

SUPPLEMENTAL MATERIAL

FIGURE S1. Different reconstituted complexes of eIF3. (A) Tif34-Tif35 complex (third lane) formed out of individual proteins (first and second lane). (B) Prt1^{181C}-Tif34-Tif35 complex. (C) Tif32-Nip1 complex. (D) Tif32-Nip1-Prt1 complex. (E) Tif32-Nip1-Prt1-Tif34-Tif35 (full eIF3, first lane) and Nip1-Prt1-Tif34-Tif35 (second lane) complexes. (F) Tif32¹⁻⁴⁹⁴-Nip1-Prt1-Tif34-Tif35 complex.

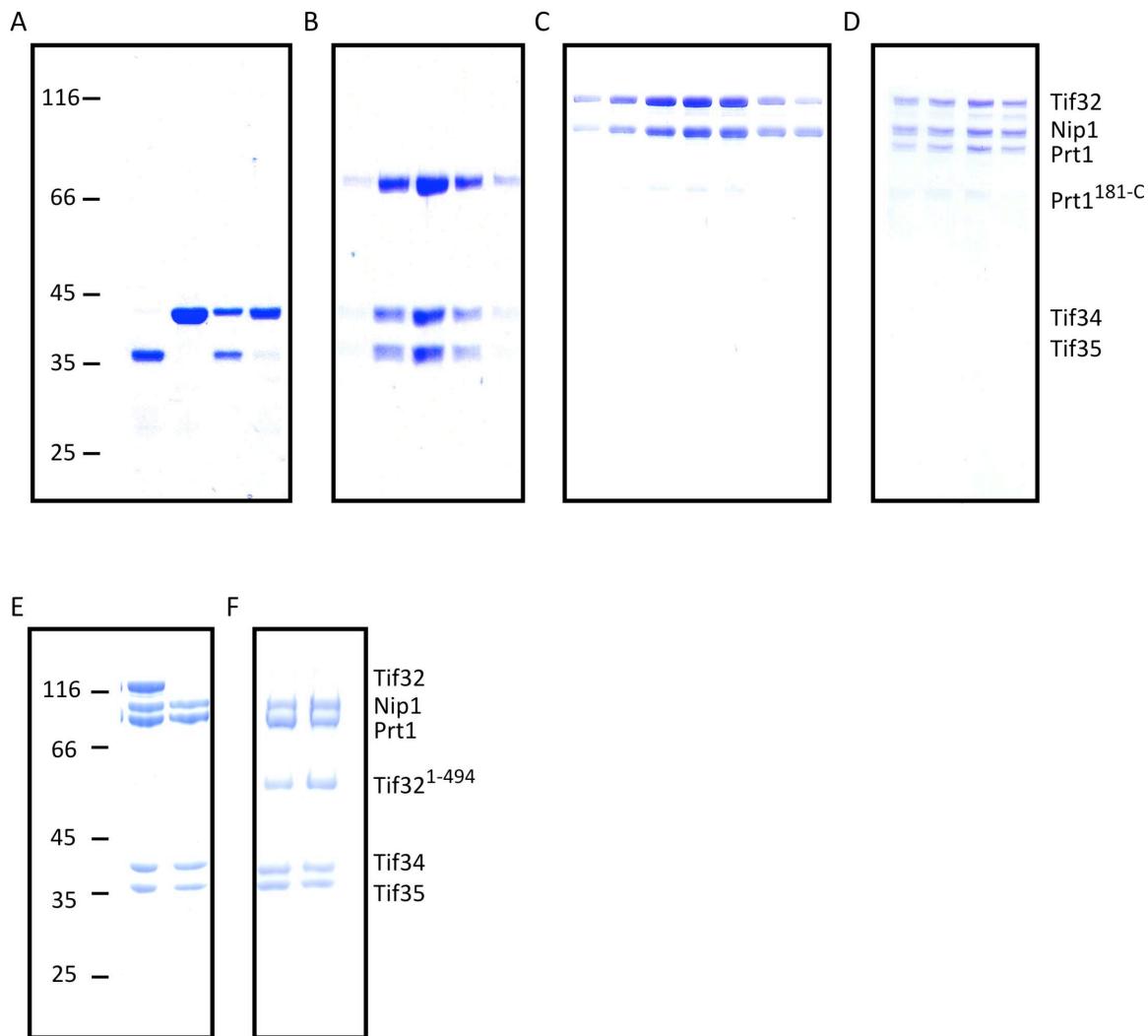


TABLE S1. List of the oligo-primers used in this work. In each case, the forward primers used for cloning of a certain protein or fragment are mentioned first (denoted F) followed by the corresponding reverse primers (denoted R).

