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# Interspecific bacterial sensing through airborne signals modulates locomotion and drug resistance

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Bacteria use chemical signals to sense each other and to regulate various physiological functions. Although it is known that some airborne volatile organic compounds function as bacterial signalling molecules, their identities and effects on global gene expression and bacterial physiological processes remain largely unknown. Here we perform microarray analyses of *Escherichia coli* exposed to volatile organic compounds emitted from *Bacillus subtilis*. We find that 2,3-butanedione and glyoxylic acid mediate global changes in gene expression related to motility and antibiotic resistance. Volatile organic compound-dependent phenotypes are conserved among bacteria and are regulated by the previously uncharacterized *ypdB* gene product through the downstream transcription factors *soxS*, *rpoS* or *yjhU*. These results strongly suggest that bacteria use airborne volatile organic compounds to sense other bacteria and to change master regulatory gene activity to adapt.

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Many bacterial species co-exist in dynamic communities where they compete or cooperate with one another<sup>1,2</sup>, in the regulation of their functions<sup>3–5</sup>. Bacteria have developed intercellular signalling systems to adapt to natural variations caused by stress, competition and countering host defenses<sup>6</sup>. Several bacterial communication pathways have been discovered<sup>7</sup>. The most widespread phenomenon is quorum sensing (QS), which is an interconnected, multi-layered regulatory network that includes the secretion of QS molecules<sup>8,9</sup> and their concentration-dependent signalling cascades<sup>10</sup>. Bacteria perform QS with chemicals, such as *N*-acyl derivatives of homoserine lactone, cyclic dipeptides and quinolones<sup>10–14</sup>, to sense and respond to external and internal signals, and environmental cues. More than 300 bacteria release a wide variety of biologically active airborne volatile organic compounds (VOCs) that can diffuse in heterogeneous mixes of solids, liquids and gasses<sup>15,16</sup>, suggesting that VOC production is a general phenomenon of bacteria. However, the mechanisms underlying VOC-mediated bacterial communication, the identities of active VOCs and the cellular behaviours that are affected remain largely unknown, except in some cases. For example, in the opportunistic human pathogen *Pseudomonas aeruginosa*, the 2-aminoacetophenone VOC modulates virulence regulator activity and promotes a bistable phenotype by stochastically silencing acute virulence functions<sup>12</sup>. More directly, VOCs modify the antibiotic resistance phenotype of *Escherichia coli*<sup>13,14</sup>. It is becoming clear that VOCs have a role in interkingdom interactions between bacteria and eukaryotic cells. For example, *Vibrio cholerae* O1 produces the gaseous compound 2,3-butanediol, which interacts with the human immune system by suppressing the induction of immune modulators<sup>17</sup>. Bacterial VOCs also have been shown to promote plant growth<sup>18</sup> and induce systemic resistance in *Arabidopsis*<sup>19</sup>. Therefore, the roles played by bacterial VOCs and their functional mechanisms need to be elucidated fully to develop a wider understanding of bacterial survival in dynamic multitrophic interactions.

The structures of Gram-negative and Gram-positive bacteria appear significantly different, but recent metagenomics studies showed that both bacterial species co-exist in any natural habitats, such as human, soil or water, with different populations<sup>20–23</sup>. The abilities of these bacteria to communicate and respond to different environmental conditions could be major factors in making a large and dynamic bacterial community. There are many potential trade-offs for communications between two bacterial groups that allow bacteria to co-exist and maintain their own survival in variable environments. However, it is technically challenging to experimentally study bacterial communications in variable environments. The two species *E. coli* and *Bacillus subtilis* are known to share similar ecological niches<sup>20–22</sup>. In this study, *E. coli* and *B. subtilis* were used as representative models for studying the roles and effects of VOCs in Gram-negative and Gram-positive species, respectively. Microarray analysis was used to examine gene expression in *E. coli* in response to exposure to VOCs from *B. subtilis*, to understand how these VOCs affect the intercellular regulation of gene expression. VOCs dramatically affected the phenotypes of receiver cells and altered the expression of more than 100 genes. Treatment with 2,3-butanediol (2,3-BD) and glyoxylic acid (GA) in nanomolar to micromolar concentrations mediated changes in global gene expression that correlated with motility and antibiotic resistance phenotypes. VOC-mediated interactions were modulated by the previously uncharacterized *ypdB* gene product and its downstream transcription factors, *soxS*, *rpoS* or *yjhU*. The results of this study strongly suggest that VOCs represent a cross-specific signal for rapid sensing between bacteria that triggers variations in gene expression.

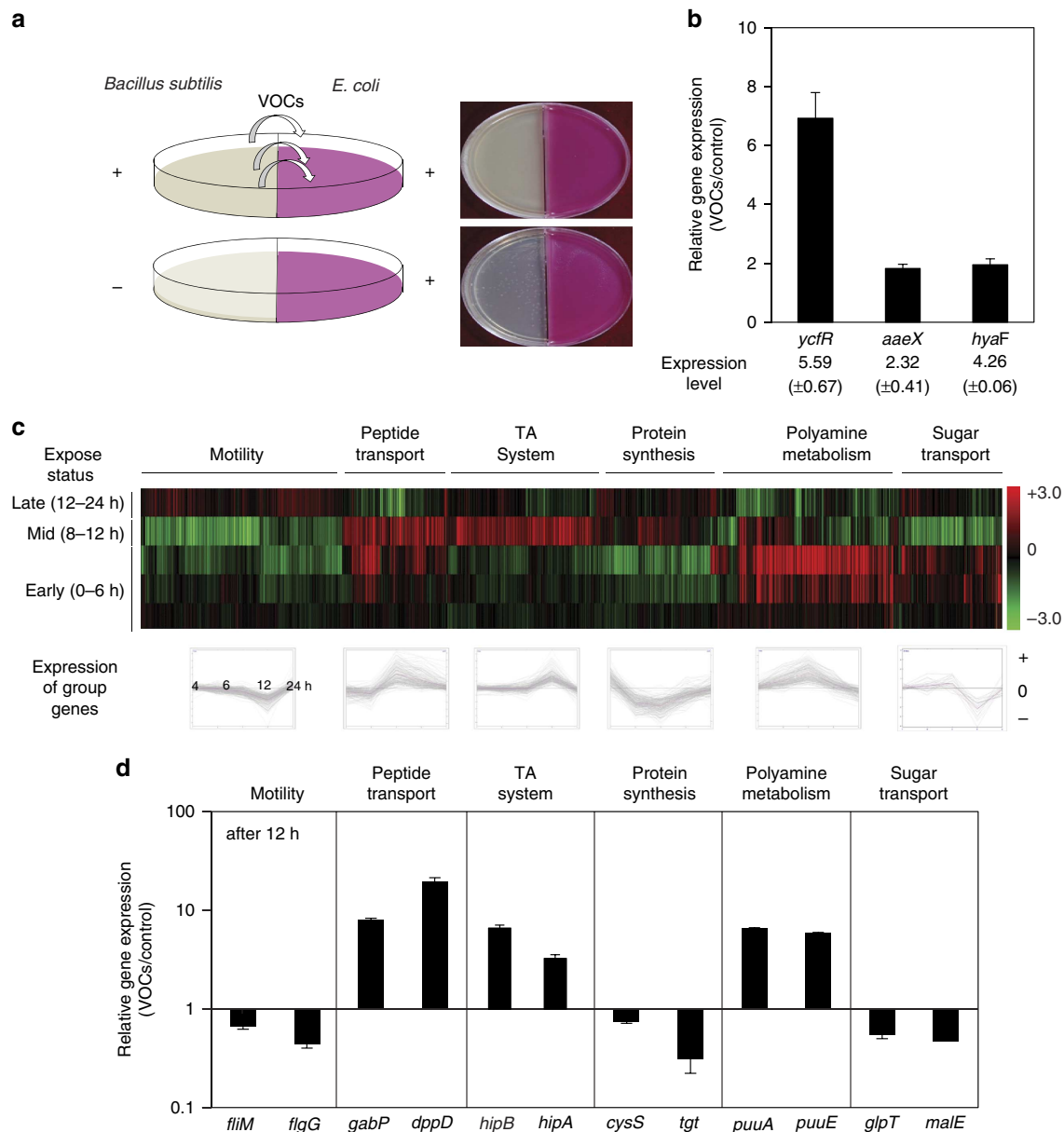
VOC-mediated bacterial communication may induce modifications that affect the self-defence and motility of physically separated bacteria.

