GTPases IF2 and EF-G bind GDP and the SRL RNA in a mutually exclusive manner

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Translational GTPases (trGTPases) are involved in all four stages of protein biosynthesis: initiation, elongation, termination and ribosome recycling. The trGTPases Initiation Factor 2 (IF2) and Elongation Factor G (EF-G) respectively orchestrate initiation complex formation and translocation of the peptidyl-tRNA:mRNA complex through the bacterial ribosome. The ribosome regulates the GTPase cycle and efficiently discriminates between the GDP- and GTP-bound forms of these proteins. Using Isothermal Titration Calorimetry, we have investigated interactions of IF2 and EF-G with the sarcin-ricin loop of the 23S rRNA, a crucial element of the GTPase-associated center of the ribosome. We show that binding of IF2 and EF-G to a 27 nucleotide RNA fragment mimicking the sarcin-ricin loop is mutually exclusive with that of GDP, but not of GTP, providing a mechanism for destabilization of the ribosome-bound GDP forms of translational GTPases.

In all domains of life protein biosynthesis on the ribosome follows the same functional cycle (for review see1). First, the ribosome associates with the mRNA and locates the start codon, which is recognized by initiator tRNA. Assembly of the initiation complex is regulated by several initiation factors; bacterial initiation factor 2 (IF2) and its eukaryotic and archaeal homologues eIF5B and aIF5B, respectively, are evolutionary conserved GTPases involved in ribosomal subunit joining and, in the case of IF2, additionally in initiator tRNA selection2. Upon formation of the initiation complex, the ribosome starts its elongation cycle: the A-site codon is decoded by the aminoacyl-tRNA delivered in complex with a GTPase protein - EF-Tu in bacteria, called eEF1A in eukaryotes - and eEF1A in archaea. Next, another GTPase - EF-G in bacteria, eEF2 in eukaryotes and aEF2 in archaea - catalyzes ribosomal translocation along the mRNA for one codon4,5. The elongation cycle repeats until the ribosome reaches the stop codon, which is decoded by a specialized release factor (RF1/RF2 in bacteria) that cleaves off the produced protein6. EF-G also has an additional role in ribosome recycling in bacteria, acting in concert with the ribosome recycling factor RRF to disassemble the post-termination complex7.

In vitro biochemical investigations have demonstrated that IF2 is switched to a functionally activated form by interactions with initiator tRNA (fMet-tRNAi), GTP, and – to much lesser extent – GDP8. At physiologically relevant temperatures off the ribosome, both bacterial IF2 and eukaryotic eIF5B have somewhat lower affinity to GTP than to GDP9,10, and binding of G nucleotides is largely insensitive to IF2 binding to initiator tRNA11,12. IF2 interacts with the 30S ribosomal subunit via two binding sites, i.e. the N-terminus and domain II13,14, and in the context of the initiation complex, the latter interaction is strongly stimulated by GTP and fMet-tRNAi14.

Ribosomal translocation has been extensively studied in the bacterial system, and relationships among the EF-G GTPase function and distinct stages of translocation are well understood (for review see5). Since in translocation GTP cannot be substituted for monoQ-purified GDP lacking trace amounts of GTP15–18, it is a prerequisite for successful translocation that while engaging the ribosome, either EF-G is in the GTP-bound form, or GDP is exchanged for GTP while EF-G is bound to the ribosome. Over the years, EF-G affinities to GTP and GDP have been measured both off and on the ribosome in several laboratories11,19–23. According to the detailed balance principle25, there should also be a dramatic increase of EF-G:GTP affinity to the ribosome, compared to apo-EF-G. Such an increase in affinity has been confirmed experimentally23,26.

The effects of the ribosome on EF-G's interaction with GDP are not so well understood. Estimates of the dissociation rate constants (k-1) demonstrated that GDP dissociation is slowed down on the ribosome about...
ten-fold, which was interpreted as a stabilizing effect of the 70S on the EF-G-GDP complex. However, in the absence of the association rate constants \( k_{+1} \), a decrease in the dissociation rate does not necessarily mean that affinity increases, since \( K_d \) is a product of the division of the two rate constants \( K_d = k_{-1} / k_{+1} \), given that the interaction has a simple one step binding mechanism. Moreover, \( k_{+1} \) estimates for GDP dissociation from the 70S:EF-G:GDP complex vary at least ten-fold between different reports, complicating the situation even further. This necessitates further investigations of the relationship between the ribosome and GDP binding to EF-G.