Tif32 ^{FL}	F-BamHI	5' C CGG GAT CCA <u>ATG</u> GCC CCC CCA CCA TTC 3'
	R-Xhol	5' CCG CTC GAG <u>TTA</u> CCT GCC CCC CTT <u>GGC</u> 3'
Tif32 ¹⁻⁴⁹⁴	R-Xhol	5' CCG CTC GAG <u>TTA</u> ATC TTT AGC GAA TGT AAC CTT AGC 3'
Tif32 ⁵⁴⁴⁻⁹¹⁹	F-BamHI	5' CCG GGA TCA <u>GAT</u> CCT GTT ATT ATC CGC AAT TCT TA 3'
Tif32 ⁵⁴⁴⁻⁹¹⁹	R-Xhol	5' CCG CTC GAG <u>TTA</u> ACC GGC TAG TCT CTG TTC <u>G</u> 3'
Nip1 ^{FL}	F-Smal	5' T CCC CCG GGT <u>ATG</u> TCC CGT TTC TTT TCG TCT AAT <u>TAC</u> 3'
Nip1 ^{240-C}	F-BamHI	5' CCG GGA TCC <u>ATT</u> TCT TCG TCT CAA GGC AAT <u>G</u> 3'
	R-Xhol	5' CCG CTC GAG <u>TCA</u> ACG ACG ATT TGA TGG TGG GTT <u>AAG</u> .3'
Prt1	F-Ndel	5' GG GAA TTC CAT <u>ATG</u> AAA AAT TTT CTT CCA CGC ACA TTG AAA <u>A</u> 3'
Prt1 ^{181-C}	F-Ndel	5' GG GAA TTC CAT <u>ATG</u> CCT ACA TTC GTT CCA TCT AGT 3'
	R-Xhol	5' CCG CTC GAG <u>TTC</u> GAC CTT TTC CTT TGT TTC <u>TTC</u> 3
Tif34	F-Ndel	5' GG GAA TTC CAT <u>ATG</u> AAG GCT ATC AAA TTA ACA GGT <u>CAT</u> G 3'
	R-Xhol	5' CCG CTC GAG <u>TTA</u> ATT AGC TTC TTG CAT GTG CTC TTT <u>A</u> 3'
Tif35	F-Smal	5' T CCC CCG GGT <u>ATG</u> AGT GAA GTT GCA CCA GAG 3'
	R-Xhol	5' CCG CTC GAG <u>CTA</u> TTC CTT AAC CTT AGG TTT GGA <u>C</u> 3'
Hcr1	F-BamHI	5' CCG GGA TCC <u>ATG</u> TCT TGG GAC GAC GAA <u>G</u> 3'
	R-Xhol	5' CCG CTC GAG <u>TTA</u> CAT AAA GTC GTC ATC ACC GAA 3'

TABLE S2. The list of expression vector and strains used in this study. This table lists the optimal combination of the vector, position and type of the tag, as well as the expression strain which was used to obtain the highest amount of soluble protein.

Protein	vector	tag	Expression strain
Tif32	pET28b	N-His	Rosetta2 (DE3)
Tif32 ¹⁻⁴⁹⁴	pET15b	N-His	Rosetta2 (DE3)
Tif32 ⁴⁹⁴⁻⁹¹⁹	pGEX-6P-1	N-GST	Rosetta2 (DE3)
Tif32 ⁵⁴⁴⁻⁹¹⁹	pGEX-6P-1	N-GST	Rosetta2 (DE3)
Nip1	pGEX-6P-1	N-GST	Bl21 (DE3)
Nip1 ^{240C}	pGEX-6P-1	N-GST	Rosetta2 (DE3)
Prt1	pET22b	C-His	Rosetta2 (DE3)
Prt1 ^{181-C}	pET15b	N-His	Rosetta2 (DE3)
Tif34	pET15b	N-His	Bl21 (DE3)
Tif35	pGEX-6P-1	N-GST	Rosetta2 (DE3)
Hcr1	pGEX-6P-1	N-GST	Rosetta2 (DE3)