## Results

***B. subtilis* VOCs alter the expression of *E. coli* genes.** A model system was designed to examine the mechanisms underlying phenotypic modifications induced by bacterial metabolites in interspecies interactions. *B. subtilis* 168 (ref. 24), and *E. coli* K12 strains were used as emitter and receiver bacteria, respectively. *B. subtilis* produces diffusible secondary metabolites, such as antibiotics and VOCs<sup>25</sup>. These products can affect plant defence systems by eliciting innate immunity against a broad spectrum of pathogens<sup>18,19</sup>.

It was hypothesized that the VOCs produced by *B. subtilis* modulate the expression profiles and functions of certain *E. coli* genes. A two-compartment Petri plate (I-plate) assay was used for concurrent growth of bacteria (Fig. 1a). In this assay, *E. coli* grown on Luria-Bertani (LB) agar was exposed to VOCs from *B. subtilis* 168 cell suspensions grown on tryptic soy broth (TSB) agar. There was no physical contact between the cultures or the bacteria. *E. coli* cells were collected after different periods of exposure to VOCs (6, 12 and 24 h), and transcription profiling of *E. coli* gene expression was performed using microarray analyses of 4,440 open-reading frames. The 6-, 12- and 24-h treatments showed differential expression of 137, 569 and 93 genes (> 3-fold change), respectively (Supplementary Table S1). The largest change in gene expression occurred after 12 h of exposure to *B. subtilis* VOCs. To confirm the responses of these genes to *B. subtilis* VOCs, three of the top 50 genes identified by microarray analysis (*ycfR*, *aaeX* and *hyaF*) were selected for further analysis. Gene expression was examined by reverse transcription quantitative PCR (RT-qPCR). As expected, all three genes showed increased expression in response to *B. subtilis* VOCs (Fig. 1b), with *ycfR* exhibiting the greatest change. Further analyses were performed using the Gene Ontology website tools<sup>26</sup> (Supplementary Table S1) or Kyoto Encyclopedia of Genes and Genomes<sup>27</sup> information. Strongly affected genes were allocated into functional groups, such as motility, peptide transport, toxin-antitoxin (TA) system, protein synthesis, polyamine metabolism and sugar transport (Fig. 1c, Supplementary Table S2). To validate the microarray data, two different transcripts were selected from each group, and expression profiles were determined by RT-qPCR. These data closely matched the Gene Ontology patterns of the microarrays (Fig. 1d). Thus, the microarray results identified *E. coli* genes modulated by *B. subtilis* VOCs. Their functional clusters may shed light on the mechanisms underlying the physiological responses of this receiver.

**VOCs change the expression of motility-related genes.** From microarray data, VOC treatments decreased the expression of all 30 genes related to chemotaxis or the motility phenotype of *E. coli* (Fig. 1d, Supplementary Tables S1 and S2). Motility-related genes are one of the key clusters associated with VOC-mediated physiological changes. Examination of the expression of some motility genes (*motA*, *cheA*, *cheW* and *fliA*) using RT-qPCR confirmed the microarray data (Fig. 2a). To verify the gene-expression changes correlated with the motility phenotype, the swimming and swarming abilities of VOC-treated *E. coli* cells were measured. Both swimming (Fig. 2b) and swarming (Fig. 2c) decreased by ~32% and ~60%, respectively. To test whether this phenotype is limited to *E. coli* or if it is a general trait of bacteria, *B. subtilis* VOCs were used to treat other Gram-negative bacteria, such as *Burkholderia glumae* BGR1 and *P. aeruginosa* PA14, and

