Cooperative Adaptation to Establishment of a Synthetic Bacterial Mutualism

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Abstract

To understand how two organisms that have not previously been in contact can establish mutualism, it is first necessary to examine the temporal changes in their phenotypes during the establishment of mutualism. Instead of tracing back the history of known, well-established, natural mutualisms, we experimentally simulated the development of mutualism using two genetically engineered auxotrophic strains of Escherichia coli, which mimic two organisms that have never met before but later establish mutualism. In the development of this synthetic mutualism, one strain, approximately 10 hours after meeting the partner strain, started oversupplying a metabolite essential for the partner's growth, eventually leading to the cooperative phenotype of the partner only after encountering the partner strain before the growth of the strain itself. By transcriptome analysis, we found that the cooperative phenotype of the strain was not accompanied by the local activation of the biosynthesis and transport of the oversupplied metabolite but rather by the global activation of anabolic metabolism. This study demonstrates that an organism has the potential to adapt its phenotype after the first encounter with another organism to establish mutualism before its extinction. As diverse organisms inevitably encounter each other in nature, this potential would play an important role in the establishment of nascent mutualism in nature.

Introduction

Mutualism is based on a mutually beneficial interaction between two organisms and is ubiquitous in nature [1,2,3,4,5,6]. Mutualisms observed in nature are thought to be the result of adaptation of each organism to the existence of the partner after their first encounter. The genetic origin and trajectory of this adaptation has been investigated via a phylogenetic approach [2,7,8]. However, tracing back established mutualisms to their origin is challenging as no intermittent states are defined with which to measure the adaptation in terms of phenotypic traits, population size and local environment [1]. To investigate the environmental conditions required to establish a nascent mutualism, one study reported a synthetically designed mutualism using two species of bacteria [9]. The findings of that study clearly demonstrated the importance of spatially structured environments for the establishment of mutualism, providing proof of principle of natural selection of cooperative behavior that has been proposed by the theoretical studies [10,11,12,13]. These types of experimental studies using microbial ecosystems to test the theories of cooperative systems have recently been reported [14,15,16,17,18]. Most of these studies focused not on the adaptation of the organisms but on the environmental conditions required for the persistence of cooperative behavior in natural selection.

Some studies have characterized the behavior of organisms in nascent mutualisms. Wintemute et al. synthetically designed mutualisms comprising certain pairs of auxotrophs of Escherichia coli and found significant metabolic synergy in 17% of 1035 such pairs tested [19], although it was unclear if any adaptation of the bacteria contributed. Shou et al. synthetically designed an obligate mutualism composed of two yeast auxotrophs [20], each of which was genetically engineered to overproduce the metabolite essential for the growth of the partner. Both of the auxotrophic strains grew to saturation without the need for external supplementation of their essential metabolites compensating for the auxotrophy. Moreover, they showed adaptation in as little as one hundred generations, where they became capable of growing from diluted cell densities or ceased growth due to weakening of the beneficial interaction. Hillesland et al. demonstrated that the growth rate of microorganisms in another synthetic mutualism increased after serial passage, even in the absence of spatially structured environment, while the extent of the adaptation was increased in a spatially structured environment [21]. These adaptations of microorganisms occurred after the establishment of nascent mutualisms, strengthening their interactions.

Can adaptation occur before the establishment of a nascent mutualism, leading to its establishment? Here we show that a strain of bacteria became more beneficial to another strain before...
their population started growing and establishing a nascent mutualism. Specifically, we synthetically designed an obligate mutualism comprising two auxotrophs of Escherichia coli. We show that one of the two auxotrophs, upon encountering the partner strain before their own population growth, adapted by oversupplying the metabolite essential for growth of the partner, which in turn permitted its own growth, leading to the successive growth of both strains. This study therefore shows the potential of organisms to adaptively respond to the first encounter with another organism, which could lead to the establishment of nascent mutualisms. As diverse organisms inevitably encounter each other in nature, this potential would play an important role in the establishment of nascent mutualisms in nature.

Results

To create a synthetic model of obligate mutualism, we constructed two different types of nutrient auxotrophs of E. coli by genetic recombination (Fig. 1A): an isoleucine (Ile) auxotroph, designated I, labeled with a red-fluorescent protein (dsred.T3), and a leucine (Leu) auxotroph, L, labeled with a green-fluorescent protein (gfpuv5) [22] (see Methods). We were able to distinguish these two strains by flow cytometry (FCM). In minimal medium without amino acid supplements, neither strain was able to grow in monoculture. However, in coculture, if the two strains supplied a sufficient amount of the essential amino acids required by the other strain, they would successively grow and thereby establish mutualism.

