

Supplemental Information for

***E. coli* translation strategies differ across nutrient conditions**

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The supplemental information contains:

Supplementary Text

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1. Microscopic model of ribosome dynamics along an mRNA

Ribosome profiling provides positional information of ribosomes occupancy on mRNA transcripts at single nucleotide resolution. In order to investigate ribosomal dynamics on transcripts, we built a microscopic model to estimate the position-dependent behavior of mRNA-bound ribosomes. As shown in the following part of this section, the modeling results lead to a classification of mRNA-bound ribosomes into two groups: “initiating” and “working”, which helps us to build a concise macroscopic model in Section 2.

A. Construction of a representative ribosome-occupancy profile.

We first combine the experimentally obtained ribosome profiling data to obtain an average picture for ribosome occupancy along a “representative” gene with length $N_{\text{aa}} = 300$ amino acids [1]. The average ribosome occupancy $O(x)$ for codon position $x = 1:301$, including the stop codon, was constructed via the following steps:

1. All genes that code for proteins longer than 100 aa and have counts larger than 10 transcripts per million (TPM) were selected. This set of genes accounts for more than 75% of total reads.
2. Total counts from the selected genes were normalized to one million (yielding reads per million, RPM) and the ribosome counts at the first and last 50 codons were obtained by summing the RPM at the corresponding position from all the selected genes (result shown in Figure 2B) and converted to fraction of total counts. This procedure effectively weights genes in the representative ribosome-count profile according to their level of expression.
3. We used the data from the first 50 codons and the last 49 codons before the stop codon to produce a smooth fit to the experimentally obtained ribosome-count profile. A gap with length 201 was inserted between the first 50 and last 49 codons before the stop codon to produce an mRNA producing 300 amino acids. In order to capture the sharp drop of ribosome counts for the first several codons, together with the slower decrease over the rest of the profile, we used a sum of two decaying exponential functions to fit the data. The experimentally measured ribosome count from the stop codon ($x = 301$) was directly taken from the data without fitting. The final results are shown in Figure S3C.

B. The microscopic model for bound ribosomes.

In order to decipher the information about ribosome dynamics contained in ribosome occupancy $O(x)$, we built a microscopic model shown in Figure S3B. In this simplified model, bound ribosomes on codon x move with a step rate $r_s(x)$, and a certain fraction $f_{\text{at}}(x)$ abort translation during this transition. At steady state, the in- and out-flux of ribosomes at every codon x must be balanced:

$$O(x-1) \cdot r_s(x-1) \cdot (1 - f_{\text{at}}(x-1)) = O(x) \cdot r_s(x). \quad \text{Eq.S 1}$$

According to Eq.S 1, under steady state, the relative occupancy of neighboring codons is determined by the step rate and fraction of aborted translation. The profile of $O(x)$ constrains the values of $r_s(x)$ and $f_{at}(x)$. We would like to infer the value of $r_s(x)$ and $f_{at}(x)$ given $O(x)$ within some assumptions. Two possibilities concerning the profile of $O(x)$ and the corresponding assumptions are listed below:

1. $O(x)$ decreases with x . This implies an increased step rate r_s and/or a non-zero fraction of aborted translation f_{at} . For simplicity, we assume a constant fraction (f_{dec}) of the decrease of occupancy ($\frac{O(x-1)}{O(x)} - 1$) can be attributed to an increased step rate ($\frac{r_s(x)}{r_s(x-1)} - 1$):

$$f_{dec} = \frac{\frac{r_s(x)}{r_s(x-1)} - 1}{\frac{O(x-1)}{O(x)} - 1}. \quad \text{Eq.S 2}$$

2. $O(x)$ does not decrease with x . This implies that the step rate r_s does not increase with x . Under this condition, the higher f_{at} , the more r_s decreases with x . According to previous research, the translation elongation rate of ribosomes generally increases from 5' to 3'[2, 3]. Therefore, we assume $f_{at} = 0$ under this condition.

With the former assumptions, we can derive $r_s(x-1)$ and $f_{at}(x-1)$ given $r_s(x)$:

$$r_s(x-1) = \begin{cases} \frac{r_s(x)}{f_{dec} \cdot \left(\frac{O(x-1)}{O(x)} - 1\right) + 1}, & \text{if } \frac{O(x-1)}{O(x)} > 1 \\ r_s(x) \cdot \frac{O(x)}{O(x-1)}, & \text{if } \frac{O(x-1)}{O(x)} \leq 1 \end{cases}, \quad \text{Eq.S 3}$$

$$f_{at}(x-1) = \begin{cases} (1 - f_{dec}) \cdot \left(1 - \frac{O(x)}{O(x-1)}\right), & \text{if } \frac{O(x-1)}{O(x)} > 1 \\ 0, & \text{if } \frac{O(x-1)}{O(x)} \leq 1 \end{cases}. \quad \text{Eq.S 4}$$

C. Calculation of step rate $r_s(x)$ and fraction of aborted translation $f_{at}(x)$.

By applying Eq.S 3 - Eq.S 4 to the ribosome occupancy $O(x)$ obtained experimentally in Section 1A, we can calculate r_s and f_{at} from $x = 300$ to $x = 1$, for assumed values of f_{dec} and the step rate at the last coding codon $r_s(300)$. The existence of aborted translation has been suggested by previous studies [2, 4], however the quantitative fraction of aborted translation under different nutrient-limitations is difficult to quantify. Given the uncertainty of the aborted translation rate, we evenly sampled values of f_{dec} from 0 to 1 and values of $r_s(300)$ from 5 to 20 codon/sec. Each combination of f_{dec} and $r_s(300)$ gives a solution for $r_s(x)$ and $f_{at}(x)$ from $x = 300$ to

$x = 1$. The results are shown in Figure S3D,E: regardless of f_{dec} and $r_s(300)$, aborted translation predominantly happens within the first 10 codons, and the step rate stays relatively flat after the first 10 codons.

D. Classification of ribosomes and mRNAs.

Guided by the above results, we classify mRNA-bound ribosomes into two groups:

1. Initiating ribosomes (symbolized by R_i): ribosomes located within codon 1 to 10 are classified as “initiating”. Given the results in Section 1C, we assume that aborted translation, if it exists at all, only happens to this group of ribosomes. As one ribosome roughly occupies 10 codons on an mRNA [5, 6], the binding of one ribosome at the initiating region excludes the association of another ribosome. Thereby mRNAs are divided into two groups: the freely initiable mRNAs (symbolized by M_f), and the un-initiable mRNAs with one ribosome occupying the 1-10 codon region (symbolized by M_b). By definition, the number of un-initiable mRNAs is equal to the number of initiating ribosomes.

2. Working ribosomes (symbolized by R_w): ribosomes located after the 10-th codon are classified as “working”. They contribute to the increase of biomass. We assume that these ribosomes elongate at a constant rate k_{el} without terminating translation prematurely. Under this assumption, there is no aborted translation in transition from the last coding codon ($x = 300$) to the stop codon ($x = 301$), and the number of ribosomes transiting from the last coding codon to the stop codon is equal to the number of ribosomes leaving stop codon in a steady state. Therefore, the ribosome occupancy at stop codon does not influence the overall protein production rate. In addition, ribosomes occupancy at the stop codon are similar across different nutrient-limitations (Figure 2B), and only take about 2% of total ribosomes. Therefore, we count ribosomes at stop codon also as “working ribosomes” instead of classifying them into another group.