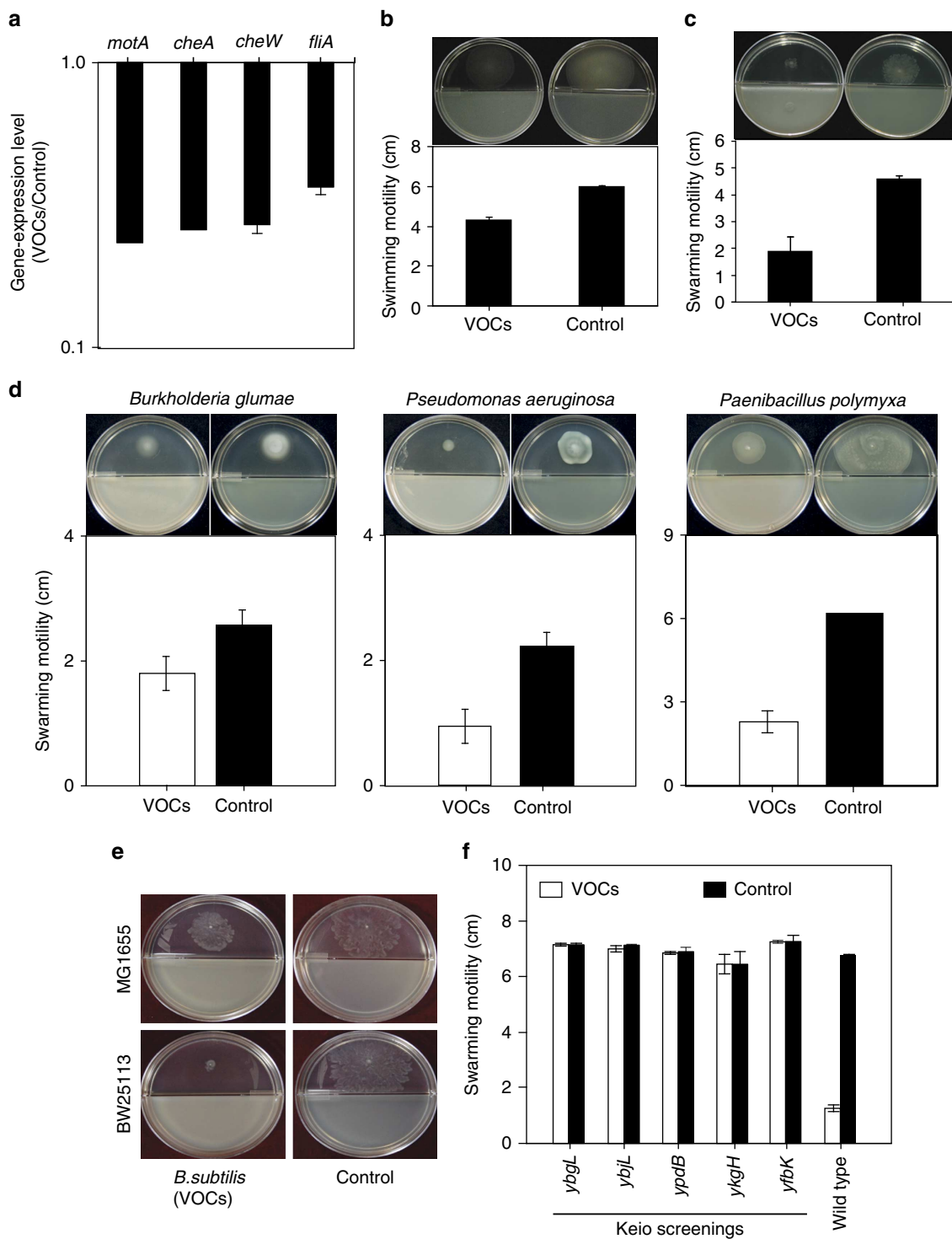


**Figure 1 | Effect of *B. subtilis* VOCs on *E. coli* gene expression.** (a) Schematic showing the two-compartment plate assay (1-plate system). For growth of *E. coli*, LB agar was used; for visualization and photography, MacConkey-agar plate was used. (b) Relative gene-expression analysis of *E. coli*. Three genes showing >3-fold changes in expression in response to *B. subtilis* VOCs in microarray data are represented as ‘Expression level’; these were randomly selected and analysed by RT-qPCR after *E. coli* cells were treated with VOCs for 12 h. Error bars are defined as s.e.m. ( $n=3$ ,  $P=0.05$ ). (c) Clustering of microarray data. The microarray data show >3-fold up- or downregulation induced by treatment with VOCs. Clustering analysis was performed using MultiExperiment Viewer<sup>45</sup>. Three treatment periods are shown (6, 12 and 24 h) (d) Gene-expression pattern of clustered groups. Two genes were randomly selected from each of the groups in c and examined by RT-qPCR using gene-specific primers listed in Supplementary Table S6. Error bars are defined as s.e.m. ( $n=3$ ,  $P=0.05$ ).

the Gram-positive bacteria *Paenibacillus polymyxa* E681. The swarming motility of each receiver bacteria was measured as a representative of motility phenotype. Figure 2d shows that swarming motility for all the species tested was decreased by VOCs, suggesting that VOCs are the general signalling molecules for altering bacterial motility. Because swarming ability was more strongly affected by VOCs than swimming ability, the changes to swarming activity were examined in greater detail.

To understand how the phenotype is modulated, the genes affecting the phenotype need to be identified. Therefore, a mutant library of *E. coli* genes<sup>28</sup> was screened to evaluate the effect of VOCs on the swarming motility. Because the Keio collection

was developed from the strain *E. coli* BW25113<sup>29</sup>, and different *E. coli* strains showed different motilities<sup>30</sup>, the effect of VOCs on the swarming motility of phenotypes using MG1655<sup>31</sup> needed to be compared with that of BW25113 to show a general phenotype of *E. coli*. Exposure to VOCs caused a significant reduction in the swarming motility of *E. coli* BW25113 and MG1655 (Fig. 2e), suggesting that the effect is a general phenotype of *E. coli*. Therefore, BW25113 cells were used to validate the motility-related genes examined in Fig. 2e. Mutants of *motA*, *cheA*, *cheW* and *fliA* showed no motility variation when the cells were exposed to VOCs, which suggests that they are essential for swarming ability (Supplementary Fig. S1).



**Figure 2 | Effect of VOCs on bacterial motility.** (a) Change of motility-related gene expression. The expression of genes *motA*, *cheA*, *cheW* and *fliA* in *E. coli* MG1655 was measured by RT-qPCR after treatment with VOCs. The gene-expression level indicates the normalized gene level of VOC-treated sample to non-treated sample (control). The data shown are the averaged values ( $n=5$ ,  $P=0.05$ ). Error bars are defined as s.e.m. VOC-induced changes in (b) swimming and (c) swarming motility of *E. coli*. (d) Conservation of swarming motility in Gram-negative and Gram-positive bacteria. (e) Comparison of swarming motility in two different *E. coli* strains. The plates are representative of data obtained in triplicate. (f) *E. coli* mutant strains in which swarming motility was unchanged by VOCs. Supplementary Table S1 contains further information about these genes.

However, the lack of motility in these mutants was similar to that of VOC-treated wild type. To identify a possible regulatory gene responsible for the VOC-mediated changes in swarming ability, mutants from the Keio collection were screened to identify knockouts of VOC-mediated genes that showed no

significant variation in swarming motility with or without VOC treatment. Five mutants (*ybgL*, *ybjL*, *ypdB*, *ykgH* and *yfbK*) were identified (Fig. 2f, Table 1). These data suggest that VOCs affect *E. coli* motility via previously uncharacterized motility-related genes.

**Table 1 | Genes affecting VOC-dependent swarming motility.**

Gene	Location in the <i>E. coli</i> genome (direction)*	Function and description of gene product
<i>ybgL</i>	744,388–745,122 (→)	Unknown, low-abundant protein, UPF0271 family
<i>ybjL</i>	887,357–889,042 (←)	Unknown, putative AAE family transporter
<i>ypdB</i>	2,498,405–2,499,139 (→)	Unknown, predicted response regulator cognate to YpdA sensor protein
<i>ykgH</i>	323,920–324,588 (←)	Unknown
<i>ompF</i>	985,117–986,205 (←)	Outer membrane porin F
<i>yfbK</i>	2,382,017–2,383,744 (←)	Unknown, predicted VMA domain lipoprotein

Abbreviations: AAE, Aspartate:alanine exchanger; VMA, von Willebrand factor type A.  
\*The coding region for each protein on the *E. coli* K12 genome. The direction of the ORF is indicated by arrows (→:clockwise; ←:counter-clockwise). This information was adapted from www.ecogene.org.

**Antibiotic-resistance phenotype is mediated by TA genes.** From the microarray data, 27 different TA-related modules (Supplementary Table S2) were upregulated by VOCs. The toxins *relE* (5.9-fold), *yafQ* (7.8-fold) and *yoeB* (6.8-fold) were strongly induced in persister cells<sup>32–34</sup>. Stationary-phase cultures of *E. coli* with *hipBA*, a representative TA system, represented a subpopulation of bacteria (~1% in the stationary state) that were viable after treatment with lethal concentrations of antibiotics<sup>35</sup>. Many bacterial VOCs modulate antimicrobial activity<sup>36–38</sup>. Therefore, we hypothesized that *Bacillus* VOCs could modulate multidrug resistance. First, the effect of VOCs on persister cells was analysed with respect to susceptibility to 240 commercial antibiotics. For this purpose, we used the Phenotype Microarray (PM) technique, which is a high-throughput technology for simultaneous testing of a large number of cellular phenotypes. It uses preconfigured well arrays of different cellular phenotypes and an automated instrument that monitors and records the response of cells in all wells of the arrays<sup>39</sup>. The effect of *B. subtilis* VOCs on the susceptibility of *E. coli* to different antibiotics varied (Supplementary Fig. S2). Although VOCs exerted a marked effect on bacterial cells exposed to the PM12 plate, they did not improve resistance to all the antibiotics tested (Fig. 3a).

