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# Bacterial growth: global effects on gene expression, growth feedback and proteome partition

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The function of endogenous as well as synthetic genetic circuits is generically coupled to the physiological state of the cell. For exponentially growing bacteria, a key characteristic of the state of the cell is the growth rate and thus gene expression is often growth-rate dependent. Here we review recent results on growth-rate dependent gene expression. We distinguish different types of growth-rate dependencies by the mechanisms of regulation involved and the presence or absence of an effect of the gene product on growth. The latter can lead to growth feedback, feedback mediated by changes of the global state of the cell. Moreover, we discuss how growth rate dependence can be used as a guide to study the molecular implementation of physiological regulation.

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## Introduction

Many bacteria can grow exponentially with a wide range of growth rates, depending on the nutrient-content of the growth medium. *Escherichia coli* for example can grow with doubling times ranging from 20 min to many hours. The study of bacterial growth has played a central role in the development of modern microbial physiology as well as in the discovery of the regulation of gene expression [1–3]. It constitutes a rather natural subject of interest for systems biology [4,5], as cell growth obviously is a systems-level phenomenon that depends on the coordinated functions of many cellular components [6], and new experimental tools allow to study growth with unprecedented control over the growth conditions [7\*\*].

Moreover, bacterial growth provides a model system for studying the coupling between individual genetic circuits and the global state of the cell. With recent progress in designing synthetic genetic circuits [8–10] and in quantitative studies of physiological gene regulation systems [11,12\*], it has become clear that genetic circuits are not strictly insulated from their ‘host’ cell, but rather are coupled to the physiological state of the cells as a whole. For example, gene expression requires RNA polymerases and ribosomes, and the concentration of these macromolecular machineries is dependent on the growth conditions [13,14]. These observations point towards the limitations of analogies between genetic and electronic circuits and of metaphors that describe the host cell of a circuit as a ‘chassis’ [15,16], a rigid framework onto which the circuit is mounted: Changes in gene expression are often desired in response to changes in the environment, which typically also affect the cell’s physiological state, for example, a slow-down of growth. Thus, changes in gene expression occur in conjunction with adaptation of the physiology of the cell as a whole. For synthetic gene circuits, such coupling may mostly be an undesired complication [17,18], but for endogenous systems, the regulatory mechanisms have likely evolved to work in such conjunction with the global physiological adaptation of the cell.

While the coupling between physiology and gene expression can be expected to be a general phenomenon, exponentially growing bacteria provide a reasonably simple model system for its quantitative study, because the global state of a cell can to a large extent be characterized by the growth rate as a key parameter [19]. In this article, we review different types of growth-rate dependent gene expression (summarized in Table 1) and discuss the use of growth-rate dependencies as a physiological guide for studying regulatory mechanisms.

## Passive dependence of gene expression on growth-rate

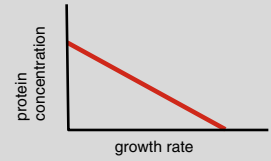
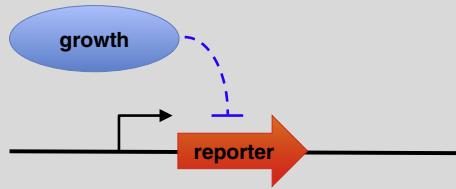
The simplest cases of growth-rate dependence are given by genes, whose products are neutral with respect to growth and present in low abundance, so that negative effects of overexpression on growth are negligible. Specifically, constitutively expressed genes can be used to separate growth-rate dependence from the effects of gene regulation [20,21\*] (Figure 1 and cases 1a, 1b in Table 1). Already in the 1970s, several studies [22,23] showed that the concentrations of the protein product of several constitutively expressed genes in *E. coli* are lower at faster growth. More recently, this dependence was

**Table 1**

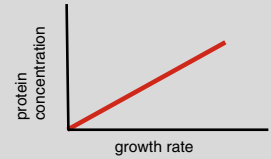
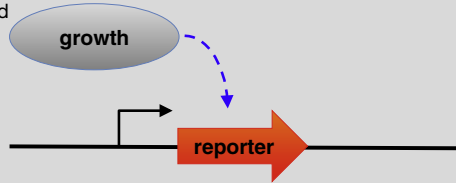
**Types of growth-rate dependencies in gene expression\***