2. Macroscopic model of ribosome dynamics among different states

A. Dynamics of ribosomes and mRNAs between different states.

In Section 1, we classified the mRNA-bound ribosomes into “initiating” and “working” states. This result allowed us to construct a macroscopic model that provides a concise picture of ribosomal dynamics. The transition of ribosomes between three states is shown in Figure 2C. As ribosomal subunits and free 70S ribosomes may convert between each other, they are treated as one group, named unbound ribosomes (symbolized by R_u). A sub-population of the unbound ribosomes can associate with freely initiable mRNAs (M_f) with an effective rate constant k_f . This reaction forms a complex composed of an initiating ribosome (R_i) and an un-initiable mRNA (M_b). The initiating ribosomes either become unbound ribosomes through aborted translation (with rate constant k_r), or proceed to the 11-th codon to become working ribosomes (R_w) (with rate constant k_p). Both processes can release the mRNA back to the freely initiable state. The working ribosomes elongate with constant rate k_{el} . At the last coding codon, a fraction of bound ribosomes ($\sim 1/N_{\text{aa}}$) enters stop codon with rate k_{el} , and the same number of ribosomes complete translation and releases mRNA and become unbound ribosomes.

The non-redundant kinetic ordinary differential equations for the number of ribosomes and mRNAs in each state are:

$$\frac{dR_i}{dt} = k_f \cdot M_f \cdot R_u - k_r \cdot R_i - k_p \cdot R_i, \quad \text{Eq.S 5}$$

$$\frac{dR_w}{dt} = k_p \cdot R_i - \frac{k_{el}}{N_{aa}} \cdot R_w, \quad \text{Eq.S 6}$$

$$\frac{dM_f}{dt} = -k_f \cdot M_f \cdot R_u + k_p \cdot R_i + k_r \cdot R_i. \quad \text{Eq.S 7}$$

Under steady-state growth conditions, the fluxes between ribosomal states should be balanced, and the right-hand sides of Eq.S 5 - Eq.S 7 should all be equal to zero (the production and degradation/dilution by growth of new ribosomes and mRNAs occurs at a negligible rate compared to their recycling). The total number of ribosomes (R_t) and the total number of mRNAs (M_t) are constant parameters for this model:

$$R_u + R_i + R_w = R_t, \quad \text{Eq.S 8}$$

$$M_b + M_f = M_t, \quad \text{Eq.S 9}$$

and by the definition in Section 1D,

$$M_b = R_i. \quad \text{Eq.S 10}$$

B. Estimation of R_t , M_t , k_{el} , and k_p from experimental measurements.

We are interested in the biological mechanisms that lead to distinct ribosomal dynamics under different nutrient conditions. Possible regulatory processes can be represented by parameters in the macroscopic model, including the total number of ribosomes and mRNAs and the rates for ribosomes to initiate, dissociate, and elongate. Some of these parameters, specifically R_t , M_t , k_{el} , and k_p , can be directly estimated from experimental measurements as follows:

1. The total number of ribosomes (R_t): The total mass of rRNA can be obtained by multiplying the RNA-to-protein ratio (RPR), protein mass in a cell (P_m), and the fraction of RNA as rRNA (f_r). Given the mass of the rRNA in a ribosome as m_r , we estimated the total number of ribosomes in a cell:

$$R_t = P_m \cdot RPR \cdot \frac{f_r}{m_r}. \quad \text{Eq.S 11}$$

2. The total number of mRNAs in a cell (M_t): The method is similar to the calculation of R_t in Eq.S 11. Given the fraction of RNA as mRNA as f_m , and the average weight of a nucleotide as m_{nuc} , we have:

$$M_t = P_m \cdot RPR \cdot \frac{f_m}{N_{aa} \cdot m_{nuc} \cdot 3}. \quad \text{Eq.S 12}$$

3. The protein synthesis rate (J_P): The amount of newly synthesized proteins in a cell per second is defined as J_P (aa/sec). J_P can be calculated by two ways: One, the growth rate μ (h^{-1}) can be calculated as the relative rate of protein mass accumulation: $\mu = \frac{J_P}{P_m/m_{aa}}$, where m_{aa} is the average mass of amino acid (g).

Therefore, J_P is linearly proportional to the growth rate:

$$J_P = \mu \cdot \left(\frac{P_m}{3600 \cdot m_{aa}} \right). \quad \text{Eq.S 13}$$

J_P can also be calculated from the contribution of working ribosomes to the growth of the total protein pool:

$$J_P = R_w \cdot k_{el}. \quad \text{Eq.S 14}$$

Combining Eq.S 13 and Eq.S 14, the growth rate μ is determined by three factors: the total number of ribosomes (R_t), the fraction of working ribosomes (ϕ_{Rw}), and the average elongation rate of the working ribosomes (k_{el}):

$$\mu = R_t \cdot \phi_{Rw} \cdot k_{el} \cdot \left(\frac{3600 \cdot m_{aa}}{P_m} \right). \quad \text{Eq.S 15}$$

The contribution of these three factors to the growth rate is illustrated by Figure 2A. The surface in the space of these three factors that corresponds to growth rate $\mu = 0.1 \text{ h}^{-1}$ is shown in Figure S5I, with the estimated values of R_t , ϕ_{Rw} , and k_{el} under various conditions indicated on this surface.

Substituting R_t in Eq.S 15 by Eq.S 11, the relationship between the four experimentally measured values – growth rate (μ), RNA-to-protein ratio (RPR), average elongation rate (k_{el}), and fraction of working ribosomes (ϕ_{Rw}) – is given by:

$$\mu = RPR \cdot \phi_{Rw} \cdot k_{el} \cdot \left(3600 \cdot m_{aa} \cdot \frac{f_r}{m_r} \right). \quad \text{Eq.S 16}$$

Eq.S 16 can be used to estimate growth rate as shown in Figure 1F. Using different definitions of ϕ_{Rw} , we show that it can lead to different estimations of growth rate. The ϕ_{Rw} we used for modeling is from the following Section 2B4. While growth rate, RNA-to-protein ratio and fraction of working ribosomes are global measurements, the elongation rate was measured using the *lacZ* induction assay.

In order to obtain a global estimate of the elongation rate k_{el} , we derived k_{el} from the measurements of growth rate, RNA-to-protein ratio, and the fraction of working ribosomes by the following steps 4-5.

4. Quantifying the fraction of different ribosomal species: The fraction of unbound ribosomes is the sum of the free 70S fraction and the subunit fraction ($\phi_{Ru} = \phi_{R70Sf} + \phi_{RS}$). The fraction of initiating ribosome is calculated by multiplying the fraction of bound ribosomes by the sum of ribosome occupancies at the first 10 codon ($\phi_{Ri} = \sum_{x=1}^{10} O(x) \cdot (\phi_{R70Sb} + \phi_{Rpoly})$). The fraction of working ribosomes is calculated by multiplying the fraction of bound ribosomes with the summation of ribosome occupancy at the 11-th to 301-th codon ($\phi_i = \sum_{x=11}^{301} O(x) \cdot (\phi_{R70Sb} + \phi_{Rpoly})$).
5. Estimating the average working ribosome elongation rate k_{el} : From Eq.S 16 and the definition of ϕ_{Rw} , k_{el} can be calculated by measurements of RPR and ϕ_{Rw} :

$$k_{el} = \frac{\mu}{RPR \cdot \phi_{Rw} \cdot 3600 \cdot m_{aa} \cdot \frac{f_r}{m_r}}. \quad \text{Eq.S 17}$$

The resulting k_{el} is within the error range of the elongation rates measured from the *lacZ* induction assay (Figure S5A).

6. The rate constant k_p for ribosomes to proceed from initiating to working: According to Eq.S 6, at steady state,

$$k_p = \frac{k_{el}}{N_{aa}} \cdot \frac{\phi_{Rw}}{\phi_{Ri}}. \quad \text{Eq.S 18}$$

C. Fraction of working ribosomes (ϕ_{Rw}) as a function of kinetic parameters.

Among the three factors in Eq.S 15 that determine growth rate, R_t and k_{el} are parameters of the macroscopic model. In contrast, the fraction of working ribosomes, ϕ_{Rw} , is the outcome of dynamic processes described by the parameters in the macroscopic model. We would like to infer the relationship between ϕ_{Rw} and these parameters, in order to identify the mechanisms that lead to the different observed ϕ_{Rw} under different conditions.