Thirteen antibiotics with Respiration Area >100 or <−100 were identified (Supplementary Table S3) from PM11 to PM20 plates. *E. coli* cells showed resistance to ten antibiotics and susceptibility to three antibiotics. Only the resistant phenotype was detected in the penicillin/ampicillin and tetracycline groups (Fig. 3b). These data suggest that resistance to antibiotics in these groups may be dependent on VOC treatment. The PM data were validated by growing VOC-treated and VOC-untreated *E. coli* on ampicillin and tetracycline. VOCs increased the number of ampicillin- and tetracycline-resistant cells by 8.7- and 2.3-fold, respectively (Fig. 3c).

Microarray data (Fig. 1, Supplementary Table S1) showed that VOCs induced expression of *hipA* and *hipB* genes by 14.5- and 21.2-fold, respectively. VOC-mediated induction of these genes was confirmed by RT-qPCR (Fig. 3d). The expression of *hipA* is known to mediate persistence<sup>40</sup>. Mutant *hipA7* cells show a very high frequency of persister formation, and overexpression of wild-type *hipA* confers persistence<sup>41–43</sup>. The effect of VOCs on ampicillin resistance in *E. coli* was examined with respect to expression of *hipA* to determine if *hipA* has a role in the VOC-mediated resistance phenotype. VOCs increased the number of wild-type cell colonies growing on ampicillin by 2.8-fold, but this effect was not observed in the *hipA*-deletion strain with control plasmid, in which

the number of ampicillin-resistant colonies was reduced by ~30% (Fig. 3e). This effect was fully complemented in the knockout strain by an overproducer of *hipA*, which suggests that *hipA* is the necessary factor for regulating VOC-induced antibiotic resistance. These results indicate that, although TA-related genes are involved in VOC-dependent antibiotic resistance, additional cellular factors must be identified to develop a full understanding of the VOC-induced TA system.

#### Identification of VOCs that affect bacterial phenotypes.

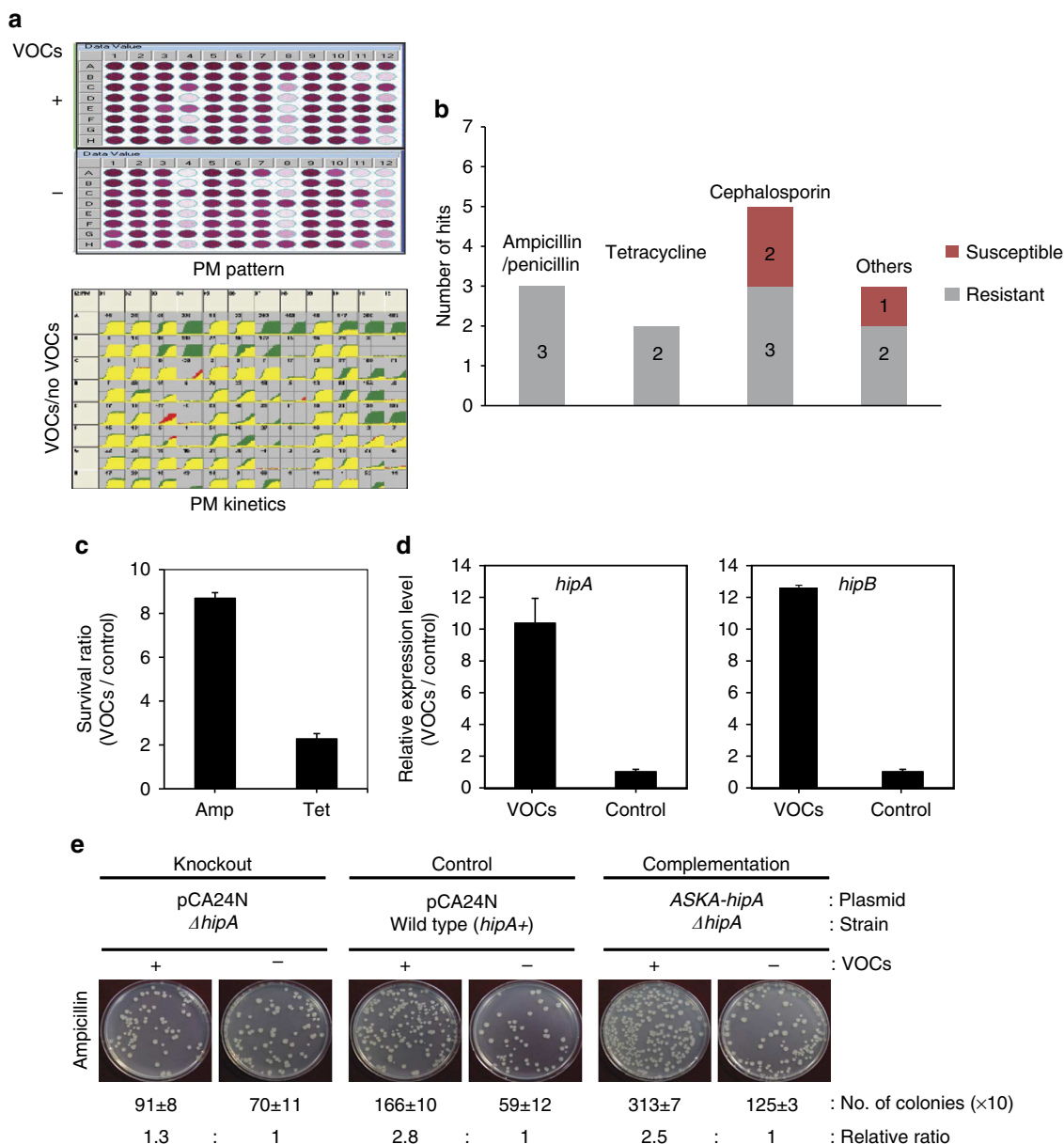
*B. subtilis* emits ~30 different VOCs<sup>44</sup>, but how many or which specific VOCs modulate newly identified phenotypes remains to be validated. Therefore, the effects of individual *B. subtilis* volatiles on the changes in motility and antibiotic resistance were tested. We selected seven commercially available VOCs of the ~30 VOCs<sup>44</sup>, and examined VOC-mediated induction of swarming motility and the antibiotic resistance phenotype. The screenings showed that exposure to 2,3-BD, one of the primary ketones synthesized in *B. subtilis* GB03, and GA, the second-most synthesized acid from *B. subtilis* GB03<sup>44</sup>, reproducibly reduced swarming motility (Fig. 4a). These VOCs elicited a resistance response to ampicillin that was similar to the total VOC-induced phenotype in both strains (Fig. 4b). The level of antibiotic resistance induced by 3M1BOH and 2M1PAOL was dramatically different in the two strains (Fig. 4b). Variations in the architecture of strains can influence motility and biofilms<sup>30</sup>, and it is possible that the architecture of these *E. coli* strains contributes to their different responses to VOCs. These data strongly suggest that VOCs, in particular 2,3-BD and GA, modulate swarming motility and antibiotic resistance in *E. coli* cells.

#### Determination of effective VOC concentrations for phenotypes.

Our data clearly showed an effect of total and individual airborne VOCs from physically separated bacteria on the phenotypes of receiver bacteria. However, how VOCs, which are gaseous molecules, affect the receiver bacteria is not clear. It is possible that gaseous VOCs are solubilized in the culture media of *E. coli*, and the amount affects their function. Therefore, a determination of the working concentrations of 2,3-BD and GA that are effective for the bacterial phenotypes will indicate the relevance of two airborne VOCs on the physiology of physically separated bacteria.