**1. Passive effects due to growth: constitutively expressed genes (gene product neutral, low protein abundance)**

1a. Constitutive expression in nutrient-modulated growth [20,23,26\*]

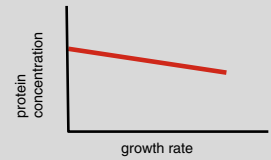
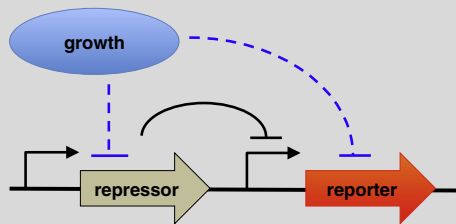


1b. Constitutive expression in translation-modulated growth [12\*]

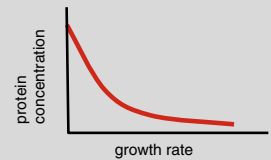
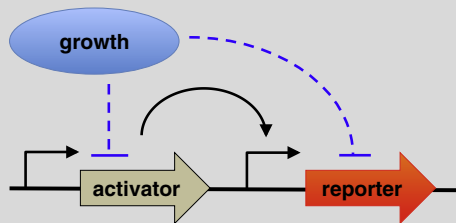


**2. Passive effects due to growth: genes regulated by specific transcription factors**

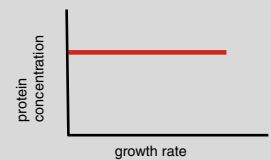
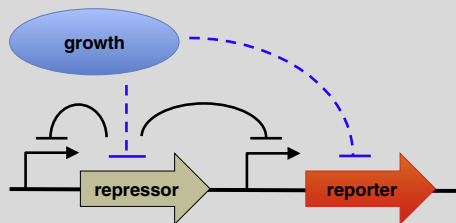
2a. Simple repression (nutrient-modulated growth) [20,26\*]



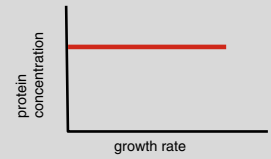
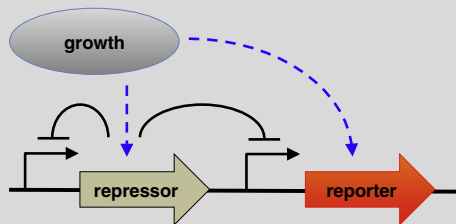
2b. Simple activation (nutrient-modulated growth) [20]



2c. Autorepression (nutrient-modulated growth) [20]



2d. Autorepression (translation-modulated growth) [12\*]



**3. Growth feedback (gene product affects growth)**

3a. Genes with toxic product (nutrient-like growth modulation) [20]

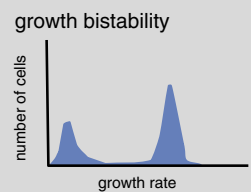
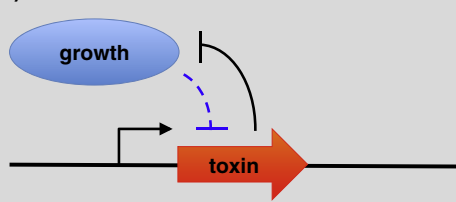
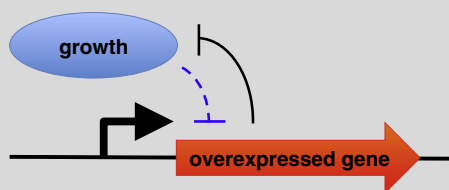
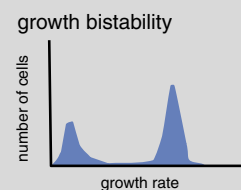
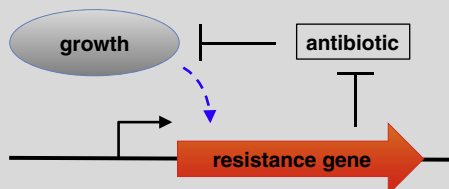


Table 1 (Continued)

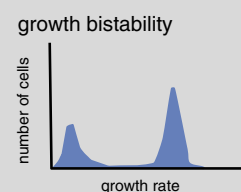
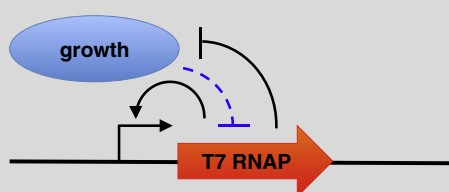
3b. High product abundance/overexpression without specific effects on growth [12\*,57]