The parameters k_r and k_f cannot be obtained experimentally. Nevertheless, we will show in the following part of this section that these two parameters work in combination to influence the value of ϕ_{Rw} , and the combined parameter can be obtained from experimental measurements.

According to Eq.S 5-Eq.S 10, in steady state, we obtained an expression for ϕ_{Rw} :

$$\phi_{Rw} = \frac{F}{2} \cdot \left(\left(\frac{1}{S \cdot (F+1)} + \frac{M_t}{R_t} \right) - \left(\left(\frac{1}{S \cdot (F+1)} + \frac{M_t}{R_t} \right)^2 - 4 \cdot \frac{M_t}{R_t} \cdot \frac{1}{F+1} \right)^{\frac{1}{2}} \right), \quad \text{Eq.S 19}$$

with

$$F = N_{aa} \cdot \frac{k_p}{k_{el}},$$

$$K_M = \frac{k_r + k_p}{k_f},$$

$$S = \frac{R_t}{K_M + R_t}.$$

Eq.S 19 depends on three combined parameters: $\frac{M_t}{R_t}$ and F, S . According to Eq.S 11 and Eq.S 12, $\frac{M_t}{R_t} = \frac{f_m \cdot m_r}{N_{aa} \cdot m_{nuc} \cdot 3 \cdot f_r}$ is a constant across different conditions under our assumptions. Physically, the combined parameter F can be interpreted as the “relative proceeding rate”, reflecting the rate for ribosomes to proceed from an initiating to a working state relative to the average elongation rate. If there were no regulation at this step, F would be constant across different conditions regardless of elongation rates. However, if there is nutrient-specific regulation on the step rate or aborted translation rate within the first 10 codons, F will have different values for different conditions. The value of F can be calculated from the ribosomal profiling data according to Eq.S 18: $F = \frac{\sum_{x=11}^{x=301} O(x)}{\sum_{x=1}^{x=10} O(x)}$. Figure S5B shows the values of F under C-, N-, P-limitations ($\mu = 0.1 \text{ h}^{-1}$) and Minimal condition ($\mu = 0.9 \text{ h}^{-1}$) for WT and the *relA* mutant.

The lumped parameter S can be interpreted as a “saturation parameter”: According to Eq.S 5-Eq.S 10 and Eq.S 13-Eq.S 14, the growth rate can be expressed as a function of R_u , mimicking the form of Michaelis-Menten:

$$\mu(R_u) = \frac{3600 \cdot N_{aa} \cdot m_{aa}}{P_m} \cdot M_t \cdot k_p \cdot \frac{R_u}{K_m + R_u}. \quad \text{Eq.S 20}$$

In analogy to enzymatic reactions, in our model, ribosome subunits can be viewed as the substrates, mRNA as the enzyme, and working ribosomes as the product, and therefore K_m is interpreted as the half-saturation concentration of unbound ribosomes. Therefore, $S = \frac{R_t}{K_m + R_t}$ positively correlates with the degree of saturation for ribosomes in translation: A value of S near zero implies a large fraction of mRNAs are initiatable, waiting for ribosomes, while S approaching one implies most mRNAs are occupied in the initiation region and increasing in ribosome number cannot substantially boost growth rate. The value of S can be calculated by quantifying the ribosomal species according to Eq.S 5, Eq.S 11, and Eq.S 12: $S = 1 / (1 + (\frac{f_m \cdot m_r}{N_{aa} \cdot m_{nuc} \cdot 3 \cdot f_r} - \phi_{Ri}) \cdot \frac{\phi_{Ru}}{\phi_{Ri}})$. Figure S5C shows the values of S under C-, N-, P-limitations ($\mu = 0.1 \text{ h}^{-1}$) and Minimal condition ($\mu = 0.9 \text{ h}^{-1}$) for wild type and the *relA* mutant.

According to Eq.S 19, ϕ_{Rw} is an increasing function of both F and S (Figure S5H). In wild type, the relative proceeding rate F and the saturation parameter S both increase from C-, N- P-limitation to Minimal condition, leading to an increasing ϕ_{Rw} . The *relA* mutant under P-limitation has a very high relative proceeding rate, consistent with the known biological role of *relA*. By contrast, the loss of *relA* does not affect the saturation parameter.

D. Growth rate μ as a function of total ribosome number R_t and total mRNA number M_t .

Ribosomes and mRNAs are two major players in translation. If the number of ribosomes were too high relative to mRNAs, most initiation regions of mRNAs would be occupied, leaving a large number of idle unbound ribosomes; if the number of mRNAs were too high, there would not be enough ribosomes for each mRNA to be translated efficiently. Given our detailed estimation of the parameters associated with translation under different nutrient conditions, we are interested in how the total number of ribosomes and mRNAs influences the protein production rate, and which one is the limiting factor for cell growth.

Combining Eq.S 15 and Eq.S 19, yields the growth rate μ as a function of R_t, M_t , and other kinetic parameters of the macroscopic model:

$$\mu(R_t, M_t) = \left(\frac{3600 \cdot m_{aa}}{P_m} \right) \cdot \frac{k_p}{2} \cdot N_{aa} \cdot \left((A \cdot K_M + A \cdot R_t + M_t) - \left((A \cdot K_M + A \cdot R_t + M_t)^2 - 4 \cdot M_t \cdot R_t \cdot A \right)^{\frac{1}{2}} \right) \quad \text{Eq.S 21}$$

with

$$A = \frac{1}{F + 1} = \frac{1}{\frac{N_{aa} \cdot k_p}{k_{el}} + 1}$$

This function resembles the shape of a Michaelis-Menten function of both R_t and M_t . Intuitively, $\mu = 0$ when R_t or M_t equal to zero. If R_t increases while M_t is held constant, μ monotonically increases and saturates at

$$\mu(\infty, M_t) = \left(\frac{3600 \cdot m_{aa}}{P_m} \right) \cdot k_p \cdot N_{aa} \cdot M_t \quad \text{Eq.S 22}$$

The half-saturation value of R_t is also linearly increasing with M_t :

$$R_{t,1/2} = \frac{1}{2 \cdot A} \cdot M_t + K_M \quad \text{Eq.S 23}$$

Figure S5 D,E show the functions $\mu(R_t)$, the half-saturation values of R_t , and the estimated values of R_t under different nutrient conditions while M_t is held constant.

If M_t increases while R_t is held constant, μ monotonically increases and saturates at

$$\mu(R_t, \infty) = \left(\frac{3600 \cdot m_{aa}}{P_m} \right) \cdot k_p \cdot N_{aa} \cdot A \cdot R_t \quad \text{Eq.S 24}$$

The half-saturation value of M_t is also linearly increasing with R_t :

$$M_{t,1/2} = \frac{A}{2} \cdot R_t + A \cdot K_M \quad \text{Eq.S 25}$$

Figure S5 F,G show the functions $\mu(M_t)$, the half-saturation values of M_t , and the estimated values of M_t under different nutrient conditions while R_t is held constant.

As shown in Figure S5 D-G, in all conditions, the estimated ribosome numbers are smaller than their half-saturation concentrations, falling in the linear region. By contrast,

the estimated mRNA number is close to or larger than mRNA half-saturation value, albeit not fully saturated. In future, it will be interesting to directly probe the degree of saturation for both ribosomes and mRNAs experimentally.

E. Predicting the growth rate after nutrient upshift to rich medium.

As mentioned in the main text, there appear to be diverse strategies that achieve the same growth rate under different nutrient limitations, but these strategies may lead to different outcomes when the environment changes. One possibility is that having extra ribosomes at slow growth rates can enable a faster recovery when nutrients become abundant again. Therefore, we expanded our macroscopic model beyond steady state to predict growth dynamics upon nutrient upshift.