We measured the supply of amino acids from each strain in monoculture to test whether the quantities were sufficient for the successive growth of both strains in coculture. Figs. 1B and 1C show the supply of Leu from I cells in monoculture, with and without the addition of 10 μM Ile, respectively, and Figs. 1D and 1E show the supply of Ile from L cells in monoculture, with and without the addition of 10 μM Leu, respectively. Before inoculation of these cultures, we washed each strain with minimal media not containing amino acids to exclude the carry-over of supplements from preculture (see Methods). Obviously, both strains did not grow without the addition of amino acids (Fig. 1C and 1E). We measured the concentrations of Leu and Ile in the culture media using a bioassay (see Methods), and expressed these as the cell concentration of L and I cells which can be produced by the amount of amino acid supplied, respectively. In every case (Fig. 1B–E), the final concentrations of an amino acid in the recipient cell were always less than the maximum concentrations of the donor cell. That is, the nutrient supply from the donor cells was insufficient to produce an equal amount of nutrients in the recipient cells, and was therefore insufficient to sustain the net growth of both strains [20]. These results implied that any adaptation to the mutualism, such as an increase in the nutrient supply, needs to occur in coculture for the successive growth of both strains.

Despite the insufficient amino acid supply in monoculture, both strains grew to saturation (around 10^6 to 10^7 cells/ml) in coculture (Fig. 1F). Initially, I cells grew (red • at <10 h), followed by L cells (green ▲ at ~20 h). Qualitatively, the initial growth of I cells was consistent with the results of amino acid supplementation in monoculture as follows. In monoculture, I cells supplied Leu only after growth and the uptake of Ile (Fig. 1B and 1C), while L cells supplied Ile regardless of growth (Fig. 1D and 1E). These results suggested that initially L cells supplied Ile promoting the growth of I cells. In addition to the initial growth of I cells, the amount of Leu was detected at time 0 in coculture, as shown in Fig. 1F (blue +). As I cells supplied Leu only when they consumed Ile, these results indicated that I cells consumed Ile supplied from L cells and then supplied Leu just after mixing but prior to sampling. However, quantitatively, the initial growth of I cells was inconsistent with the results from monoculture. I cells grew to greater than twice the concentration of L cells (Fig. 1F, red • at <10 h). That is, the Ile supply from L cells was sufficiently high to produce a greater concentration of I cells than L cells, which was different from the results of monoculture described above (Fig. 1D and 1E). The final concentration of Ile was also significantly higher than that of L cells in coculture (Fig. 1F, orange ▲). The inconsistency in the quantity of Ile supplied by L cells in coculture and monoculture suggested the enhanced supply of Ile from L cells on encountering I cells. It should be noted that L cells did not show significant growth before nine hours in coculture when the Ile supply from L cells already appeared to be enhanced (Fig. 1F), which indicates that enhancement of the Ile supply from L cells did not require the population growth of strain L itself. Also, enhancement of the Leu supply from I cells was detected in coculture (Fig. 1F), as discussed later.

We investigated the growth kinetics of the cocultures at various initial cell concentrations of strains I and L (Fig. 2). The cells entered stationary phase at around 20–30, 40–120, and 300–600 h when the initial cell concentration of strain L was 10^4, 10^5, and 10^6/ml, respectively. Cell growth was not observed when the initial cell concentration of strain L was 10^3/ml. On the other
hand, clear dependency on the initial cell concentration of strain \( I^- \) was not observed. The difference in the dependencies on the initial concentration of \( I^- \) and \( L^- \) cells was consistent with the differences in the features of nutrient supply found in the monocultures; only \( L^- \) cells supplied Ile even in the absence of amino acids in monoculture as described above (Fig. 1E). These results suggested that \( L^- \) cells initiated the first steps towards establishing mutualism in coculture.

The time courses of the cocultures also showed another feature. In some cases, the initial growth of \( I^- \) cells reached a concentration of up to approximately 10-fold greater than that of \( L^- \) cells (Fig. 2, depicted by arrows). This suggested that the Ile supply from \( L^- \) cells was insufficient to produce this concentration of \( I^- \) cells. As the Ile supply from \( L^- \) cells was insufficient to produce an \( I^- \) cell concentration equal to that in monoculture, which was equivalent to the initial lag time in coculture, the sufficient supply of Ile from \( L^- \) cells in coculture suggested that \( L^- \) cells changed to a "high supplier" phenotype prior to growth. Indeed, a mathematical model assuming the change in \( L^- \) cell phenotype could explain the time course of coculture (Fig. S1). It should also be noted that the lag period was dependent on the initial cell concentration of \( L^- \) cells in the coculture, which suggested that the interactions between cells were required for the change in \( L^- \) cell phenotype.