A powerful method for investigating thermodynamics of molecular interactions is Isothermal Titration Calorimetry (ITC) (for a review see), which we have successfully applied in the study of complex formation between GTPases and their ligands, such as nucleotides, RNAs and proteins. However, studying interactions with the ribosome presents a challenge for this approach due to technical reasons, such as molecular aggregation. So far, only experiments with ribosomal subunits have been successful. One possible way of overcoming this limitation is to use isolated ribosomal components that mimic different functional centers—an approach that has been successfully applied to various biochemical and structural investigations of the translational apparatus.

While these partial systems necessarily overlook most of the complexity of the in vivo system, they have the advantage of allowing us to decouple one specific facet of the interactions from the others.

The sarcin-rinic loop (SRL), a part of the ribosomal RNA that is targeted by sarcin and ricin toxins (for review see), is one of the most well-established oligonucleotide mimics. In the native ribosomal complex, it directly interacts with EF-G and is crucial for activation of GTP hydrolysis, though recent data suggest that is acts merely as an anchoring point for the EF-G on the ribosome and is not involved directly in the GTPase activation. Similarly, the SRL was documented as crucial element for IF2 function on the ribosome: the interaction was first shown using chemical probing, and later validated by several cryo-electron microscopy studies. Using a gel-retardation assay, a 27 nucleotide-long SRL RNA fragment was shown to bind to EF-G. Surprisingly, SRL-EF-G complex formation is strongly inhibited by the presence of GDP and is insensitive to GDPNP, a non-hydrolysable GTP analogue. Differential effects of GDPNP and GDP, together with mutagenesis data suggest that the model reaction described by Munishkin and Wool reflects a relevant partial reaction rather than non-specific interaction between EF-G and the RNA oligonucleotide. Several questions, however, remain unanswered in the original report. First, GDPNP, even though it is a widely used mimic of GTP, does not have exactly the same properties and often has considerably lower affinity, therefore sometimes failing to represent the effects of GTP faithfully. Since the SRL RNA oligonucleotide does not induce GTP hydrolysis on EF-G, it is principally possible to perform ITC experiments with GTP rather than GDPNP. Second, by the detailed balance argument, inhibition of EF-G-SRL complex formation by GDP suggests that SRL binding to EF-G should result in a strongly decreased affinity to GDP. This prediction has never been tested experimentally. Third, cross-talk between binding of G nucleotides and the SRL to other translational GTPases such as IF2 has never been investigated.

**Results**

In the absence of nucleotides, EF-G associates with the SRL with a \( K_d \) of 5.9 \( \mu \)M at 5°C and 4.2 \( \mu \)M at 25°C (Figure 1A, Table 1), which is in satisfactory agreement with the \( ~7 \) \( \mu \)M \( K_d \) reported by Munishkin and Wool. The complex formation has stoichiometry close to unity, indicating its specificity. The interaction is entropy driven (\( \Delta H = 4.94 \text{ kcal/mol} \) vs \( T \Delta S = 11.57 \text{ kcal/mol} \), suggesting a hydrophobic nature. The addition of GTP at 500 \( \mu \)M had no significant effect on the interaction between EF-G and the SRL (\( K_d = 3.7 \) vs 4.2 \( \mu \)M, Table 1). However, in the presence of 500 \( \mu \)M GDP, no complex formation was detected, again in good agreement with the results of Munishkin and Wool.

After validating the inhibitory effect of GDP on SRL-EF-G complex formation, we tested the effects of the SRL on the binding of GDP and GTP to EF-G. In the case of GTP, the presence of 60 \( \mu \)M SRL had an insignificant effect on the interaction between EF-G and the nucleotide (\( K_d = 2.4 \) vs 2.7 \( \mu \)M, Table 1). Binding of GDP, however, was considerably inhibited by the SRL, with apparent \( K_d \) increasing four times in the presence of 60 \( \mu \)M SRL (Table 1). The addition of 200 \( \mu \)M SRL to EF-G resulted in almost complete inhibition of GDP binding (Figure 1B), demonstrating the competition directly. To validate the specificity of the SRL-EF-G interaction, we employed an SRL G2655U mutant that displays dramatically weaker binding to EF-G as judged by gel-shift analysis. Using 25 \( \mu \)M EF-G in the cell and 250 \( \mu \)M SRL G2655U in the syringe, we failed to detect binding at 4°C and 25°C (data not shown). This indicates that the mutation indeed severely affects the Gibbs free energy and/or enthalpy of the interaction, in line with the results of Munishkin and Wool.