The rate of cell growth positively correlates with the rate of protein synthesis. Among newly synthesized proteins, a certain fraction of mass is allocated to ribosomal proteins ($\Psi_R(t)$). The steady-state value of this mass fraction (Ψ_R^*) under a given condition (i) can be calculated from the estimated number of total ribosomes:

$$\Psi_{R,i}^* = R_t \cdot N_{Raa} \cdot \frac{m_{aa}}{P_m}. \quad \text{Eq.S 26}$$

N_{Raa} is the number of amino acids in a ribosome, and m_{aa} is the average mass of an amino acid in *E. coli* protein. The time-dependent growth rate $g(t)$ of a bacterial population can be defined in terms of the increase in total cell volume:

$$g(t) = \frac{V'(t)}{V(t)}. \quad \text{Eq.S 27}$$

Assuming the concentration of protein (C_p) remains constant, the rate of increase of total volume will be linearly related to the rate of increase of protein mass, which is proportional to the concentration of working ribosomes $r_b(t)$ and the translation completion rate k_{el}/N_{aa} :

$$V'(t) = V(t) \cdot k_{el} \cdot r_b(t) \cdot \frac{m_{aa}}{C_p}. \quad \text{Eq.S 28}$$

Based on Eq.S 27 and Eq.S 28, the growth rate can be expressed as:

$$g(t) = k_{el} \cdot r_b(t) \cdot \frac{m_{aa}}{C_p}. \quad \text{Eq.S 29}$$

However, during the process of reaching steady state, $\Phi_R(t)$ might be regulated, e.g. to accelerate cell growth, and therefore could be a time-dependent function.

Applying the chain rule of differentiation to intracellular concentrations, $\frac{d(\frac{X}{V})}{dt} = \frac{1}{V} \cdot \frac{dX}{dt} - \frac{X}{V} \cdot \left(\frac{1}{V} \cdot \frac{dV}{dt}\right)$, the rates of change in the concentration of different ribosome species can be expressed as:

$$\frac{dr_t(t)}{dt} = \Psi_R(t) \cdot \frac{k_{el}}{N_{aa}} \cdot r_w(t) - g(t) \cdot r_t(t), \quad \text{Eq.S 30}$$

$$\frac{dr_i}{dt} = k_f \cdot M_f \cdot (r_t(t) - r_i(t) - r_w(t)) - k_r \cdot r_i(t) - k_p \cdot r_i(t) - g(t) \cdot r_i(t), \quad \text{Eq.S 31}$$

$$\frac{dr_w(t)}{dt} = k_p \cdot r_i(t) - \frac{k_{el}}{N_{aa}} \cdot r_w(t) - g(t) \cdot r_w(t). \quad \text{Eq.S 32}$$

We are interested in the growth dynamics after switching cells from C-, N-, or P-limitation to rich medium. We therefore solved Eq.S 30-Eq.S 32 for parameter values obtained under rich condition. The initial states of r_t , r_i , and r_w were taken as their steady-state values (r_t^* , r_i^* , and r_w^*) under C-, N-, and P-limitations, respectively.

The fraction of newly synthesized ribosomal proteins $\Psi_R(t)$ reflects the regulation of protein allocation by the cell. Eventually this value needs to reach the steady-state Ψ_R^* in rich media. Nevertheless, during the transition period, different strategies of regulating $\Psi_R(t)$ yield different growth dynamics. In our model, we tested three different control strategies for $\Psi_R(t)$: Bang-Bang control

$$\left(\Psi_R(t) = \begin{cases} 1 & (\text{if } r_t < r_{t,\text{rich}}^*) \\ \Psi_{R,\text{rich}}^* & (\text{if } r_t = r_{t,\text{rich}}^*) \\ 0 & (\text{if } r_t > r_{t,\text{rich}}^*) \end{cases} \right), \quad \text{Eq.S 33}$$

smooth control ($\Psi_R(t) = 2 \cdot \Psi_{R,\text{rich}}^* / (\frac{r_t}{r_{t,\text{rich}}^*} + 1)$), and steady-state control ($\Psi_R(t) = \Psi_{R,\text{rich}}^*$). Interestingly, we found that a Bang-Bang control strategy gave the prediction closest to our experimentally observed dynamics (Figure S8). It has been shown that Bang-Bang control of regulation can maximize the accumulated increase of cell volume [7-9].

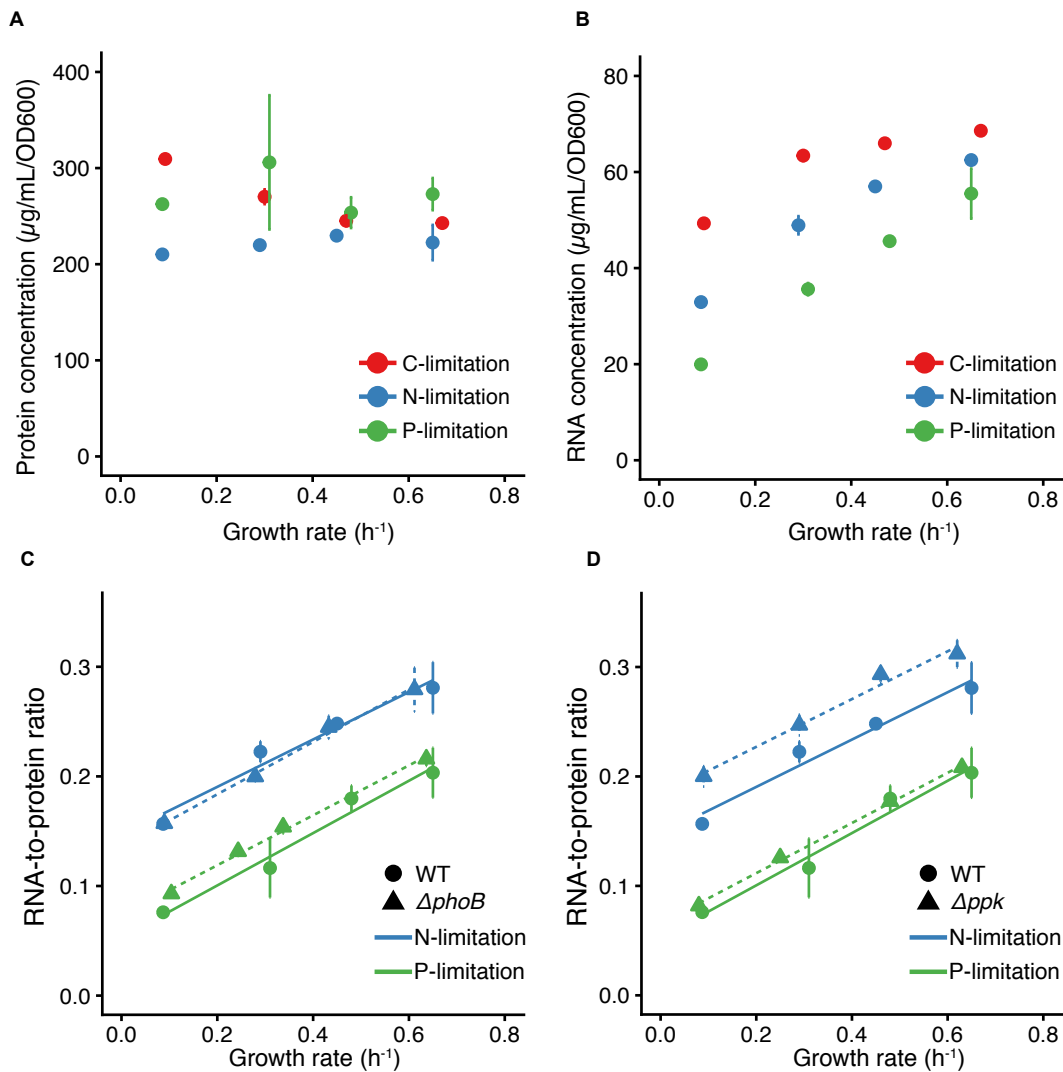


Figure S1. (Related to Figure 1) Lower RNA-to-protein ratio under P-limitation results from lower RNA concentration, and phosphorus metabolism-related genes are not involved in the regulation. (A) Total protein ($\mu\text{g/mL/OD600}$) and **(B)** RNA ($\mu\text{g/mL/OD600}$) concentrations were measured. These data were used to generate Figure 1B of the RNA-to-protein ratio. Each data point represents three technical replicates with standard deviation plotted as error bars. **(C)** RNA-to-protein ratio for chemostat cultures of wild-type and ΔphoB cells upon N-, and P-limitations at different growth rates. Each data point represents three technical replicates with standard deviations after error propagation shown as error bars. **(D)** RNA-to-protein ratio for chemostat cultures of wild-type and Δppk cells upon N-, and P-limitations at different growth rates. Each data point represents three technical replicates with standard deviations after error propagation shown as error bars.