To determine the index of the VOC effect, we measured the effective concentration of 2,3-BD and GA by natural vaporization of *B. subtilis* VOCs to *E. coli* cells. For this purpose, swarming motility was measured after treatment with different concentrations (10 nM to 100 mM) of 2,3-BD or GA for 16 h. As seen in Fig. 4c, application of 10 nM of 2,3-BD or GA by the method described in Fig. 1 modified the swarming motility (~22–25% decrease compared with non-treated controls), whereas total VOCs decreased motility by ~50%. This information defines tentative volatility as the percentage change of motility by time of VOC treatment (% h<sup>−1</sup>). Therefore, the volatility of 2,3-BD and GA is ~22–25% h<sup>−1</sup>, and ~50% h<sup>−1</sup> for total VOCs. To determine the amount of 2,3-BD or GA in the media that directly affects the phenotype, either 2,3-BD or GA from 1 nM to 1 μM was directly added to the *E. coli* cell cultures. With 1 nM of 2,3-BD or GA, there was a linear decrease in swarming motility. At 1 μM of 2,3-BD or GA, there was a volatility of ~62% h<sup>−1</sup> or ~45% h<sup>−1</sup>, respectively (Fig. 4d). In those conditions, the growth of *E. coli* was not affected by the increased concentration of individual VOCs (Supplementary Fig. S3). *E. coli* were resistant to ampicillin, even when picomolar final concentrations of 2,3-BD and GA were added (Supplementary Fig. S4).

These results indicate that both gas and liquid phases of VOCs affect motility and susceptibility to antibiotics when receiver



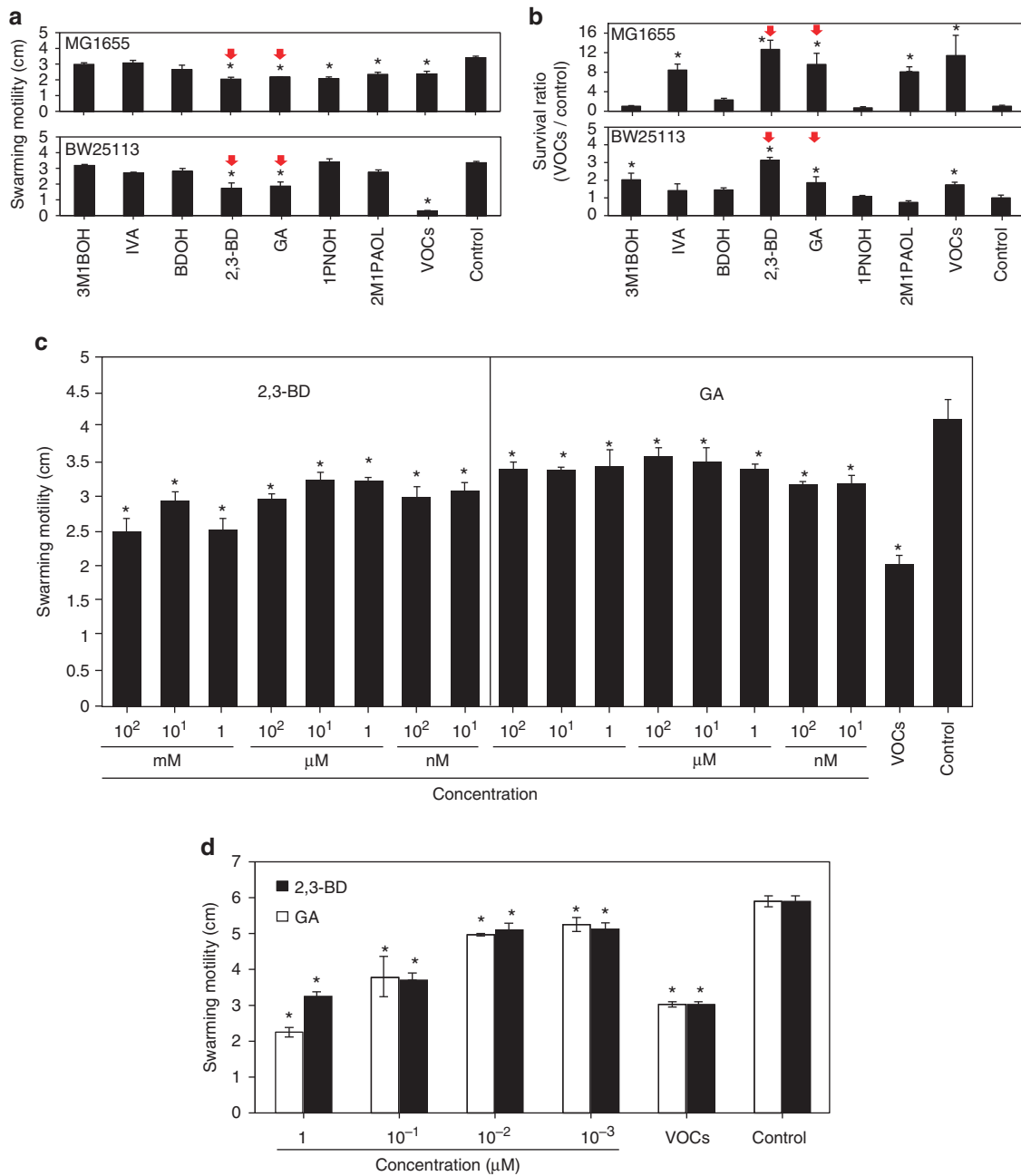
**Figure 3 | Modulation of susceptibility to antibiotics by VOCs.** (a) Representative kinetic analysis of antibiotic susceptibility of *E. coli* cells from  $n = 3$  with  $P = 0.05$ . PM pattern (left) and PM kinetics (right) of a PM12 plate. (b) Quantitative analysis of PM kinetics from PM11 to PM20. (c) Effect of VOCs on antibiotic resistance in *E. coli*. The relative survival ratio of VOC-treated to non-treated *E. coli* in ampicillin or tetracycline is shown. (d) Expression of *hipA* and *hipB* in the presence and absence of VOCs. Gene expression was monitored by RT-qPCR. Error bars are defined as s.e.m. from triplicate experiments. (e) Growth on ampicillin is dependent on *hipA* expression. VOCs did not improve survival in the *hipA*-knockout strain, whereas VOCs improved survival in the complemented strain. The plasmids pCA24N and ASKA-*hipA* were used<sup>50</sup>.

bacteria are treated with VOCs in concentration ranges of pM to  $\mu$ M, which are physiologically relevant concentrations.

**Mechanistic studies on the VOC-induced pathways.** The data indicated that VOCs affect the expression of groups of genes related to cellular phenotypes of bacteria. Another question is how VOCs affect the phenotypes. Because the cellular pathways are generally mediated by complex systems, it might be postulated that a master regulatory gene controls the VOC-mediated phenotypes by modulating transcription factors. To identify the master regulators, knockout mutants of VOC-modulated genes that no longer exhibited VOC-related phenotypes, such as swarming motility or antibiotic resistance, were selected for

further investigation. VOCs exerted no change in the swarming motility of five candidate knockout mutants (*ybgL*, *ybjL*, *ypdB*, *ykg* and *yfbK*; Fig. 2e, Table 1). When the effect of VOCs on the antibiotic resistance of these mutants was studied, only the absence of *ypdB* resulted in reduced antibiotic resistance (Fig. 5a). The *ypdB* mutation showed the same effect on bacterial growth as in wild-type cells (Fig. 5b).