3c. Constitutively expressed resistance gene (translation-modulated growth) [36\*]

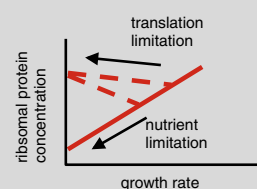
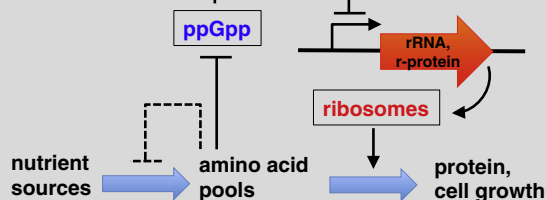


3d. Positive autoregulation with growth feedback (T7 RNA polymerase, mechanism of growth modulation unknown) [37]

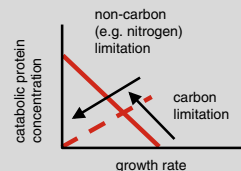
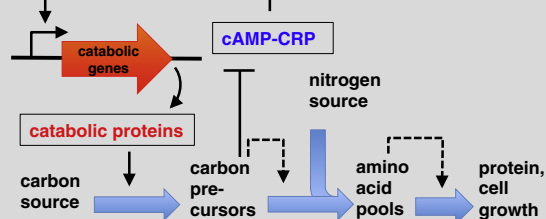


#### 4. Genes controlled by growth-related physiological signals (coordination with metabolic fluxes)

4a. Ribosomal protein genes [12\*]



4b. Genes controlled by catabolite repression [49\*\*]

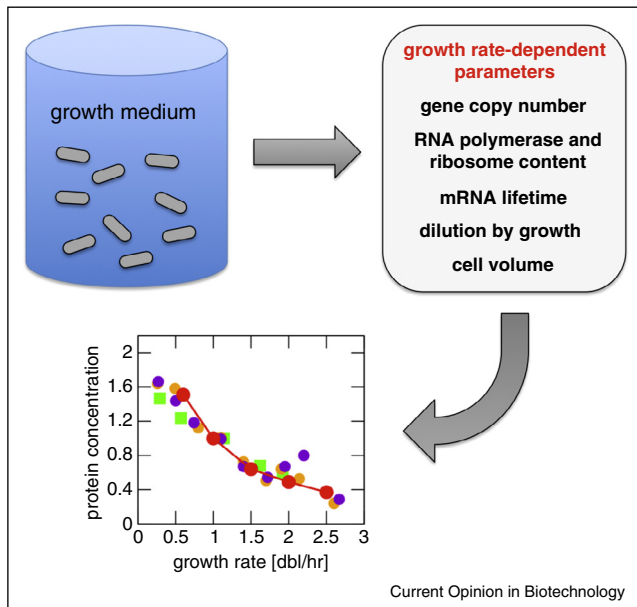


\* In the schematic circuit diagrams, black arrows indicate specific regulation, dashed blue arrows effects of the global state of the cell. Bold blue arrows depict nutrient fluxes.

shown to arise from the combination of the growth-rate dependencies of several cellular parameters, some of which tend to increase expression at faster growth (transcription rate, gene copy number) and some of which decrease it (dilution rate, cell volume) [20,21\*]. Within a phenomenological top-down view based on proteome partitioning this reduction can be interpreted as a consequence of shifting ribosomal activity to making more ribosomal protein and hence generally away from non-ribosomal proteins [12\*,24]; see also the discussion below. For genes controlled by transcriptional regulators, the growth-rate dependence of the regulator levels also contributes to the growth-rate dependent expression [20]. Some examples are listed in Table 1 (cases 2a–2d).

The growth rate dependence is affected by the position of a gene on the chromosome, which determines the timing of its replication. In rapid growth, genes close to the origin of replication are amplified relative to genes close to the replication terminus [13,25]. A recent study has tested and confirmed these position-dependent effects for constitutive and repressed genes at different chromosomal positions (with some additional effects for specific positions and for close proximity of the repressor gene and its target) [26\*]. For genes on plasmids, the growth-rate dependence is affected by the plasmid copy number, which in turn depends on the growth-rate dependencies of parameters of its replication control system [27]. The copy number of some plasmids changes strongly as a

Figure 1



Growth-rate dependence. The global state of a cell, which depends on the growth medium and can often be characterized by the growth rate, influences the quantitative level of gene expression via several parameters. As a result, the concentration of a gene product is typically growth-rate dependent, illustrated here with data for constitutive expression [20].

function of growth rate [28], resulting in strongly growth-rate dependent expression of plasmid-encoded genes [20].