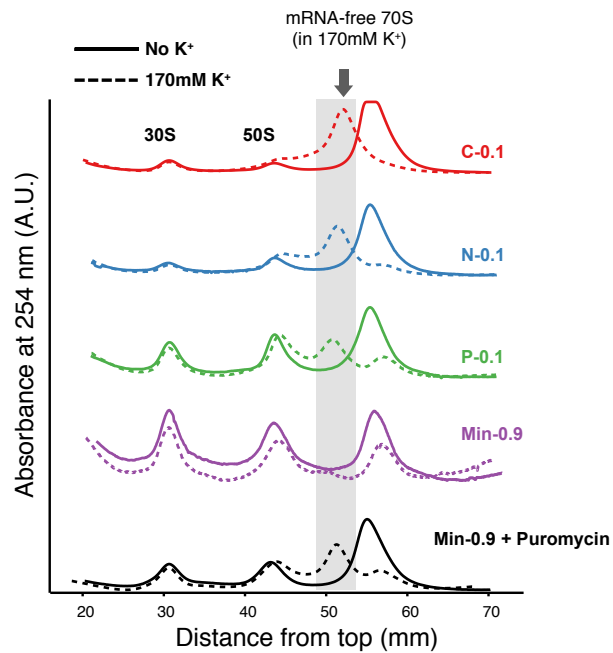


Figure S2. (Related to Figure 1) Elevated $[K^+]$ distinguishes mRNA-free from mRNA-bound 70S ribosomes. The plot shows the separation of these two distinct 70S species under different growth conditions using 170 mM K^+ . The A_{254} values are vertically shifted for each condition for easier visualization. Puromycin is used as a positive control for free ribosome accumulation and fast-growing cells in glucose minimal media (Min-0.9) are used as a negative control for cells expected to lack free ribosomes.

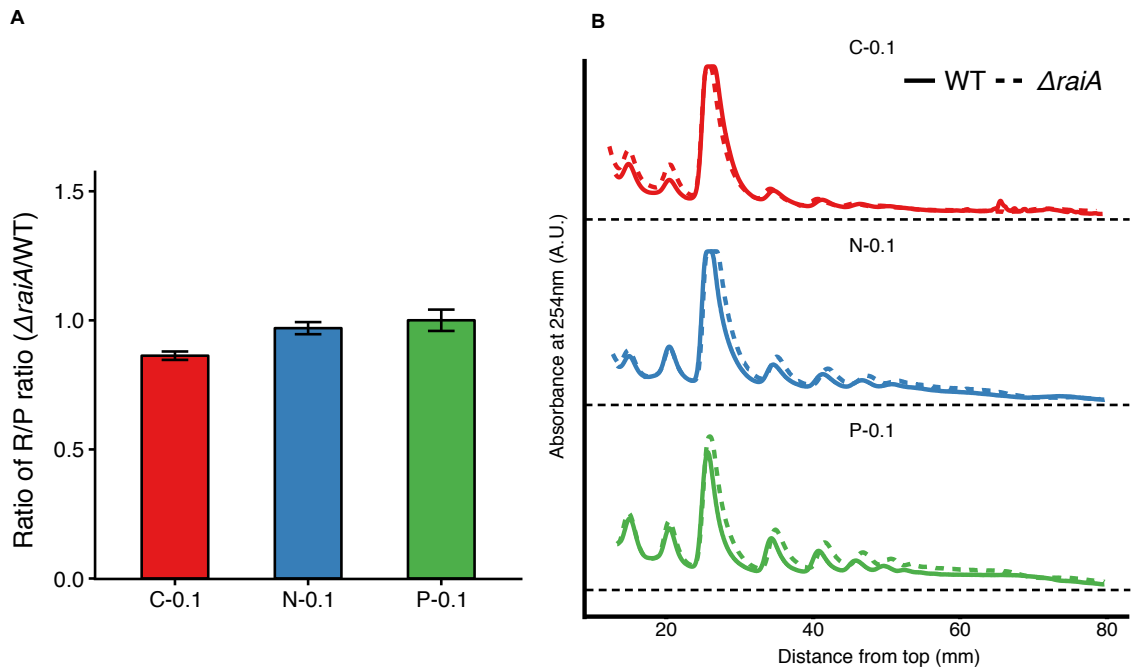


Figure S3. (Related to Figure 1) Deletion of *raiA* does not alter ribosome dynamics. (A) RNA-to-protein ratio of wild type and $\Delta raiA$ at growth rate 0.1 h^{-1} under different nutrient limitations. Each data point represents three technical replicates with standard deviations after error propagation shown as error bars. **(B)** Polysome profiles of wild type and $\Delta raiA$ cells. Same amount of RNA from WT and $\Delta raiA$ was loaded for comparison. Deletion of *raiA* has no effect on the size of 70S peak. 100S peak from ribosome dimerization was not detected under any condition. The thin dashed line marks the average baseline.