To directly observe the change in \( L^- \) cells to a high supplier phenotype, we tested reconstituted cocultures (re-coculture) using \( L^- \) cells prepared from mid-coculture (Fig. 3). To prepare \( L^- \) cells separately from \( I^- \) cells from mid-coculture, we inoculated each strain into media separated by a membrane, which was permeable to amino acids but not to \( E. \ coli \) (membrane coculture) (see Methods). The time course of the membrane coculture was almost the same as that of coculture without membrane separation (Fig. 3A). For the re-coculture, we used both strains harvested from three different culture conditions: (i) at the log phase in monoculture with the addition of the required amino acids, which is the same as the initial state (0 h) of membrane coculture \( I^-_{\text{ini}} \) and \( L^-_{\text{ini}} \), (ii) at 23 h of membrane coculture, when \( I^- \) cells had grown to a concentration approximately 10-fold higher than that of the \( L^- \) cells (Fig. 3A) \( I^-_{\text{co}} \) and \( L^-_{\text{coi}} \), and (iii) at 23 h of membrane monoculture in the absence of amino acids, when both strains were not growing \( I^-_{\text{mono}} \) and \( L^-_{\text{mono}} \). Before inoculation of the re-cocultures, we washed each strain with minimal media to exclude supplements carried over from the first membrane cultures. Fig. 3B–E show the time courses of the re-cocultures comprising \( I^-_{\text{ini}} \) and \( L^-_{\text{ini}} \), \( I^-_{\text{co}} \) and \( L^-_{\text{coi}} \), \( I^-_{\text{ini}} \) and \( L^-_{\text{ini}} \), \( I^-_{\text{co}} \) and \( L^-_{\text{co}} \), and \( I^-_{\text{ini}} \) and \( L^-_{\text{ini}} \), respectively. Only the re-coculture containing \( L^-_{\text{co}} \) cells showed initial growth of \( I^- \) cells without a lag phase (Fig. 3D and 3E, arrows). These results indicated that \( L^-_{\text{co}} \) cells were high suppliers of Ile at time 0 in the re-coculture, in contrast to \( L^-_{\text{ini}} \) cells. It is worth noting that \( L^-_{\text{ini}} \) cells represent the initial state of \( L^-_{\text{co}} \) cells in the first membrane coculture, i.e., \( L^- \) cells change to a high supplier phenotype in the first membrane coculture. Recoculture containing \( L^-_{\text{mono}} \) cells exhibited a lag phase before the initial growth of \( I^- \) cells (Fig. 3F and 3G). These results indicated that \( L^-_{\text{mono}} \) cells were not high suppliers, like \( L^-_{\text{co}} \) cells, and the change to a high supplier phenotype was dependent on coculturing. A significant change in \( I^- \) cells was not detected (Fig. 3B and 3C) in the re-coculture, although the oversupply of Leu from strain \( I^- \) was observed in Fig. 1F. These results experimentally confirmed that \( L^- \) cells changed to a high supplier phenotype in coculture prior to their own growth.

These findings raised the question: how does gene expression change in the two strains during coculture? To investigate this, we carried out a comprehensive analysis of gene expression in these two strains. To harvest each strain separately from coculture, we again employed membrane coculture. Using a DNA microarray, we measured and compared the expression intensities of all 4345 genes.
genes of each strain cultured under three different conditions. These three conditions were: (i) at the log phase in monoculture, which is the same as the initial state (0 h) of membrane coculture, as described above (I ini and L ini); (ii) at the stationary phase (45 h) of membrane coculture (I st,co and L st,co), and (iii) at the stationary phase (45 h) of monoculture after growth in the presence of the required amino acids (I st,mo and L st,mo). As E. coli is known to substantially change its gene expression depending on the growth phase [23], samples were taken at 45 h (not 23 h) to identify coculture-specific changes by comparing samples at the same phase (stationary phase). In the I strain, the changes in gene expression from I ini to I st,co, strongly correlated with those from I ini to I st,mo (Fig. 4A), that is, the dominant changes were dependent on the growth phase. This correlation was also observed in L cells and the slope of linear regression was smaller than that of I cells (Fig. 4B), which may have been because the time after entering stationary phase was shorter in L st,co cells than in I st,co cells (Fig. 3A). More importantly, in strain L, the correlation coefficient was smaller than that in strain I (Fig. 4B). These results indicated that the change in gene expression of L cells in coculture was more coculture-specific than that of I cells.