Next, we investigated SRL interactions with IF2 (Table 2). In the absence of nucleotides, IF2 and the SRL form a considerably tighter complex than the SRL and EF-G (\( K_d = 0.67 \) \( \mu \)M vs 4.2 \( \mu \)M at 25°C). This interaction was, as with EF-G, mostly entropy-driven (\( AH = -2.27 \text{ kcal/mol} \) vs \( T \Delta S = 6.14 \text{ kcal/mol} \). Complex formation was largely insensitive to the GTP nucleotide (\( K_d = 1 \) \( \mu \)M in the presence of 500 \( \mu \)M GTP), and, again, just as in the case of EF-G, addition of 500 \( \mu \)M GDP inhibited the interaction between IF2 and SRL completely, both at 5°C and 25°C. Binding of GDP in the presence of 200 \( \mu \)M SRL was not detected, again indicating strong mutual competition between GDP and SRL binding.

In our recent ITC investigations of IF2 complex formation with fMet-tRNA, we observed no significant inhibitory effect of GDP, indicating the SRL:IF2 interaction is distinct from other IF2:RNA interactions. To further investigate the specificity of the SRL:IF2 interaction, we analyzed IF2 binding to the SRL G2655U mutant. The G2655U mutation resulted in a modest two-fold increase in the \( K_d \) of IF2:SRL complex formation, but partitioning of the Gibbs free energy between the enthalpic and entropic components was dramatically altered (\( AH = -6.0 \text{ kcal/mol} \) and \( T \Delta S = 2.03 \text{ kcal/mol} \) for wt SRL vs \( AH = -6.0 \text{ kcal/mol} \) and \( T \Delta S = 6.14 \text{ kcal/mol} \) for the G2655U mutant). This effect, known as enthalpy-entropy compensation, is a hallmark of biological molecular interactions and is observed when a system is perturbed by temperature or ionic strength changes, mutations, binding of allosteric ligands etc.

Another ribosomal element that interacts with translational GTPases is ribosomal protein L7/L12, which on the ribosome interacts with the G domain of EF-G via its CTD domain. The interaction between isolated L7/L12 and the apo-form of EF-G has been shown to be extremely weak (\( K_d = 0.4 \) ± 0.1 \( \mu \)M), but it has been suggested that it could be affected by G nucleotides, suggesting a possibility that interaction with L7/L12 should be responsible for the dramatic increase in EF-G affinity to GTP. Therefore we have analyzed L7/L12 binding to EF-G in a wide range of temperatures at increasing concentrations of the interacting partners. Even when EF-G at 400 \( \mu \)M was titrated with 4.7 \( \mu \)M L7/L12, the interaction signal was mainly dominated by the dilution heat (Supplementary Text S1: Supplementary Figure S1), and was not detectably stimulated by addition of 1 \( \mu \)M GTP and GDP (data not shown).

**Discussion**

In this report we provide an in-depth thermodynamic analysis of the interaction of bacterial translational GTPases EF-G and IF2 with the SRL RNA element. Recent investigations suggest a possibility that the SRL acts as an anchoring point for GTPase binding to the ribosome and is not implicated in GTPase activation per se. Our complementary quantitative analysis demonstrates that binding of
GDP and the SRL to GTPases IF2 and EF-G are mutually exclusive, while the thermodynamic profiles of complex formation between apo- and GTP-bound GTPases and the SRL are almost indistinguishable (Tables 1 and 2, Figure 2). Taken together, this suggests that apo- and GTP-bound forms of both EF-G and IF2 are efficiently discriminated by the SRL to the exclusion of the GDP-bound forms. The GDP and the SRL to GTPases IF2 and EF-G are mutually exclusive, while the thermodynamic profiles of complex formation between apo- and GTP-bound GTPases and the SRL are almost indistinguishable (Tables 1 and 2, Figure 2). Taken together, this suggests that apo- and GTP-bound forms of both EF-G and IF2 are efficiently discriminated by the SRL to the exclusion of the GDP-bound forms. The

Table 1 | Thermodynamic parameters of SRL, GDP and GTP binding to EF-G determined by isothermal titration calorimetry