Swarming motility in the *ypdB* knockout was similar to the untreated control, and did not show any significant change following exposure to 2,3-BD, GA or total VOCs (Fig. 5c). The survival ratio of the *ypdB*-knockout cells was similar to that in wild-type cells in the absence of VOCs. However, the survival ratio of *ypdB*-knockout cells was lower than that in wild-type cells exposed to 2,3-BD and total VOCs (Fig. 5d). VOC-induced

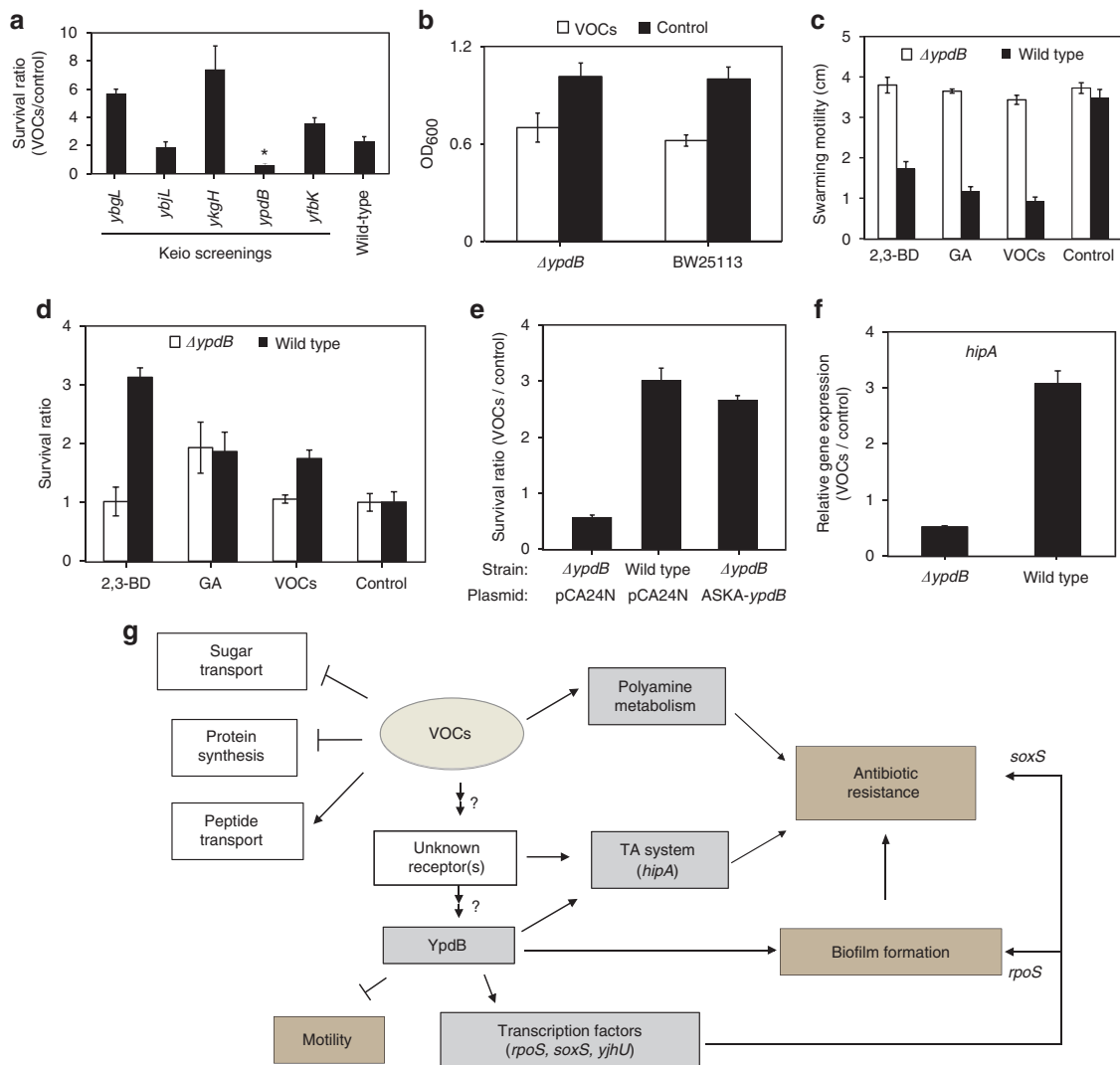


**Figure 4 | Identification and characterization of individual VOCs affecting *E. coli* phenotypes.** Effect of individual VOCs on (a) swarming motility and (b) ampicillin resistance in *E. coli* MG1655 and BW25113 ( $n = 3$ ,  $P = 0.05$ ). The antibiotic resistance phenotype was expressed as the survival ratio of VOC-treated and VOC-untreated *E. coli* grown on ampicillin. Control represents no VOC treatment. Determination of effective concentration of individual VOCs by (c) natural vaporization (gaseous phase) or by (d) direct addition (aqueous phase). Asterisks (\*) indicate statistically significant data ( $P = 0.05$ ) and arrowheads indicate individual VOCs that show the same phenotype as total VOCs in both BW25113 and MG1655 cells. Error bars are defined as s.e.m.

antibiotic resistance was eliminated by the *ypdB* deletion and complemented by ectopic induction of *ypdB* in the knockout strain (Fig. 5e). The expression of *hipA* was severely reduced in the *ypdB* knockout (Fig. 5f), which suggests that the *ypdB* gene product might function as an upstream activator or transcription factor for *hipA* expression. Therefore, *ypdB* may be the master regulator of VOC-mediated effects in *E. coli*.

There have been many reports that chemicals work as signals for the transcription of specific genes<sup>45</sup>. We hypothesized that gene-expression changes induced by VOCs reflect the modified activity of transcription. To test our hypothesis, we examined the

transcription activity (that is, promoter activity) of some genes that are upregulated by VOCs. For this purpose, chromosomal promoter-green fluorescent protein (GFP) fusions of three genes (*ycfR*, *aeX* and *hyaF*) that show >3-fold changes in transcript levels induced by VOCs (Fig. 1) were generated. Then, the expression of hybrid-GFP proteins in *E. coli* cells exposed to *B. subtilis* VOCs and untreated controls (no *B. subtilis*) were measured. All selected genes showed increased expression of GFP in response to *B. subtilis* VOCs (Supplementary Fig. S5), suggesting that VOCs could affect the transcription of target genes by modulating promoter activity.



**Figure 5 | Master regulation of VOC-mediated phenotypes by the *ypdB* gene product.** (a) Identification of VOC-regulated genes involved in antibiotic resistance. Survival ratio of VOC-treated to VOC-untreated Keio collection mutants grown in the presence of ampicillin ( $n=3$ ). Asterisk (\*) indicates the statistical significance ( $P=0.05$ ). (b) Growth phenotypes of wild-type and *ypdB* knockout by VOCs. The growth (OD<sub>600</sub>) was monitored by the absence or presence of VOCs by natural vaporization. Comparison of the effect of 2,3-BD and GA on swarming motility (c) and antibiotic resistance (d) in a *ypdB*-knockout and wild-type *E. coli*. (e) Survival ratio of VOC-treated to VOC-untreated cells in a *ypdB* knockout, the complemented knockout and wild-type cells grown in the presence of ampicillin. (f) Relative change in *hipA* expression with and without VOC treatment in wild-type *E. coli* and *ypdB*-knockout cells ( $n=3$ ,  $P=0.05$ ). (g) Model for VOC-mediated changes to *E. coli* physiology. The model is based on data from this study and previous reports. Error bars are defined as s.e.m.