The next question raised was: which categories of genes were involved in the coculture-specific changes in expression in L cells shown in Fig. 4B? Initially, we focused on genes that showed significantly induced or repressed expression in coculture compared to monoculture, i.e., those in which the ratio of gene expression in L st,co to that in L st,mo cells was greater than three or lower than one-third. We statistically screened the up-regulated and down-regulated gene categories to which the significantly induced or repressed genes belonged (Table 1). For the gene categories, we adopted the “cellular processes” category in the Gene Ontology (GO) database [24], and the categories for genes regulated by sigma factors in the database, RegulonDB [25]. In the “cellular processes” category of the GO database, 14 categories out of 124 were found to be up-regulated, and most of these up-regulated categories were related to anabolism, such as the biosynthesis of amino acids (tryptophan, proline, methionine, phenylalanine, leucine, cysteine, and chorismate, which is a precursor of tyrosine, phenylalanine and tryptophan), polyamines and proteins. In contrast, of the nine down-regulated categories, most were related to catabolism, such as the various energy cycles (glyoxylate and tricarboxylic acid cycles), fatty acid oxidation, and the catabolism of amino acids, aminobutyrate and carbohydrates. Some of these up- and down-regulated categories were also identified when comparing L st,co and L ini cells (Table 1, indicated by arrows). Although L cells over-supplied Ile in coculture, no significant increase was found in the expression of genes related to Ile biosynthesis or Ile transport in L st,co cells compared with both L st,mo and L ini cells (Fig. S2). It is worth noting that the results of liquid chromatography showed that the predominant supplement from L cells required by I cells was Ile (Fig. S3). Among the genes regulated by sigma factors, we detected the down-regulation of genes regulated by Sigma 70, the housekeeping sigma factor [26], during the change from L ini to L st,co cells (arrows, Table 1) and the change from L ini to L st,mo cells. These results were consistent with the change in growth phase to stationary phase. Although the down-regulation of Sigma 70 genes occurred in both L st,co and L st,mo cells, the gene expression significantly differed between L st,co and L st,mo cells (Fig. 4B). Therefore both up- and down-regulation was found when comparing L st,co and L st,mo cells. Down-regulation of the glutamine biosynthesis gene category correlated with the down-regulation by Sigma 54, the sigma factor controlling nitrogen usage [26,27]. As glutamine biosynthesis opposes glutamate biosynthesis leading to the biosynthesis of other amino acids, the down-regulation of glutamine biosynthesis is not inconsistent with the up-regulation of the anabolic categories. As above, we found that the coculture-specific changes in gene expression in L cells were not related to the local activation of the biosynthesis and transport of isoleucine, but were related to the global activation of anabolic metabolism.
Table 1. The categories of up-regulated and down-regulated genes in L\textsubscript{st,co} cells.

| Database  | Up-regulated categories | Down-regulated categories |
|-----------|-------------------------|---------------------------|
| GO        | 0000162 tryptophan biosynthesis ↑ | 0006071 glycerol metabolism |
| Cellular  | 0000270 peptidoglycan metabolism | 0006097 glyoxylate cycle ↓ |
| processes | 0006310 DNA recombination | 0006099 tricarboxylic acid cycle ↓ |
|           | 0006412 protein biosynthesis | 0006542 glutamine biosynthesis ↓ |
|           | 0006561 proline biosynthesis | 0009063 amino acid catabolism ↓ |
|           | 0006596 polyamine biosynthesis | 0009450 aminobutyrate catabolism |
|           | 0006790 sulfur metabolism | 0016052 carbohydrate catabolism ↓ |
|           | 0009086 methionine biosynthesis ↑ | 0019395 fatty acid oxidation |
|           | 0009094 L-phenylalanine biosynthesis | 0042594 response to starvation ↑ |
|           | 0009098 leucine biosynthesis (except leuB) ↑ | |
|           | 0009243 O antigen biosynthesis | |
|           | 0009257 10-formyltetrahydrofolate biosynthesis | |
|           | 0009423 chorismate biosynthesis ↑ | |
|           | 0019344 cysteine biosynthesis | |
| Sigma     | Sigma 70 ↓ | Sigma 54 ↓ |
| factors   | Sigma 70 ↓ | |

The categories were screened by comparing between L\textsubscript{st,co} and L\textsubscript{st,mo} cells as coculture-specific changes. In the screened categories, upward and downward arrows (↑ or ↓) at the end of the category name represent up-regulated or down-regulated categories, respectively, in L\textsubscript{st,co} cells relative to L\textsubscript{st,mo} cells.

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Discussion

In our synthetic model of obligate mutualism comprising two auxotrophs of E. coli, strains I\textsuperscript{−} and L\textsuperscript{−}, the increase in the Ile supply from L\textsuperscript{−} cells occurred before the population growth of L\textsuperscript{−} cells, and both strains grew successively thereafter in coculture. We found that the increase in the Ile supply from L\textsuperscript{−} cells depended on coculture with I\textsuperscript{−} cells and was accompanied by coculture-specific changes in the gene expression of L\textsuperscript{−} cells. This change in L\textsuperscript{−} cells in coculture was not related to the local activation of the biosynthesis and transport of isoleucine, but was related to the global activation of anabolic metabolism.

What is the mechanism behind the phenotypic change in L\textsuperscript{−} cells to become “high suppliers” of isoleucine? There are two possibilities: (i) a fraction of high suppliers preexists in the initial population and their fraction in the L\textsuperscript{−} population increased in the coculture (natural selection), (ii) L\textsuperscript{−} cells changed their phenotypes in response to the changes in the environment from monoculture to coculture (phenotypic plasticity). We can rule out neither possibility completely.