Several groups have explored the coupling between gene regulation and global adaptation further by considering dynamic changes in gene expression during growth transitions [29,30,31]. In contrast to steady state exponential growth, no mechanistic description of the change of microscopic parameters (such as transcription and translation rates, which are modulated by the RNA polymerase and ribosome content of the cell) is currently available for growth transitions and thus it is not straightforward to attribute changes in reporter activity directly to specific molecular mechanisms. However the global effects can still be addressed by comparing the dynamics of the gene of interest to that of a constitutively expressed reference gene. This approach has recently been used to study the slow-down of growth of *E. coli* cells during exhaustion of glucose in the growth medium, where a surprisingly large part of the changes in expression of genes related to carbon metabolism could be attributed to global effects rather than to specific regulation [30]. Another recent study characterized the ‘promoter activity’ (change in reporter fluorescence) as a function of steady-state growth rate and used these dependencies in a model for the dynamics of gene expression during a growth transitions

[31]. Good correspondence with measured data was found for several constitutive promoters and for growth on several different media. We note that this approach requires that the state of a cell during a growth transition can (at least approximately) be described by a continuum of quasi-steady states of growth.

Even though the growth rate is a key parameter of the physiological state of the cell, it is not the only such parameter. Different modes of growth modulation can lead to different growth rate dependencies [12]. All studies discussed so far modulated growth by changing the nutrients in the growth medium, the most common experimental situation. In that case, the concentration of constitutively expressed proteins decreases with increasing growth rate (case 1a in Table 1). For growth modulation by antibiotics targeting the ribosomes, different growth rate dependencies have been observed and, in that case, constitutive expression leads to a protein concentration that increases with increasing growth rate (case 1b in Table 1) [12]. For growth-rate dependencies based on growth-related regulators (discussed below), different modes of growth-rate dependence are obtained for different types of limiting nutrients, for example, carbon vs. nitrogen.

### Growth-mediated feedback

If the product of a gene has a (positive or negative) effect on growth, growth provides a feedback mechanism for the expression of that gene [20]. Whether the feedback is positive or negative depends on the functional form of the growth rate dependence of gene expression (determined by the regulation of that gene) and on the mode of growth-limitation. The effect on growth may be gene-specific (e.g. because the gene product is toxic or because it is required to process a limiting nutrient) or generic (e.g. growth suppression by overexpression), see cases 3a–3d in Table 1.

Positive feedback is obtained for constitutive expression of a toxic protein in nutrient-limited growth (Table 1, case 3a): An increase in the concentration of the toxic protein leads to a reduction in growth rate, which in turn result in a further increase of the toxin concentration (provided that the toxin’s effect modulates growth in similar fashion as nutrient depletion). Bistability arises if positive feedback is sufficiently cooperative and is reflected in bimodal distributions of gene expression levels [32,33]. For growth-mediated feedback, the two subpopulations also exhibit different growth rates. Such feedback may, for example, be induced by expressions of toxins from chromosomal toxin–antitoxin systems. The resulting coexistence of rapidly growing and slow-growing cells is observed in bacterial persistence, phenotypic tolerance to antibiotics [34], where toxin–antitoxin systems indeed play a key role [35]. Another instance of generic growth feedback arises for antibiotic resistance

genes in growth modulated by translation-targeting antibiotics: Increased expression of such a gene reduces the antibiotic level in the cell, thus enhancing growth, which in turn increases the concentration of the gene product (Table 1, case 3c) [36\*].

Growth-mediated feedback may function in conjunction with feedback mediated by transcription factors, effectively making the latter more cooperative [20,37]. This was seen in an autoregulatory T7 RNA polymerase circuit (Table 1, case 3d), where the effective cooperativity induced by growth modulation resulted in bistability in a nominally non-cooperative system [37], as well as in a synthetic positive autoregulation system in yeast [38]. Effective cooperativity is also induced generically when there are several growth feedback systems, because they are coupled through the growth modulation. For example, if one toxin slows down growth, it may enhance expression of other toxins as well. Such toxin cooperativity has recently been proposed [39\*] to explain persister frequencies in strains with deletions of multiple toxin–antitoxin systems [40].