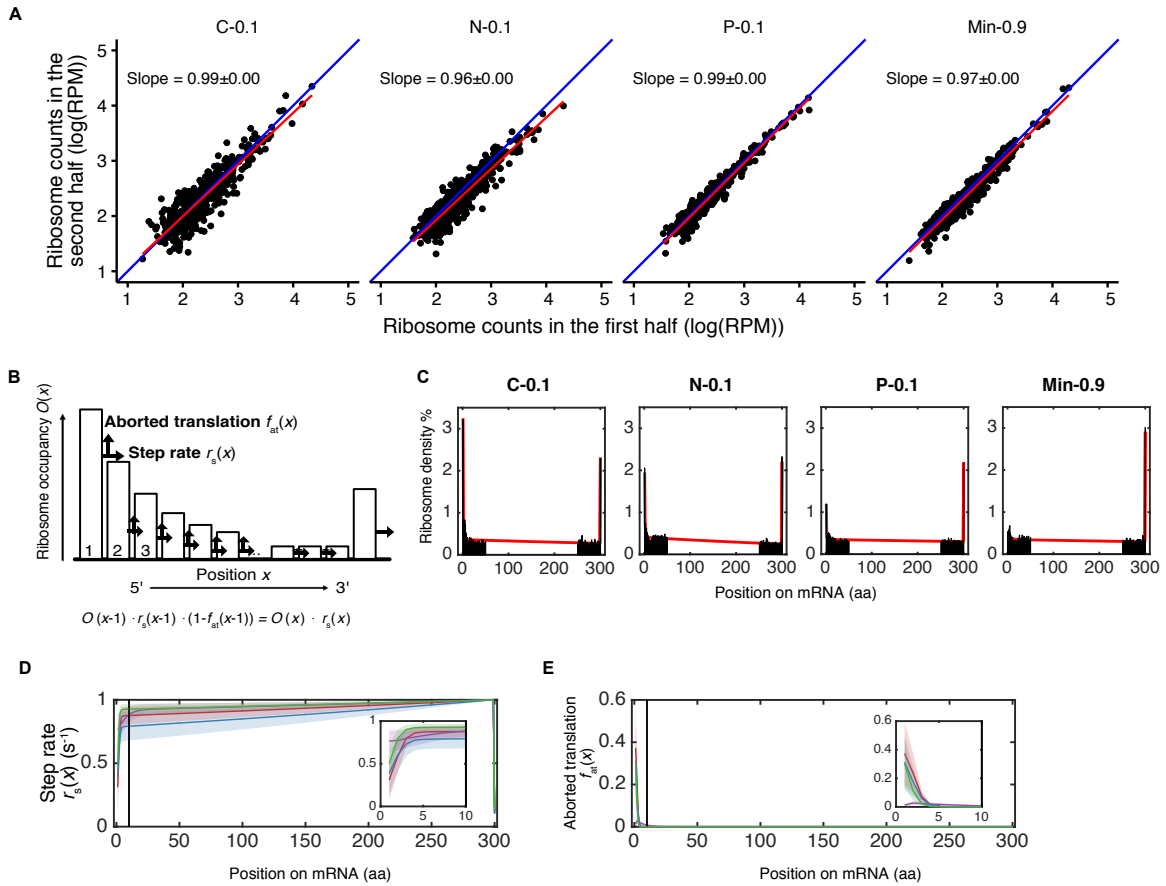


Figure S4. (Related to Figure 2) Modeling ribosome dynamics on an mRNA with a microscopic model. (A) Comparison of ribosome density between first and second halves of genes. Scatter plot of summed RPM (reads per million) in the first and second halves of each gene. Blue line has a slope of one. Red line is the fitting result with the value of slope marked in the graph. **(B)** Illustration of the microscopic model for bound ribosome dynamics on mRNA, and the corresponding flux balance equation. **(C)** Dark bars are the normalized ribosome count in the first 50 and the last 50 codons from ribosome profiling studies. Red curves are the smoothed ribosome occupancies obtained by fitting the normalized ribosome count in the first 50 and the last 49 codons to two-term exponential functions, and copying the observed count number at the stop codon. **(D-E)** The position-dependent normalized step rate (s^{-1}) and fraction of aborted translation that best fit the smoothed ribosome occupancies in (C). Insets are magnifications for codon position 1-10.

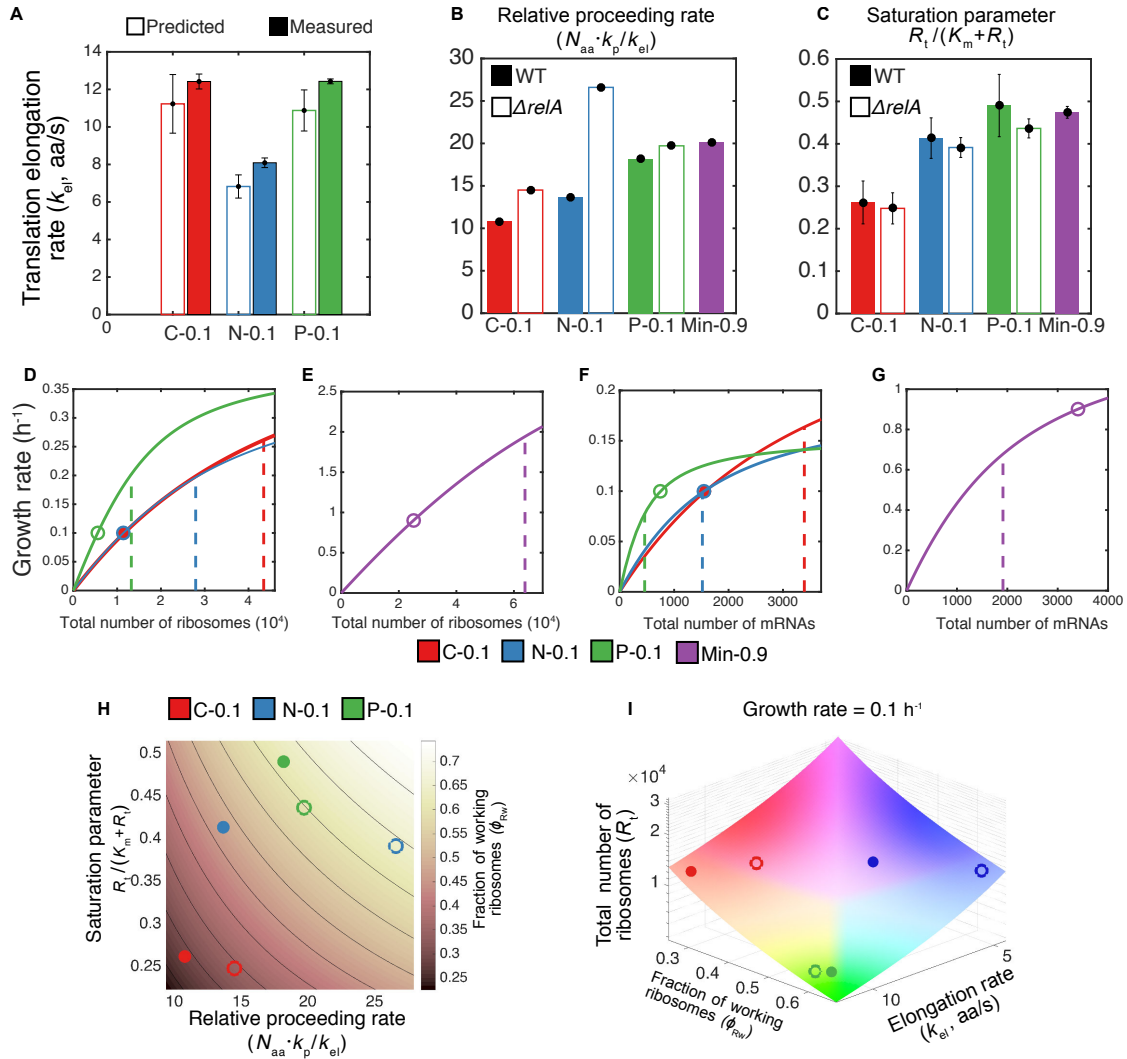


Figure S5. (Related to Figure 2) Parameters of the macroscopic model relate translation activity and growth. (A) Comparison of predicted elongation rates from the macroscopic ribosome dynamic model with experimentally measured ones. **(B-C)** The relative proceeding rate ($N_{aa} \cdot k_p / k_{el}$) and saturation parameter ($R_t / (K_m + R_t)$) under various conditions. **(D-G)** Model relationships between total number of ribosomes (D-E) or mRNAs (F-G) and growth rate, while the total number of mRNAs or ribosomes is held constant. Dashed lines indicate the half-saturation concentration, and dots mark the estimated values of ribosomes or mRNAs under each condition. **(H)** Fraction of working ribosomes as a function of relative proceeding rate and saturation parameters. Color indicates the fraction of working ribosomes (ϕ_{rw}) as a function of relative proceeding rate and saturation parameter. Filled circles indicate the values for C-, N-, P-limited wild type cells, and open circles indicate corresponding values in $\Delta relA$ mutants. **(I)** The relationship between elongation rate (k_{el}), fraction of working ribosomes (ϕ_{rw}), and total number of ribosomes (R_t) that leads to the same growth rate of $0.1 h^{-1}$. Filled circles indicate the values for C-, N-, P-limited wild type cells, and open circles indicate the corresponding values for the $\Delta relA$ mutant.