We searched for transcription-related factors that were related to the phenotypes. For this purpose, we introduced a fluorescent promoter library<sup>46</sup> for the screening of factors on the phenotypes. From the library, 60 available promoter reporters were selected as transcription-related libraries after filtering by keywords of transcription or DNA-binding transcription regulation using both EcoCyc<sup>47</sup> and gene descriptions of microarray data (Supplementary Table S4). A 96-well plate was used to screen reporters with natural vaporization from 1  $\mu$ M of 2,3-BD or GA or *B. subtilis* VOCs (from 10<sup>5</sup> cells) for 14 h at 37 °C. The promoter activity as fluorescence from *E. coli* promoter reporters was monitored as fluorescence/OD<sub>600</sub>. From the results of 60 different promoters (Supplementary Table S4), the activity of three different genes (*yjhU*, *soxS* or *rpoS*) was reproducibly induced by all tested VOCs (2,3-BD, GA and *B. subtilis* VOCs) (Table 2). Therefore, these genes could be either upstream or downstream genes of VOC-mediated responses by the *ypdB*

product. This hypothesis was further validated by analysis of the steady-state level of those transcription factors using RT-qPCR in the presence or absence of *ypdB* or individual VOCs (2,3-BD). We found that the levels of three genes (*soxS*, *yjhU* and *rpoS*) were increased by treatment with 2,3-BD (Table 2). The gene expression of all targets decreased in the absence of *ypdB*, suggesting that those genes are activated by the *ypdB* gene product and are located downstream of *ypdB*. These data indicate that the *ypdB* product and its downstream transcription factors mediate the regulatory cascades of VOC-induced phenotypes.

## Discussion

Our study showed that airborne VOCs are important molecules in sensing distant bacteria even without physical contact. We found that gaseous VOCs released from *B. subtilis* severely affect neighbouring Gram-negative bacteria either as gas or liquid molecules, resulting in



**Table 2 | Effect of VOCs on promoter activity\* or transcript level†.**

Genes	Fluorescence reporter screenings (fluorescence/OD <sub>600</sub> )			RT-qPCR (relative amount)	
	<i>B. subtilis</i> VOCs	2,3-BD	GA	WT + 2,3-BD	$\Delta ypdB$ + 2,3-BD
<i>yjhU</i>	1.28 ± 0.13	1.22 ± 0.08	1.48 ± 0.09	1.30 ± 0.06	0.88 ± 0.03
<i>soxS</i>	1.42 ± 0.08	1.33 ± 0.02	1.53 ± 0.04	1.61 ± 0.10	1.03 ± 0.12
<i>rpoS</i>	1.44 ± 0.03	1.24 ± 0.08	1.58 ± 0.09	1.13 ± 0.04	0.63 ± 0.02
<i>ypdB</i>	N.T.	N.T.	N.T.	1.21 ± 0.01	N.D.

Abbreviations: N.D., not detected by the experiments; N.T., not tested due to the absence of the gene-fusion in the library.

Fold change to control cells (without VOC treatment) were shown from three independent experiments.

\*For fluorescence measurements, *B. subtilis* VOCs (from 10<sup>5</sup> cells) or individual VOCs (1 μM) each were subjected to natural vaporization, exposed to *E. coli* cells containing individual gene promoter reporters for 14 h at 37 °C, and the fluorescence/OD<sub>600</sub> values were measured using a Multiplate reader. All data of reporter screenings are shown in Supplementary Table S2.

†Transcript level of each gene was measured by RT-qPCR from *E. coli* cells treated with 2,3-BD (1 μM) at OD<sub>600</sub> = 0.2 until OD<sub>600</sub> reached 1.0 for 1 h at 37 °C. Data are the averaged values from *n* = 3, *P* = 0.05.

a change of gene-expression profiles that alter both motility and antibiotic-resistance phenotypes. The VOC-mediated motility phenotype is conserved in both Gram-negative (*B. glumae* and *P. aeruginosa*) and Gram-positive (*P. polymyxa*) bacteria. The 2,3-BD and GA from total VOCs of emitter bacteria are major signalling molecules that affect motility, antibiotic resistance<sup>48</sup> and increased biofilm formation (a general trait relating to antibiotic resistance<sup>49</sup> and *rpoS* mediation<sup>50</sup>) (Supplementary Fig. S6). The VOC-mediated phenotypes are linked to a previously uncharacterized *ypdB* gene product and downstream transcription factors. Therefore, these VOC signalling cascades of *ypdB*/transcription factors could rapidly alter gene-expression profiles and result in multiple phenotypic changes involved in bacterial growth and adaptation to environmental stresses.

VOCs from emitting bacteria might function as effector molecules that target transcription-related factors of interacting bacteria, resulting in the rapid alteration of gene-expression profiles. This seemed to be an energetically efficient way to simultaneously control many genes. This hypothesis was supported by the results that VOCs increased the promoter activity of genes that were upregulated in the microarray analysis (Fig. 5a), including several known stress-responsive transcription factors (*soxS*, *rpoS* and *yjhU*). The gene expression of such transcription factors could function as the downstream genes to *ypdB*, considering the decreased gene expression in the absence of *ypdB* when 2,3-BD is present (Fig. 5). Our results indicate that VOC-mediated phenotypes are mediated by the master regulation of the *ypdB* gene product and its downstream transcription factors. Therefore, the genes identified in this study could be useful as targets for the development of antimicrobials.

Although many bacteria produce VOCs<sup>25</sup>, little is known about their physiological functions. Several recent reports indicate that bacteria are responsive to VOCs, including ammonia. Ammonia is a valuable nutrient for the production of proteins and nucleic acids for bacteria. The Burgess group recently showed that the soil bacterium *Bacillus licheniformis* was sensitive to the presence of ammonium sulphate<sup>51</sup>, which is metabolized and converted to ammonia, resulting in the induction of biofilm formation. The mechanisms underlying the roles played by VOCs in bacterial communication were unknown. Ghigo and colleagues<sup>14</sup> also showed that ammonia from *E. coli* influenced antibiotic resistance in aeri ally exposed Gram-positive and Gram-negative bacteria. The authors discovered a previously uncharacterized antibiotic-resistance mechanism and showed that this effect contributed to

the induction of ammonia- and polyamine-related genes during the stationary phase. In our unpublished preliminary results, we did not identify ammonia as a major signal from *B. subtilis* as a source of VOCs. Therefore, we did not perform experiments with polyamine as a major VOC. This suggests that the source of the two VOCs was completely independent from volatile amines. Our study provides the first evidence of a mechanism underlying VOC-modified gene expression in interacting bacteria each other.

The volatility of gaseous materials is an important factor for understanding the fate of individual VOCs. We determined the tentative volatility of 2,3-BD and GA as the percentage change of motility by time of VOC treatment (% h<sup>-1</sup>) (Fig. 4), which was ~62% or 45% h<sup>-1</sup> for 2,3-BD or GA, respectively. However, this is not a general index and many people use Henry's constant (k<sup>o</sup><sub>H</sub>, mol kg<sup>-1</sup> · bar) instead, which states the amount of gas that dissolves in liquid at equilibrium and constant temperature. By comparing the k<sup>o</sup><sub>H</sub> in water at 25 °C, the volatility of 2,3-BD (k<sup>o</sup><sub>H</sub> = 74)<sup>52</sup> is much higher than that of GA (k<sup>o</sup><sub>H</sub> = 9,000)<sup>53</sup>. These data suggest that 2,3-BD will most likely evaporate, whereas GA is effectively dissolved in water. In our study, the solubilizing amount of individual VOCs was not a decisive factor for phenotypes, considering that both gas and liquid phases of VOCs at pM to μM concentrations were sufficient for the phenotypes.