Let us first assume that (i) is true, and estimate an approximate range of the fraction of preexisting high suppliers in the initial population (f\textsubscript{f0}). At first, the supply of Ile from high suppliers was about 10-fold higher than that of “normal” L\textsuperscript{−} cells. Therefore, f\textsubscript{f0} would be less than 10%, otherwise the supply of Ile from L\textsuperscript{−} cells at the initial state or in the monoculture would be higher than the experimental results. Second, the shortest time until the initial growth of I\textsuperscript{−} cells was about 10 hours. By then the high suppliers had already been a majority in the L\textsuperscript{−} population and their concentration was approximately 10\textsuperscript{7}/ml (Fig. 1F). For the concentration of the high suppliers to become 10\textsuperscript{7}/ml in 10 hours from the initial concentration 10\textsuperscript{7}/f\textsubscript{f0}/ml, f\textsubscript{f0} \geq e^{-10gH} must be satisfied, where g\textsubscript{H} is the growth rate of the high suppliers. Even if only high suppliers grew at a maximum growth rate of L\textsuperscript{−} cells (g\textsubscript{H} = 0.4/h, Table S1), f\textsubscript{f0} \geq 2% was required. f\textsubscript{f0} thus can be estimated as 2% < f\textsubscript{f0} < 10%. Note that there is no reason why only high suppliers grew at a maximum rate in the environment, where the normal L\textsuperscript{−} cells did not grow. In the mixed liquid culture, all of the L\textsuperscript{−} cells were considered to acquire Leu from the media (not from I\textsuperscript{−} cells directly) in a homologous environment (actually, physical contacts were negligible; Fig. 3A). Moreover, f\textsubscript{f0} must be kept in this range in monoculture because the time until the initial growth of I\textsuperscript{−} cells were reproducible even when we used another clone of L\textsuperscript{−} cell for the preparation of the initial population.

We then discuss about the possibility (ii). E. coli is known to alter its phenotype in response to environmental changes, such as amino acid starvation, and this is known as a stringent response [28] and represents a kind of phenotypic plasticity. As L\textsuperscript{−} cells were subject to Leu starvation at the initial in coculture, they would have changed their phenotype as the stringent response. This response might have been preserved even after their growth in which they had already been released from Leu starvation. Indeed, the up-regulation of amino acid synthesis, which is known to occur in the stringent response [28], was observed in L\textsubscript{st,co} relative to L\textsubscript{st,mo} (Table 1). However, although L\textsuperscript{−} cells in monoculture without Leu (L\textsubscript{monom}) were subject to Leu starvation, they did not change to the high supplier phenotype (Fig. 3F and 3G). In our experiments, L\textsuperscript{−} cells changed to a high supplier phenotype only in coculture, and the genes related to Ile biosynthesis and transport were not significantly induced in these cells (Fig. S2), in contrast, these genes are induced during the stringent response [28]. It is known that Ile uptake is increased and amino acid permeability is decreased during the stringent response [28], which seems to oppose the extracellular leakage of Ile. Our results might therefore indicate that the phenotypic change in L\textsuperscript{−} cells was related not only to the known stringent responses, but also to other responses due to the interaction among strains via the media. As both the I\textsuperscript{−} and L\textsuperscript{−} strains were constructed by a single gene deletion from the same original strain, DH1 [see Methods], the substances supplied by them via cell leakage would be expected to be almost the same. Thus, the interaction between these strains is unlikely due to the expression of a specific substance, as is the case in quorum sensing [29], but is
more likely due to a global change of the composition of multiple substances [30,31]. This might be consistent with the observed global activation of expression of genes involved in anabolic metabolism (Table 1). It is worth noting that a similar phenomenon was observed in a synthetic mutualism comprising $\Gamma$ cells and an uracil auxotroph (Fig. S4), therefore, the change in L– cell phenotype is not due to the similarities between the metabolism of Ile and Leu. Further studies are required to fully elucidate the mechanism behind the phenotypic change in strain L–.

It is unknown to what extent such an adaptation to a first encounter contributes to the establishment of a nascent mutualism in nature. However, this potential would provide insight into the positive factors required for the establishment of natural mutualisms. We do not believe that an adaptation, such as that described in this study, results in the establishment of nascent mutualism in every case, because we actually failed to establish mutualism with some combinations of auxotrophs (such as a glutamine auxotroph and an uracil auxotroph). Synthetic mutualism also failed to be established in other organisms without the introduction of metabolite-overproducing mutations [9,20]. However, due to the great variety of organisms in nature [3,32], organisms inevitably encounter other kinds of organisms and have the opportunity to establish a nascent mutualism, such an adaptation to this first encounter can facilitate this process. It is worth noting that such an adaptation to first encounter might be a kind of the phenotypic plasticity in response to a new environment [33,34,35,36,37,38,39,40], and the adaptation in response to a first encounter might have evolved because the organisms possessing this potential should survive in the bio-diversified nature. Field studies are required to fully investigate the contribution of such an adaptation to the establishment of nascent mutualism in nature.