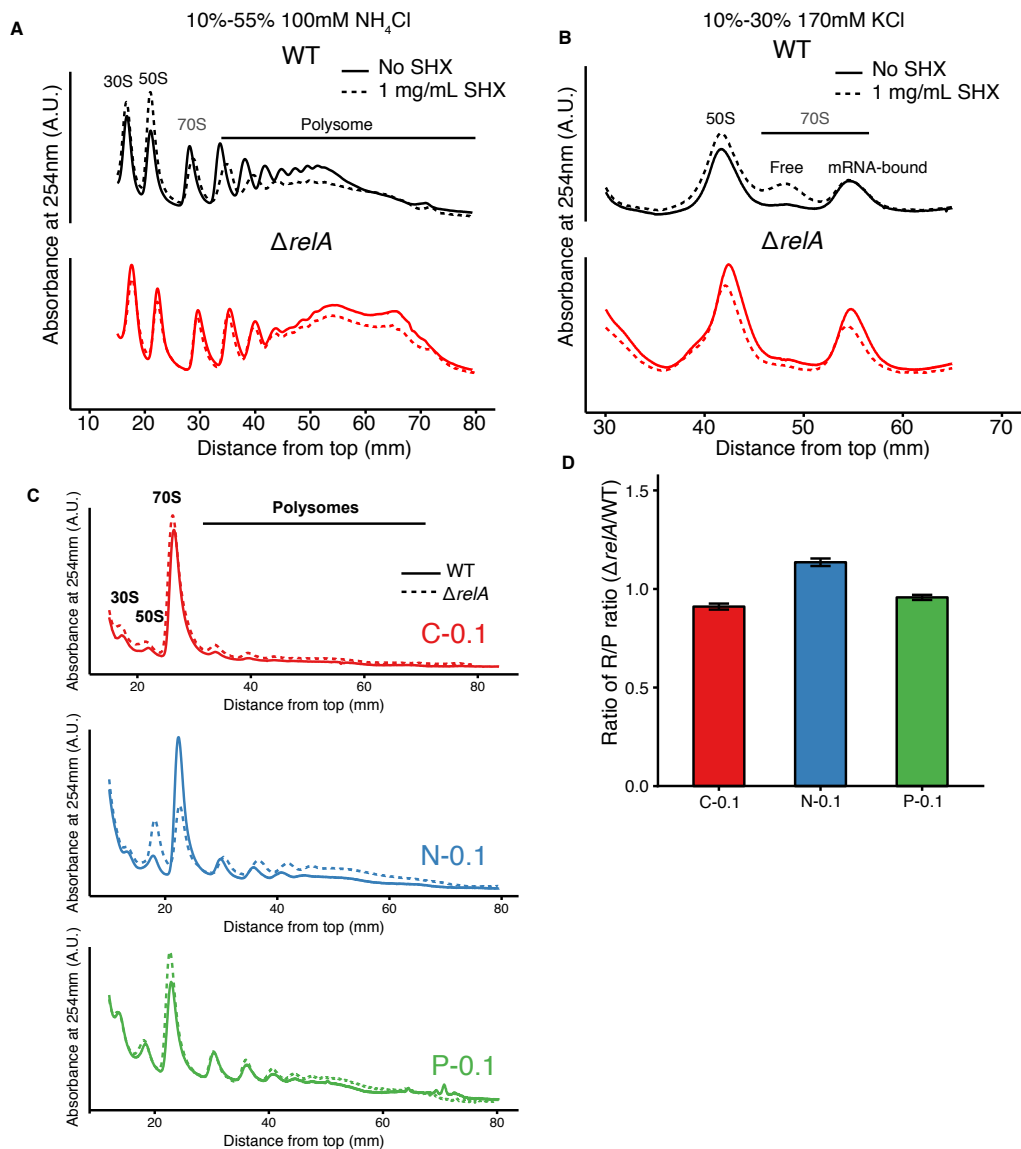


Figure S6. (Related to Figure 3) *relA*-dependent ppGpp production leads to free ribosome accumulation under stringent response and nitrogen limitation. (A) Polysome profile of wild type and $\Delta relA$ cells with and without treatment of serine hydroxamate (SHX) for ten minutes. **(B)** Free-ribosome profiling using 170 mM KCl to distinguish mRNA-free and mRNA-bound ribosomes. **(C)** Polysome profile of wild type and $\Delta relA$ cells at growth rate of 0.1 h^{-1} . **(D)** Ratios of RNA-to-protein ratios of wild type and $\Delta relA$ at growth rate of 0.1 h^{-1} . Three technical replicates were collected and standard deviation was plotted as error bars.

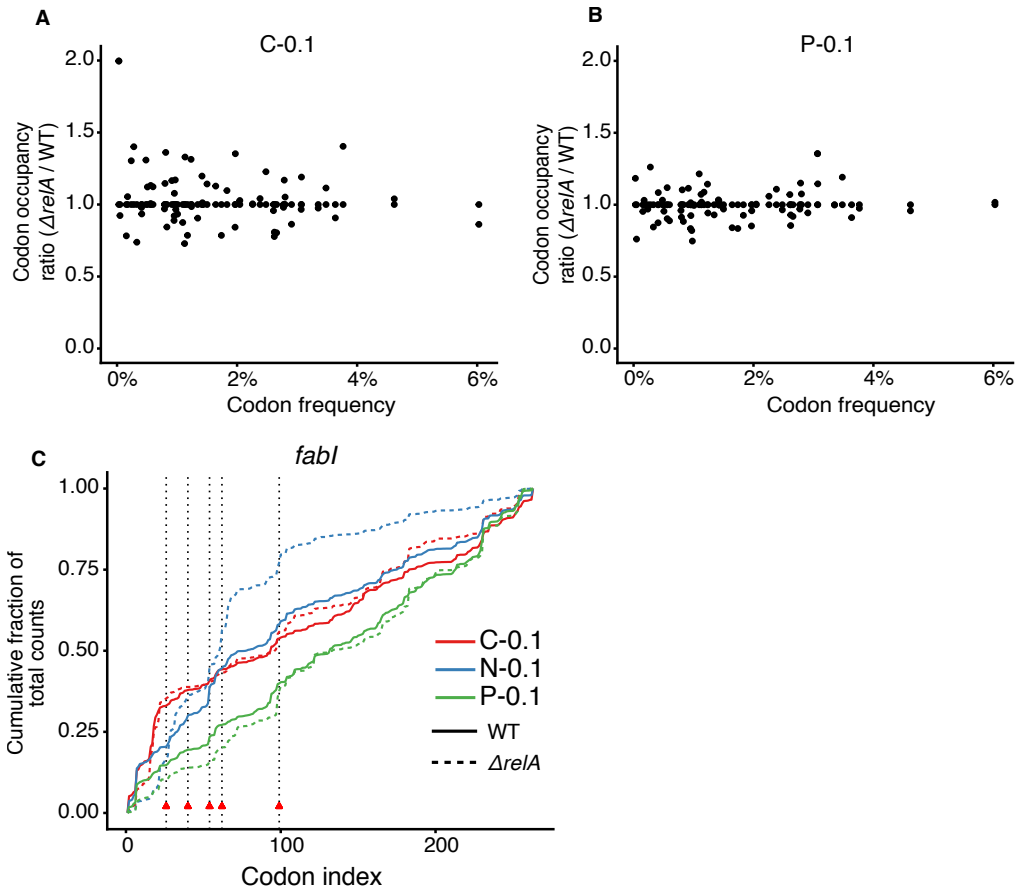


Figure S7. (Related to Figure 3) Deletion of *relA* disrupts translation under nitrogen limitation. (A) Ratio of codon occupancy between $\Delta relA$ and wild type cells under carbon limitation at 0.1 h^{-1} . (B) Ratio of codon occupancy between $\Delta relA$ and wild type cells under phosphorus limitation at 0.1 h^{-1} . (C) Cumulative fraction of ribosome counts of *fabI* in wild type and $\Delta relA$ under C-, N- and P-limitations at 0.1 h^{-1} . The vertical dotted lines and red triangles mark the positions of glutamine codons.