Once a genetic circuit displays growth bistability, its dynamics couples to the population dynamics. While the rate at which cells switch between the two phenotypes is determined by the gene circuit, the percentage of these phenotypes in the population also depends on the two growth rates [38,41]. For example, the fraction of persister cells in an exponentially growing population (in steady-state, i.e. after the high persisters levels left over from the stationary phase have been diluted out) is given by the balance of normal cells outgrowing the persisters and persister generation by (one-way) phenotype switching [41]. Indeed, this switching rate was found to be the best predictor of observed persister fractions, when comparing different *E. coli* strains [42]. A related effect has been observed for a synthetic gene circuit in yeast that is turned on as a memory of DNA damage. In this case, cells with the circuit turned on were found to grow more slowly and thus to be outgrown by the cells with the circuit off [43].

### Growth-rate dependence based on growth-related physiological signals

Finally, growth-rate dependence of gene expression can arise from regulation by growth-related physiological signals that coordinate expression of certain genes with metabolic fluxes. Even though such regulation may be complex, it can lead to surprisingly simple growth-rate dependencies that can be understood based on the economic rationale underlying that regulation. Perhaps the best known such case is the linear growth-rate dependence of ribosomal protein obtained for cells growing in media with different nutrients (case 4a in Table 1) [12\*,19,44,45]. On a microscopic level it depends on the complex mechanisms controlling ribosome synthesis,

with the regulatory nucleotide ppGpp as key physiological signal that adjusts ribosome synthesis to the availability of amino acids [46,47]. From a macroscopic viewpoint, this relation reflects the allocation of ribosomes to making different types of proteins. Specifically, the autocatalytic activity of ribosomes making ribosomal proteins is a requirement for exponential growth that increases with increasing growth rate [12\*,44,48], but must be balanced with the need for making proteins that import and process nutrients. Thus, if the ribosomal capacity is saturated (with almost all ribosomes engaged in translation), the increased ribosome content (proteome fraction) at rapid growth comes at the expense of a reduced content of other proteins, such as the constitutively expressed ones. The observation of a second linear relation under translation inhibition, for example, by antibiotics, has allowed the development of this idea into a quantitative and predictive theory [12\*,24]. A particular success of that theory was the quantitative prediction of the growth reduction due to overexpression of unnecessary protein (case 3c in Table 1).

Another set of strikingly simple growth-rate dependencies is found for catabolic genes under carbon-dependent and non-carbon-dependent growth limitation [49\*\*]: a linear decrease or increase with increasing growth rate, respectively (Table 1, case 4b). These linear dependencies can again be understood as reflecting a trade-off in resource allocation, this time between catabolic proteins on the one hand and biosynthetic proteins (anabolic and ribosomal proteins) on the other hand. Within a quantitative theoretical description [49\*\*], the linear relations can be interpreted as reflecting Pareto surfaces [50,51\*], resulting from the optimization of competing requirements. The physiological signal underlying these growth-rate dependencies is another regulatory nucleotide, cAMP (in a complex with the transcription factor CRP), long known as the effector of catabolite repression [52]. The central physiological role of cAMP-CRP as a coordinator of carbon metabolism with competing metabolic demands has, however, only been identified based on the quantitative analysis of these growth-rate dependencies [49\*\*]. Moreover, via the specific case of coordination between carbon and nitrogen metabolism, this analysis also suggested that feedback by precursors at the interface between carbon and nitrogen metabolism may provide the molecular implementation of that coordination. This hypothesis was subsequently confirmed and  $\alpha$ -ketoacids (including  $\alpha$ -ketoglutarate, which had previously been shown to control the uptake of both carbon and nitrogen sources [53,54]) were identified as the physiological trigger for cAMP-CRP regulation [49\*\*], thus playing the role of the unknown ‘catabolites’ in Magasanik’s original catabolite repression hypothesis [55]. This discovery of a molecular link that had been elusive for decades provides an example for how the quantitative study of physiological

growth-rate dependencies can guide the understanding of regulatory strategies and their molecular implementation [49,56].

In summary, gene expression and cell growth are inter-related in multiple ways. An understanding of these relations can improve the quantitative understanding and prediction of gene circuit behavior and provide guidance towards understanding the economic principles as well as the molecular implementation of physiological regulation.

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