The simplicity of the synthetic model of mutualism used in this study enabled us to identify unexpected and quantitative changes in the organisms. Experimental ecosystems not only provide empirical proof of theories but also highlight unexpected phenomena, such as the unknown potential of organisms which may lead to novel theories. For example, in another bacterial system, Fiegna et al. found that a single point mutation changed a cheater into a cooperator with a tolerance to exploitation by the cheater [41,42]. Without the simplicity of the system, it would have been impossible to detect such a phenomenon. For future studies, synthetically-constructed experimental ecosystems combining naturally non-interacting species [21,43,44,45,46] and recombining interactions using genetic modifications [16,19,20,47,48,49,50,51,52,53], would be invaluable for the empirical proof of theories but also highlight unexpected phenomena.

The simplicity of our synthetic model of mutualism will enable further studies to experimentally resolve some of the remaining questions, such as the molecular mechanisms behind the observed adaptation in L– cells and the evolutionary pathway of this mutualism. Our findings may also contribute to the study of adaptation in response to a new environment [33,34,35,36,37,38,39,40], and the adaptation to this first encounter can facilitate this process. It is worth noting that such an adaptation to first encounter might be a kind of the phenotypic plasticity in response to a new environment [33,34,35,36,37,38,39,40], and the adaptation in response to a first encounter might have evolved because the organisms possessing this potential should survive in the bio-diversified nature. Field studies are required to fully investigate the contribution of such an adaptation to the establishment of nascent mutualism in nature.

Culture conditions

All cultures were grown at 37°C in well-mixed minimal media modified with M63 (pH 7.0, 62 mM K2HPO4, 39 mM KH2PO4, 15 mM ammonium sulfate, 1.6 mM FeSO4-7H2O, 15 μM thiamine hydrochloride, 0.2 mM MgSO4-7H2O and 22 mM glucose; mM63 [36]). Amino acids were added to the media when appropriate. Before culturing, we washed E. coli strains with the minimal media without amino acids to exclude the carry-over of supplements from preculture. For the membrane culture, we used cell culture inserts with a pore size of 0.45 μm at a density of 106/cm2, and used six-well cell culture companion plates for the inserts (BD Falcon, Franklin Lakes, NJ, USA). The initial concentration of $\Gamma$ and L– cells are depicted at time 0 in the figures or described in the figure legends.

Measurement of cell concentrations

We measured the cell concentration relative to a known concentration of fluorescent beads (Fluospheres YG Microspheres, 5 μm; Polysciences Inc., Warrington, PA, USA) using a Cytomics TM FC500 Flow Cytometer (Beckman Coulter, Inc., CA, USA) by loading culture samples mixed with the beads. A 488 nm argon excitation laser was employed and band-pass filters of 515–535 and 610–630 nm were used to measure green and red fluorescence, respectively. Clusters of red and green cells, and the fluorescent beads, were clearly segregated (Fig. S5), and each cell concentration was calculated from these counts.

Measurement of amino acid concentrations using a bioassay

To measure the Ile concentration of a culture, the culture was passed through a 0.2 μm filter and the supernatant was supplemented with a one-sixth volume of the mM63 media and inoculated with I– cells at 105/ml. Then the Ile concentration was obtained by multiplying the saturation concentration of I– cells (≥18 h) by six. The Leu concentration of a culture was obtained using the same method for strain L–. It is worth noting that when Ile was added to the monoculture of I– cells in mM63 media, the concentration of added Ile and the saturation concentration of I– cells was proportional, with a constant of 9.8 (cells/mll/μM).

The DNA fragment was amplified by PCR using the template, pPROTe.E333-tetT, in which the dsrT3 fragment (in the presence of the plasmid pPROTe.E333-lacZ, Clontech, using the following primers, recativeXρr (5’-AACAAATCTGGCCTGAGCCTAGCCCGCCCTGCCACT-3’) and ilvE-T3act-r (5’-TAAATGGGAGCCGTGGTGTCCGTCCTTTTTTGTTTATATTACG-AGGAACAGGTGG-3’). Homologous recombination was performed as described previously [55,56]. The E. coli strain DHIΔildB::gfprv5-Kmr, designated L–, was constructed from the E. coli strain DH1, by replacing the chromosomal leuB::gfprv5-Kmr, comprising a reporter gene (gfprv5) and the kanamycin resistance gene (Kmr). The DNA fragment was amplified by PCR using the template, pGAG-2 [57] and primers, leuB-kanIG-f (5’-GCTCACAAGCGAAAGCA-CAAGGAAACCGTGTGATTAGAAAAACTCATCGAGCA-3’), and leuB-Gkan-r (5’-CGTGGACAATTTTCTGATATC-GTCTTAGCCATGAATTATCGTACGCTCA-3’).