| Sample          | Ligand | T, °C | $K_a$, M$^{-1}$ | $K_d$, μM | $\Delta H$, kcal/mol | $\Delta S$, kcal/mol | $\Delta G$, kcal/mol |
|-----------------|--------|-------|----------------|-----------|---------------------|---------------------|---------------------|
| EF-G            | SRL    | 5     | $1.7 \times 10^5$ | 5.9       | 4.94                | 11.57               | -6.63               |
|                 | SRL    | 25    | $2.4 \times 10^5$ | 4.2       | 1.75                | 9.09                 | -7.34               |
| GDP             | 4      | 6.0 x 10$^5$ | 1.7 | -8.29 | -0.97 | -7.32 |
|                 | 25     | $5.6 \times 10^5$ | 1.8 | -8.58 | -0.74 | -7.84 |
| GTP             | 5      | $1.2 \times 10^4$ | 8.3 | 1.73 | 8.17 | -6.44 |
|                 | 25     | $3.7 \times 10^4$ | 2.7 | -1.84 | 5.75 | -7.59 |
|                 | GTP    | 5      | $1.8 \times 10^4$ | 5.6       | 1.65                | 8.31                 | -6.66               |
|                 | GDP    | 25     | $4.2 \times 10^4$ | 2.4       | -1.67               | 6.00                 | -7.67               |
|                 | ADI    |        |                |           |                    |                     |                     |
| EF-G:SRL$_{500}$ | 5      | 3.7 x 10$^3$ | 6.7 | -4.84 | 2.22 | -7.06 |
|                 | GDP    | 25     | ADI            | ADI       |                    |                     |                     |
| EFG-GDP$_{500}$ | SRL    | 25     | ADI            | ADI       |                    |                     |                     |
| EFG-GTP$_{500}$ | SRL    | 25     | 2.7 x 10$^4$ | 3.7       | 1.72                | 9.13                 | -7.41               |

*All measurements were performed three to five times in phosphate buffer (5 mM K$_2$HPO$_4$, 10% glycerol, 95 mM KCl, 1 mM DTT and 5 mM MgCl$_2$, pH 7.5).

$K_a$ – affinity constant; standard deviation did not exceed ±20%.

$K_d$ – dissociation constant; calculated as 1/$K_a$.

$\Delta H$ – enthalpy variation; standard deviation did not exceed ±10%.

$\Delta S$ – entropy variation; calculated from the equation $\Delta G = \Delta H - T \Delta S$.

$\Delta G$ – Gibbs energy; calculated from the equation $\Delta G = -RT \ln K_a$.

The lowercase number indicates the concentration in μM. Concentration of EF-G was kept constant at 24 μM.

ADI – indicates Absence of Detectible Interaction rather than absence of measurements.
active role of GDP in regulation of the interaction of EF-G and IF2 with the SRL provides a mechanism for selective destabilization of GDP-bound forms of translational GTPases on the ribosome after GTP hydrolysis.

Comparison of cryo-EM reconstructions of ribosome-bound IF2-GDP and IF2:GDPNP shows that IF2 has extensive contacts with the SRL only in the GDPNP-bound complex. This observation is in agreement with our thermodynamic data demonstrating an inhibitory effect of GDP on the IF2:SRL interaction. The relatively low resolution of this cryo-EM reconstruction does not allow us to discern the molecular details of the IF2:SRL interface. However, the crystal structure of EF-G blocked on the ribosome in complex with GDP and the magnesium ion is well documented. Recent investigations of bacterial initiation factor IF2 showed profound structural rearrangements in the GDP-bound as compared to the apo-state, in agreement with complementary biochemical experiments demonstrating the functional differences between the two. However, in the case of EF-G, an absence of significant structural rearrangements in EF-G in the presence of G-nucleotides was suggested on the basis of Small Angle X-ray Scattering (SAXS) and analysis of the changes in the Solvent Accessible Area (SAA) inferred from the ITC data. However, it should be noted that both of these methods are intrinsically incapable of detecting structural rearrangements that do not lead to changes in the radius of gyration (SAXS) or SAA (ITC). In conclusion, we suggest that GDP is likely to exert its effects on SRL binding via alteration of conformational entropy by affecting protein motions rather than structure. Direct validation of this prediction requires high-end NMR investigations focused on protein side chain dynamics.

Further investigations are required in order to put the SRL effect into a kinetic framework of sequential interactions of IF2 and EF-G GTPases with the ribosome during their functional cycle. In the case of IF2, GDP was documented to have a weak activating effect on subunit joining, which is likely connected to GDP-mediated rearrangements. Time-resolved FRET studies of IF2 contacts with various components of the initiation complex – SRL, L7/12, fMet-tRNA - are necessary in order to uncover the functional role of the SRL and GDP competition for IF2 binding.

