SUPPLEMENTARY DATA

Figure S1. Secondary structure prediction done with Network Protein Sequence Analysis Server (43);

A: secondary structure prediction for GluRS: residues 181-205; For the amino acids 181-200 there is a random coil region predicted and for 201-205 an extended strand; B: secondary structure prediction for MetRS: residues 141-210; Amino acids 160-172 are a random coil region, 173-175 an extended strand and 176-195 are again a random coil region.

А

	150	160	170	180	190	200	210
	I	1	1		1	I	1
UNK 208330	KVDVNVSRWYTLLE	MDPIFGEAH	DFLSKSLLELH	KKSANVGKKKE	THKANFEIDI	LPDAKMGEVVI	RFPPEP
DSC	ccccccc hh ccccc	ccc hhhhhh l	hhhhhhhhhh	hhhccceee			eccecc
MLRC	ccccchhhhhhhh	ccchhhhhhl	hhhhhhhhh	hccccccc			eccccc
PHD	ccchhhhhhheee					cccccceeee	eccccc
Predator	haddddddeeee	secceccehl	hhhhhhhhhh	hhhcccccch	hhhhhccco	cccccceeeee	eecccc
Sec.Cons.	ccccc??hhheee	ccc??h??hl	hhhhhhhhhh	hh??cccccc		cccccceeee	eccccc

В

	150	160	170	180	190	200	210
		I	I.	I	I	I	1
UNK_228150	KFPELPSKVHNAVA	LAKKHVPRD	SSSFKNIGAV	KIQADLTVKPH	KD SEILPKPNI	SRNILITSALE	YVNNVP
DSC	ccccchhhhhhhh	hhhhccccc	cchhhhhhhe	eeehhhhccco		cceeeeecccc	cccccc
MLRC	chchhhhhhhhhhh	hhhcccccc		eecccccccc		ceeeeeecco	cccccc
PHD	hachhhhhhhhhad	cccceeecc		ecccccccc	cecececee	cceeeeecccc	cccccc
Predator	cccccccchhhhhh	hhhhccccc		c hhhhhh ccco		ceeeeeecco	cccccc
Sec.Cons.	ccc??hhhhhhhhh	hhh?ccccc		eec????cccc		cceeeee?ccc	cccccc

Figure S2. Guinier plots of experimental scattering data;

1: Pentameric complex MetRS:GluRS:Arc1p:tRNAs (grey); 2: Trimeric complex of MetRS:GluRS:Arc1p (turquoise); 3: GluRS:Arc1p (lilac); 4: MetRS:Arc1p complex (yellow); 5: GluRS (red); 6: MetRS (blue); 7: Arc1p (green).

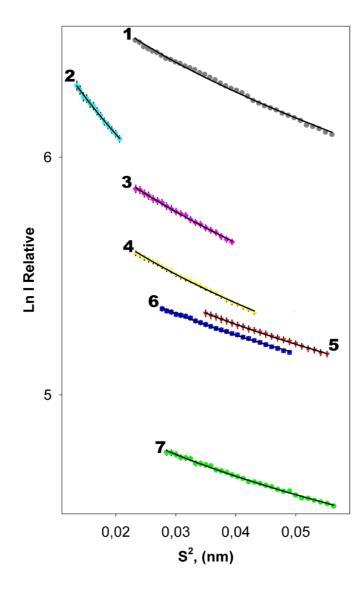
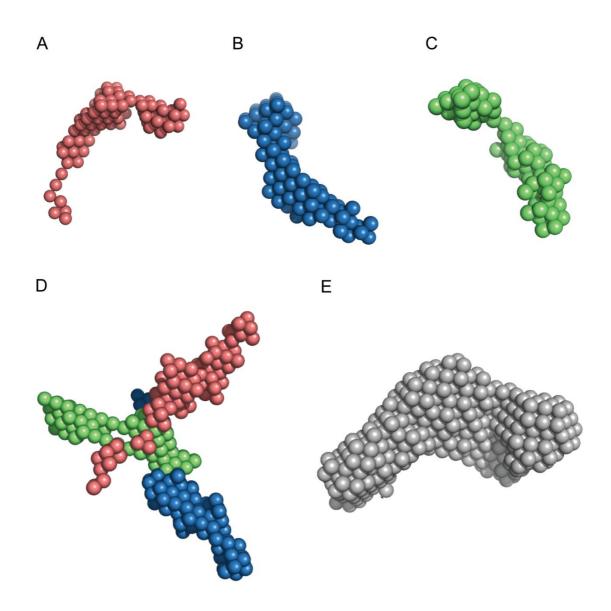


Figure S3. Ab initio Models;A: GluRS (red), B: MetRS (blue), C: Arc1p (green), D: trimeric complex andE:pentameric complexThe models were obtained by analyzing the SAXS data with the Program DAMMIN. Proteins are shown with densely packed beads.



SUPPLEMENTARY METHODS

EMSA assay

To analyze the binding of the tRNAs to the trimeric complex, the trimeric complex (MAG) was used in a 10 μ M concentration (1 μ I) in a total reaction volume of 20 μ I. Each unlabeled tRNA was applied in a 1:1 molar ratio to the complex as the highest concentration (1.5 μ I). The ³²P labeled tRNAs were added in the same concentration steps (0.5-1.5 μ I) as the unlabeled ones (Figure 3). The reactions were incubated on ice for 30 minutes and then loaded on a 6% native gel.

SUPPLEMENTARY REFERENCES

- 43. Combet,C., Blanchet,C., Geourjon,C. and Deléage,G. (2000) NPS@: network protein sequence analysis. *Trends Biochem Sci.*, **25**, 147-50.
- 44. Simader,H. and Suck,D. (2006) Expression, purification, crystallization and preliminary phasing of the heteromerization domain of the tRNA-export and aminoacylation cofactor Arc1p from yeast. *Acta. Crystallograph Sect. F: Struct. Biol. Cryst. Commun.*, **62**, 346-349.