Materials and Methods

Construction of E. coli strains

The E. coli strain DHIΔildB::dsrT3-catI, designated $\Gamma$, was constructed from the E. coli strain DH1 (National Bioresource Project, National Institute of Genetics, Shizuoka, Japan), by replacing the chromosomal ilvE::gene with a foreign DNA fragment, P_{all}::dsrT3-P_{all}catI, comprised of a reporter gene (dsrT3) and the chloramphenicol resistance gene (cat). The dsrT3 gene and the cat gene were transcribed from their promoters in opposite directions.
Gene expression analysis

E. coli gene expression was examined using a GeneChip® E. coli Genome Antisense Genome Array according to the Expression Analysis Technical Manual (Affymetrix, 2004). The expression analyses of co-culture samples were performed with two technical replicates using two different target cDNAs separately prepared for each sample. The expression level of each gene was computed according to the FH model [58]. The estimated expression levels were normalized using a quantile normalization method [59]. For the analysis of the gene categories, we used three as the threshold for the ratio of gene expression to determine whether the expression of a gene had changed. When we calculated the ratio of gene expression for each of the 4345 genes between the two replicates of L_{st,co} cells in individual co-cultures, the ratios were less than three for 90% of genes. To screen the categories that were significantly up- or down-regulated, we used a one-side binomial test at the significance level of 0.01.

Supporting Information

Figure S1 Mathematical model of the growth kinetics of the coculture. (A) Mathematical model of the growth kinetics. [X] indicates the concentration in the culture media of X. We defined active cells of strains Γ and L’ as Γ_{act} and L’_{act}, respectively, for the following reason. When we measured the cell concentration as colony forming units (cfu) under starvation conditions, the concentration determined by cfu was less than the concentration determined by analysis of fluorescent particles by flow cytometry (Fig. S5B and S5C). Although it was difficult to determine whether cells were alive or dead, we defined a cell being able to form a single colony as an active cell. This model neglects the decrease in [Γ] and [L’] because it was slow (Fig. S5B and S5C). The symbol S represents glucose as a carbon source in the minimal media, which only determines the saturation concentration (we set 10^7/ml for the simulation in Fig. S1B). The explanations and the values of the parameters are shown in Table S1. This model is based on the Monod model with the maintenance rate [60]. The specific character of this model is the heterogeneity of the supply function of the amino acid between Γ and L’ cells, as experimentally shown in Fig. 1. A mathematical model assuming these two types of nutrient supply has been reported for another obligate mutualism comprising two bacteria isolated from soil microcosms [61]. As Γ_{act} cells supplied Leu only after growth, αI was defined as the number of L’_{act} cells produced in the presence of Leu from a single new Γ_{act} cell until its death in the culture. In this model, a mathematically solved necessary condition for the stable growth of both strains is αI/K_l = M > 1. K_l/M represents the number of Γ_{act} cells produced in the presence of Ile from a single new L’_{act} cell until its death in the culture. In our experiments, K_l/M was less than one in monoculture (Fig. 1D and 1E) and at the lag phase in coculture (Fig. 2 and 3), but was nearly 10 after the lag phase in coculture (Fig. 2 and 3). (B) Comparison between the simulation results of the model and the experimental results shown in Fig. 1F. The simulation also roughly fit to the experimental results regarding the amino acid concentrations.

Figure S2 The change of the expression of genes related to Ile biosynthesis and transport in L’ cells. The black and gray bars show the ratio of the expression of each gene in L’_{st,co} and L’_{st,mo} cells compared to that in L’_{ini} cells, respectively, where L’_{st,co}, L’_{st,mo}, and L’_{ini} represent the state of L’ cells at the stationary phase of coculture, the stationary phase of monoculture and the growth phase as their common initial state, respectively (see text for details). Genes related to Ile biosynthesis and transport are depicted as “Biosynthesis” and “Transport,” respectively, under their gene names. None of the Ile biosynthesis-related genes were induced in L’_{st,co} cells compared with L’_{ini} cells or in L’_{st,co} cells compared with L’_{st,mo} cells, using three-fold (red line) as the significant threshold (see Methods). None of the Ile transport-related genes were significantly changed between any two of the three conditions.