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**Table 2 | Thermodynamic parameters of SRL, GDP and GTP binding to IF2 determined by isothermal titration calorimetry**

| Sample       | Ligand   | T, °C | $K^b$, M$^{-1}$ | $K^a$, μM | $\Delta H$, kcal/mol | $\Delta S$, kcal/mol | $\Delta V$, kcal/mol |
|--------------|----------|-------|----------------|-----------|----------------------|----------------------|----------------------|
| IF2          | SRL      | 25    | $1.5 \times 10^4$ | 0.67      | $-2.27$              | $-3.42$              | $-7.88$              |
| IF2          | GDP      | 25    | $6.1 \times 10^4$ | 6.7       | $-18.72$             | $-11.67$             | $-7.05$              |
| IF2          | GDP      | 25    | $1.0 \times 10^4$ | 1.0       | $-2.24$              | $5.96$               | $-8.20$              |
| IF2          | GTTP$_{500}$ | 25    | $1.7 \times 10^4$ | 5.9       | $-17.11$             | $-9.98$              | $-7.13$              |
| IF2          | SRL      | 5     | ADI            | ADI       | ADI                  | ADI                  | ADI                  |
| IF2:GDP$_{500}$ | SRL      | 25    | $7.3 \times 10^4$ | 1.4       | $-6.00$              | $2.03$               | $-8.03$              |
| IF2:GTP$_{500}$ | SRL      | 25    | $5.7 \times 10^4$ | 1.8       | $-6.00$              | $1.90$               | $-7.90$              |
| IF2          | GDP      | 25    | ADI            | ADI       | ADI                  | ADI                  | ADI                  |
| IF2          | SRL$_{G2655U}$ | 25   | $2.8 \times 10^4$ | 3.6       | $-3.90$              | $3.28$               | $-7.18$              |
| IF2:GDP$_{500}$ | SRL$_{G2655U}$ | 5    | ADI            | ADI       | ADI                  | ADI                  | ADI                  |

*Symbols, abbreviations and other details are given in the Table 1 footnote. Concentration of IF2 was kept constant at 20 μM.

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**Figure 2 | A schema of the interplay among G nucleotide and SRL binding to bacterial GTPases EF-G and IF2.** The affinity constants for different trGTPase complexes (EF-G in red and IF2 in blue) are shown, as measured at 25°C.

**Figure 3 | The SRL contacts three domains of EF-G.** EF-G is coloured by domains, as labeled on the figure. The SRL is shown as space-fill, in grey, while GDP and the magnesium ion are in yellow and green, respectively. The structure is taken from PDB IDs 2WRI and 2WRJ (the crystal structure of EF-G on the ribosome).
Methods
EF-G, IF2, L7/12 and SRL preparations. The HPLC-purified SRL RNA oligonucleotide (5′-GGGCCUCUAUACGAGGAGCCGAGGAGGAGG-3′) was purchased from DNA Technology. The 6His tag E. coli EF-G and IF2 cloned in the expression construct described for Forster and colleagues were overexpressed as described for the non-tagged IF2, and purified in essentially the same way, with an addition of a Ni-NTA purification step before the rest of the chromatographic procedures. Cloning, overexpression and purification of E. coli L7/L12 is described in Supplementary Text S1: SI Methods.

Isothermal titration calorimetry. The thermodynamic parameters of IF2 and EF-G binding to G nucleotides and the SRL RNA oligonucleotide were measured using a MicroCal ITC200 instrument (MicroCal, Northampton, MA) as described. Experiments were carried out at 5 or 25 °C in phosphate buffer (5 mM K2HPO4, 10% glycerol, 1 mM DTT, 95 mM KCl and 5 mM MgCl2, pH 7.5). 2.5-µl aliquots of ligands were injected into the 0.2-ml cell containing the protein solution to achieve a complete binding isotherm. Protein concentration in the cell ranged from 5 to 40 µM and ligand (G nucleotide or SRL RNA oligonucleotide) concentration in the syringe ranged from 50 to 500 µM. Investigations of SF-G binding to L7/L12 used 50 to 400 µM EF-G in the cell and up to 4.7 µM L7/L12 in the syringe. The heat of dilution was measured by injecting the ligand into the buffer solution or by additional injections of ligand after saturation; the values obtained were subtracted from the heat of interactions to obtain the effective heat of binding. The resulting titration curves were fitted using MicroCal Origin software using one binding site model. Interactions were characterized by stochiometry close to unity, indicating high activity and homogeneity of the protein and SRL preparations. Affinity constants (Kd) characterized by stoichiometry close to unity, indicating high activity and homogeneity of the protein and SRL preparations. Affinity constants (Kd) were determined by a non-linear regression fitting procedure. IdLE GTP hydrolysis by GTPTases during titration experiments was assessed by TLC and was not exceeding 1–2%.

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Author contributions

VH conceived the project, coordinated the study and drafted the paper with contributions from VAM, VS, GCA, TT and AAM. VAM, VS, IYuP, AS, VH performed experiments. GCA analyzed the structural data. AAM and TT coordinated the study and contributed materials and reagents.

Additional information

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