown to have more than one role, not only in the decoding center (9), but also taking part in 20S pre-rRNA processing (4). The fact that this critical subunit is only loosely associated and is located close to eIF3b:g:i and the observation that subunit eIF3i is also labile implies that dissociation of eIF3j, with or without its neighboring subunits, may well be associated with its regulatory role in translation initiation and in ribosome biogenesis (4).

Overall, in this study, we have derived a subunit interaction map for a 13-subunit complex using an emerging MS approach (13, 14, 27). If we consider our interaction map with the five-lobed architecture revealed by cryo-EM (8), it is tempting to speculate that the subcomplexes frequently observed in our MS experiments may correspond to the lobes identified within the EM structure. The assignment of these subcomplexes to the different domains of the 3D EM structure remains to be completed. Overall, the subunit architecture presented here, together with knowledge of the ease of dissociation and interactions that link the three stable modules provide an intriguing insight into the relationships between subunits and their possible functional significance in this complex initiation factor.

Experimental Methods

The eIF3 sample was purified from HeLa cell cytosol as described (8, 15).

LC-MS Analysis of eIF3 Subunits. Capillary HPLC-MS analysis of individual eIF3 subunits was carried out by using a LC-Packings Ultimate System (Dionex)

equipped with a capillary UV detector set at 214 and 280 nm. eIF3 was prepared in a 1:1 (vol/vol) of 0.1% TFA and 1 μ l of sample applied to a capillary PS-DVB reverse-phase monolithic column (200 μ m i.d. \times 5 cm; Dionex) equilibrated at 90% solvent A (0.05% TFA) and 10% solvent B (0.04% TFA, 90% ACN). A linear gradient was used of 10–70% solvent B in 25 min at a flow rate of 3 μ l/min. The column effluent was analyzed by both electrospray and MALDI MS (see *SI Text*).

MS of Intact eIF3 and Subcomplexes. For MS of the intact complex, 20 μ l of purified eIF3 (1 μ g/ μ l) was buffer exchanged into AmAc buffer (pH 7.5) containing 1 mM DTT by using micro biospin 6 columns (Bio-Rad). 2- μ l aliquots were electrosprayed from gold-coated borosilicate capillaries prepared in house. Spectra were recorded on a QSTAR XL (Applied Biosystems) modified for high mass detection (28) and adjusted to preserve noncovalent interactions (29). MS experiments were performed at a capillary voltage of 1,200 V and declustering potentials of 40 V and 15 V. In tandem MS experiments, ions were isolated in the quadrupole and subjected to collision-induced dissociation (acceleration energy up to 170 V). To generate subcomplexes, a 7.5 M AmAc solution (Sigma) was diluted stepwise and added to the eIF3 solution at a ratio of 1:10 (vol/vol) to give the required AmAc concentration and incubated on ice for 10 min before analysis.

HCV-IRES Binding to eIF3. Wild-type HCV-IRES was transcribed *in vitro* and gel-purified (30). eIF3 was incubated with HCV-IRES at a molar ratio of 1:1 for 3 min at 37°C in 20 mM Tris (pH 7.5), 100 mM KCl, 3 mM Mg(OAc)₂, and 1 mM DTT. A control solution of eIF3 alone in the same buffer was prepared. Both the control and the experiment were then buffer-exchanged into 150 mM AmAc (pH 7.5) containing 3 mM Mg²⁺ acetate and 1 mM DTT.

Expression, Purification, and MS of Yeast eIF3. *S. cerevisiae* eIF3 was purified from TAP-tag yeast strains (Open Biosystems), modified to express endogenous proteins fused with a C-terminal tandem affinity tag on eIF3b or c (21). See *SI Text* and *Fig. S5*. Spectra were recorded on the QSTAR XL at a capillary voltage of 1,200 V and declustering potentials of 150 V and 15 V.

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