Figure S3 The ratio of the concentration of amino acids determined by HPLC to those determined by a bioassay. We determined the quantity of Ile and Leu in the supernatants of the cultures in which each strain reached saturation phase in the presence of the required amino acid (1 μM Ile or 1 μM Leu for Γ or L’ cells, respectively) using two different methods: HPLC and a bioassay. The results indicated that the supplied nutrient from L’ cells that compensated for the Ile auxotrophy of Γ cells consisted mainly of Ile, while the supplied nutrient from Γ cells that compensated for the Leu auxotrophy of L’ cells consisted of substances other than Leu. Methods for the bioassay are described in the text and the methods for HPLC are described below. We added Ile and Leu to the supernatants (both 0.05 μM) to raise the concentrations in the supernatants above the detection range of HPLC. As an internal standard, norleucine was also added to 0.25 μM in the supernatant. The resultant solutions were derivatized by phenylisothiocyanate (Wako, Osaka, Japan) and applied to a reverse phase HPLC on a Waters LC Module 1 (Waters Corporation, MA, USA) with a column of Wakosil-PTC (4.0×250 mm, Wako, Osaka, Japan). The column was soaked in a circulating water bath at 40°C. The mobile phase comprised 60 mM sodium acetate (pH 6.0) and acetonitrile (94:6) as eluant A; eluant B consisted of 60 mM sodium acetate (pH 6.0) and acetonitrile (40:60). Gradient elution was employed according to the following linear program: time 0, 0% eluant B; 20 min, 70% eluant B; 21 min, 100% eluant B. The flow rate was 1 ml/min. Amino acid derivatives were detected by their absorbance at 254 nm.

Figure S4 Basic design and cell growth of the synthetic mutualism comprising Γ cells and a Ura auxotroph (U’). (A) Schematic diagram of the synthetic mutualism. Two auxotrophs of E. coli, strains Γ and U’, supply nutrients to each other to form a potential mutualism. (B-E) Cell growth and nutrient release properties of the monocultures. The concentration of Ura was determined by a bioassay, as was the concentration of Ile and Leu. The concentration of Ura or Ile is indicated as the density of U’ or Γ cells which can be produced by that amount of Ura or Ile, respectively. When the nutrient concentration was not detected (under the detection limit 10^5/ml), we plotted it at 10^5/ml. (B and C) The time courses of the concentration of Ura (blue square) and Γ cells (red circle) in monoculture. (B) 10^5/ml of Γ cells were
inoculated into minimal media along with 10 μM of Ile. (C) 10²/ml of I’ cells were inoculated into minimal media along with Ile. (D) The time courses of the concentration of Ile (orange square) and U’ cells (green circle). (E) 10²/ml of U’ cells were inoculated into minimal media without the addition of Ura. These results were similar to the results of the mutualism with I’ and U’ cells shown in Fig. 1. The final concentrations of nutrients were always less than the maximum concentrations of the donor cell in monoculture (B-E), which meant that the nutrient supplies from these strains in monoculture were insufficient for the continuous growth of both strains in coculture. Despite the insufficient level of nutrient supply in monoculture, both strains grew to saturation in coculture with all of the initial cell concentrations used (F). Strain U’ (DH1ΔaenB::gfpus5ΔKmR) was constructed from DH1 cells, as was strain L’, by replacing the chromosomal pyrE gene with a foreign DNA fragment comprising a reporter gene (gfpus5) and the kanamycin resistance gene (KmR).

**Figure S5** Measurement of the cell concentrations by flow cytometry (FCM). (A) The dot plot of the data collected from the coculture by FCM. I’ cells (red dots), L’ cells (green dots), and the calibration beads (yellow dots) were clearly segregated. The definitions of these three particles were as follows: x > 1.6 (red dashed line), y > 15 (red dotted line), and Log₁₀y > Log₁₀0.43 (red solid line) for I’ cells, where x and y represent the green and red fluorescent intensities (x, y), respectively; x > 15 (green dashed line) and y < 0.036x + 10 (green solid line) for L’ cells; x > 540 (yellow dashed line), y > 0.036x + 10 (green solid line), and Log₁₀y < 0.256 (Log₁₀y²) (yellow solid line) for the beads. (B and C) The difference between the concentration determined by FCM (closed symbols) and colony forming units (cfu) (open symbols) for I’ (B) and L’ cells (C), 10²/ml (black circles) or 10⁵/ml (blue squares) of the cells were inoculated into minimal media without the addition of any amino acid, or 10²/ml of the cells were inoculated into minimal media with 10 μM of the required amino acid (Ile for I’ cells and Leu for L’ cells) (red triangles). Although there was little difference between the concentration determined by FCM and the concentration determined by cfu at time 0, the concentration determined by cfu decreased more quickly than the concentration determined by FCM. Therefore, although it is difficult to determine whether a cell is alive or dead, we defined an active cell as a cell that was able to form a single colony in the mathematical model in Fig. S1A.

(EP3)

**Table S1** Explanations and values of the parameters used in the mathematical model.

(DOC)

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**Author Contributions**

Conceived and designed the experiments: HK AK TY. Performed the experiments: HK SS YY AK KM. Analyzed the data: HK NO. Contributed reagents/materials/analysis tools: YY YS AK. Wrote the paper: HK SS TY.

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