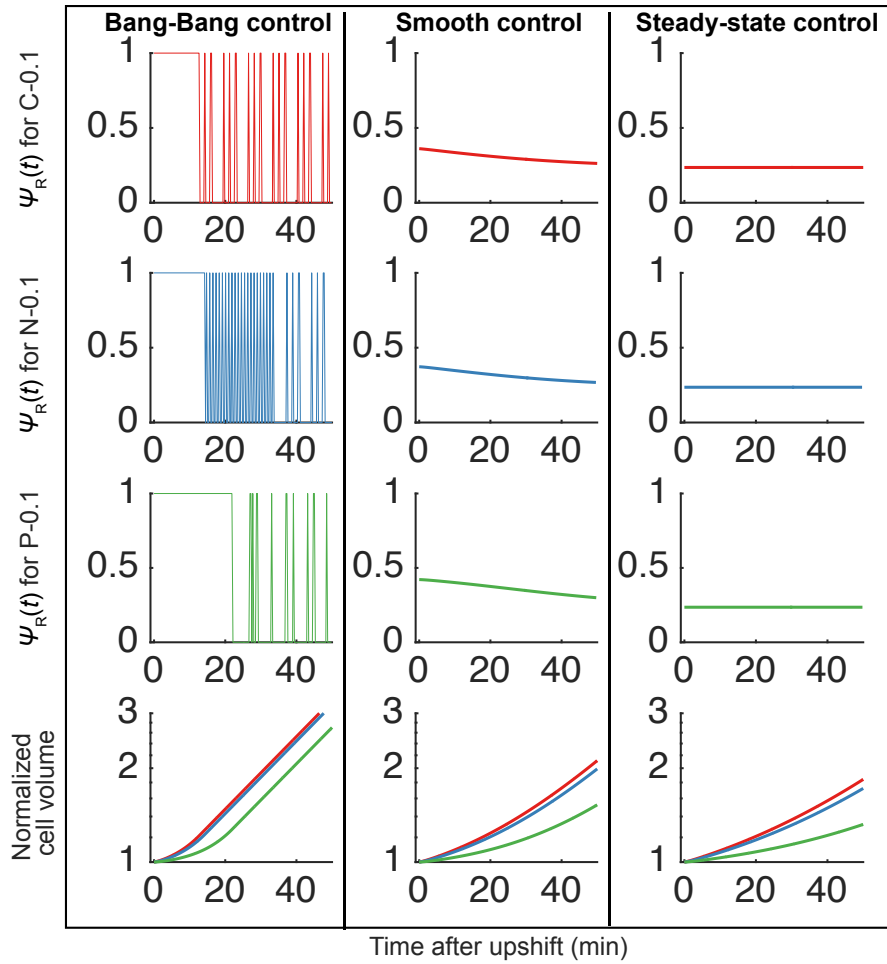


Figure S8. (Related to Figure 4) Different allocation strategies determine growth dynamics upon modeled nutrient upshift to rich medium. After nutrient upshift to a rich condition, different regulatory strategies for $\psi_R(t)$, the fraction of new protein synthesis allocated to ribosomal proteins (first three rows for C-, N-, and P-limitations, respectively), and their resulting post-upshift growth curves (last row).

Table S1. Definitions and values of parameters used in ribosome dynamics models.

Symbol	Definition	Value		Source
RPR	Mass ratio of total RNA to total Protein	WT C-limit	0.16	This work
		WT N-limit	0.16	
		WT P-limit	0.08	
		<i>relA</i> C-limit	0.16	
		<i>relA</i> N-limit	0.16	
		<i>relA</i> P-limit	0.08	
		WT Min	0.34	
		WT Rich	0.56	
μ	Steady-state growth rate (h^{-1})	WT C-limit	0.09	This work
		WT N-limit	0.09	
		WT P-limit	0.09	
		<i>relA</i> C-limit	0.09	
		<i>relA</i> N-limit	0.09	
		<i>relA</i> P-limit	0.09	
		Min	0.93	
		Rich	1.6	
ϕ_{Ru}	Fraction of ribosomes that are not bound to mRNA	WT C-limit	0.69	This work
		WT N-limit	0.49	
		WT P-limit	0.35	
		<i>relA</i> C-limit	0.64	
		<i>relA</i> N-limit	0.34	
		<i>relA</i> P-limit	0.37	
		WT Min	0.34	
ϕ_{Ri}	Fraction of ribosomes locating at the first 10 codon of mRNA, may abort translation prematurely	WT C-limit	0.03	This work
		WT N-limit	0.03	
		WT P-limit	0.03	
		<i>relA</i> C-limit	0.02	
		<i>relA</i> N-limit	0.02	
		<i>relA</i> P-limit	0.03	
ϕ_{Rw}	Fraction of ribosomes locating after the first 10 codon of mRNA, contribute to the protein production in constant elongation rate	WT C-limit	0.29	This work
		WT N-limit	0.47	
		WT P-limit	0.62	
		<i>relA</i> C-limit	0.34	
		<i>relA</i> N-limit	0.64	
		<i>relA</i> P-limit	0.60	
		WT Min	0.63	
f_r	Fractional mass of rRNA among total RNA	0.85		[10]
f_m	Fractional mass of mRNA among total RNA	0.02		[11]
V_c	Cell volume (m^3)	10^{-18}		[12]

C_p	Concentration of proteins (g/m ³)	2.4×10^5	[13]
m_r	Mass of the rRNA component in a ribosome (g)	2.8×10^{-18}	[14]
m_{nuc}	Average mass of a nucleotide in RNA in <i>E. coli</i> (g)	5.4×10^{-22}	[15]
m_{aa}	Average mass of an amino acid in <i>E. coli</i> protein (g)	1.8×10^{-22}	[16]
N_{Raa}	Number of amino acids in the ribosome	7459	[17]
N_{aa}	Average mRNA length (aa)	300	[1]
P_m	Total protein mass in a cell (g)	$C_p \cdot V_c$	
$O(x)$	Average occupancy of ribosomes at the x -th codon	Obtained by fitting the data from ribosome profiling	
$r_s(x)$	Step rate of ribosomes from codon x to the next codon		
$f_{at}(x)$	Fraction of ribosomes that abort translation during transition from codon x to the next codon		
R_t	Total number of ribosomes per cell	$P_m \cdot RPR \cdot \frac{f_r}{m_r}$	
R_i	Initiating ribosomes: ribosomes located within the first 10 codons	$R_t \cdot \phi_{Ri}$	
R_w	Working ribosomes: ribosomes located within codon 11 to 301.	$R_t \cdot \phi_{Rw}$	
R_u	Unbound ribosomes, including 70S free ribosomes and subunits	$R_t \cdot \phi_{Ru}$	
M_t	Total number of mRNAs per cell	$P_m \cdot RPR \cdot \frac{f_m}{N_{aa} \cdot m_{nuc} \cdot 3}$	
M_b	Un-initiable mRNAs, with one ribosome bound within the first 10 codons	R_i	
M_f	Freely initiable mRNAs, with no ribosome bound within the first 10 codons	$M_t - M_b$	
k_f	Effective rate constant for unbound ribosomes and free mRNAs to initiate translation (1/s)		
k_r	Rate constant for initiating ribosomes to abort translation		

	(1/s)	
k_{el}	Elongation rate for working ribosomes (aa/s)	$\frac{\mu}{RPR \cdot \phi_{Rw} \cdot 3600 \cdot m_{aa} \cdot \frac{f_r}{m_r}}$
k_p	Rate constant for initiating ribosomes to transition into working ribosomes (1/s)	$\frac{k_{el} \cdot \phi_{Rw}}{N_{aa} \cdot \phi_{Ri}}$
J_P	Total rate of protein synthesis (aa/s)	$J_P = \mu \cdot \left(\frac{P_m}{3600 \cdot m_{aa}} \right)$ $J_P = R_w \cdot k_{el}$
F	Relative proceeding rate	$\frac{N_{aa} \cdot k_p}{k_{el}}$
S	Saturation parameter	$\frac{k_r + k_p}{k_f} + R_t$ R_t
$V(t)$	Volume of the population at time t	
$g(t)$	Growth rate of the population at time t	
$\Psi_R(t)$	Fraction of newly-synthesized proteins allocated to ribosomal proteins	
r_t	Concentration inside cells of total number of ribosomes	
r_i	Concentration inside cells of initiating ribosomes	
r_w	Concentration inside cells of working ribosomes	

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