Using gas chromatography-mass spectrometry solid phase microextraction (GC-MS SPME) profiling of rhizobacterial volatiles, 2,3-BD was released at 77 μg per 24 h, whereas GA was released at ~1 μg from *B. subtilis* GB03 (ref. 44). The release of 2,3-BD was less in Gram-negative bacteria, such as *E. coli* (DH5α) (14 μg per 24 h) and *Pseudomonas fluorescens* 89B61 (7 μg per 24 h), whereas *Bacillus amyloliquefaciens* IN937a released 38 μg per 24 h. The production of the large amount of 2,3-BD from *Bacillus* species was still an enigma, but we showed that 2,3-BD (even with lower amounts as either gas or liquid phase) affects specific biological processes.

The VOC 2,3-BD occurs naturally in alcoholic beverages and is added to some foods to give a buttery flavour and aroma. Food flavourings containing 2,3-BD are used in microwave popcorn and other snack foods, pet foods, candies, baked goods and other food products<sup>54</sup>. It is an antimicrobial compound produced by bacteria such as lactic acid bacteria, which produce antimicrobial compounds including hydrogen peroxide, carbon dioxide, 2,3-BD, various uncharacterized compounds and high molecular mass compounds, such as bacteriocin<sup>55</sup>. However, its effectiveness as an antimicrobial agent among bacteria seems not to be a generalized phenomenon, but is context dependent. For example, 5.8–29 mM of 2,3-BD in skimmed milk is more effective against Gram-negative bacteria than Gram-positive bacteria, with lactobacilli having the highest resistance<sup>55,56</sup>. The minimum-inhibitory concentration of 2,3-BD among bacteria differs according to pH (ref. 55). In our study, *E. coli* growth was severely decreased by the addition of 2,3-BD (>1 mM) (Supplementary Fig. S3). This appeared to be the minimum susceptible concentration of 2,3-BD as antibiotics for *E. coli*, which are matched to the Hedgecock and Jay's results<sup>55,56</sup>. However, at extremely low dosage (nM to μM concentration) of 2,3-BD, the effect is completely reversed to that of higher dosage.

The low amount of 2,3-BD that is effective to induce VOC-mediated phenotypes is comparable to the concentration of 2,3-BD in wines for the taste threshold: 0.2 mg l<sup>-1</sup> (2.4 μM) for Chardonnay wine, 0.9 mg l<sup>-1</sup> (10.8 μM) for Pinot Noir and 2.8 mg l<sup>-1</sup> (33.6 μM) for Cabernet Sauvignon<sup>57</sup>. How a low dosage of 2,3-BD affects bacterial diversity is related to human health, although a higher dose of 2,3-BD induces respiratory toxicity in mammals<sup>58–60</sup>. One of the easy samples will be the wine microbiome<sup>61</sup>. We could identify the negative or positive selection pressure of bacterial diversity using 2,3-BD.

GA is produced as an intermediate of the glyoxylate cycle, which is a variation of the tricarboxylic acid cycle that synthesizes carbohydrates from C2 compounds, such as ethanol and acetate<sup>62</sup>. This product as a VOC occurs in many bacteria<sup>44,62,63</sup>, such as *Bacillus*, *E. coli* and *Rhodobacter* species. Therefore, our results that GA functions as a signal molecule to modulate multiple phenotypes might be correlated with general bacterial acetate metabolism. Further work will identify VOC-responsive targets in the pathway, and these will be good candidates for controlling the antibiotic-resistance pathways.

Although 2,3-BD and GA were effective at modulating the expression profiles of genes related to bacterial motility and antibiotic resistance in *E. coli*, many biological questions remain unanswered. These include the identification of additional biogenic volatile molecules and the complete mechanistic behaviour of long-range interactions. Similar to the recent progress made in identifying the interactions mediated by indole<sup>6</sup>, which was a known bacterial product for over 100 years, the biological roles of VOCs are only now beginning to be elucidated. However, it is clear that the VOC-mediated pathways are an important method of communication between physically separated bacteria. They induce changes in gene expression that may facilitate survival, such as increased antibiotic resistance. Therefore, VOC-mediated bacterial communication may promote modulation of self-defence phenotypes and motility changes in physically separated bacteria.

## Methods

**Strains.** *B. subtilis* 168 (ref. 24) (Bacillus Genetic Stock Center), *E. coli* MG1655 (ref. 31) or BW25113 (ref. 29) were used as parental strains. For the mutant analysis, Keio collections<sup>28</sup> were used. *B. glumae* BGR1 (ref. 61), *P. aeruginosa* PA14 (ref. 64) or *P. polymyxa* E681 (ref. 65) were used for the swarming motility assay.

**Chemicals.** High-purity individual VOCs (>99%) were purchased from Sigma-Aldrich. Each VOC was expressed as an acronym: 3M1BOH (3-methyl-1-butanol), IVA (3-methylbutanoic acid; isovaleric acid), BDOH (butane-2,3-diol), 2,3-BD (butane-2,3-dione; 2,3-butanedione or diacetyl), GA (oxoethanoic acid or glyoxylic acid), 1PNOH (pentan-1-ol; 1-pentanol) and 2M1PAOL (2-methyl-1-propanol).

**Cell culture and exposure to VOCs.** To perform the experiments on the effect of *B. subtilis*, VOCs on *E. coli*, *B. glumae*, *P. aeruginosa* or *P. polymyxa* cells on the I-plate (Falcon, No. 351003) assay were used. Each culture was prepared by inoculating *B. subtilis* or *E. coli* MG1655, BW25113, *B. glumae*, *P. aeruginosa* or *P. polymyxa* in different media as follows: for *B. subtilis* and *P. polymyxa*, TSB (Difco); for *E. coli* and *B. glumae*, LB broth; for *P. aeruginosa*, Nutrient Broth (NB). The culture condition for each bacteria was different: *E. coli*, *B. subtilis* and *P. aeruginosa* were cultured at 37 °C for 16–24 h; *B. glumae* and *P. polymyxa* were cultured at 30 °C for 24–48 h. For further analysis of VOC effects, ~10<sup>5</sup> cells of each bacteria were spread onto TSB-, LB- or NB-agar plates. All growth was monitored by using Ultrospec 2100 pro (AP Biotech).

**Transcriptome analysis.** The analysis was performed using *E. coli* chips described in the KRIBB Genome Encyclopedia of Microbes<sup>66</sup>, except that genes showing VOC-dependent differences in expression levels of ≥2-fold or ≤0.5-fold were considered to be differentially expressed genes. Clustering analysis was performed by MultiExperiment Viewer<sup>67</sup>.

**Construction of single-copy promoter-GFP fusions.** The preparation of backbone plasmid pGEM-T-GFPcm was performed in three consecutive steps. First, the GFP region was amplified from pDSK-GFPuv<sup>68</sup> using primers GFP-F. Second, the chloramphenicol (Cm) cassette fragment was amplified using primers GFP-P1-F and GFP-P2. Finally, GFP-Cm products were prepared by the fusion PCR method as described in a previous report<sup>69</sup>, followed by cloning of the PCR product into the pGEM-T-easy vector (Promega). This resulted in pGEM-T-GFPcm. Chromosomal *yefR*, *aaeX* or *hyaF* promoter fusions were performed by *E. coli* single-gene knock-out method<sup>29</sup>, with pGEM-T-GFPcm and the following primers for each gene: *yefR*-HL, *yefR*-HR, *aaeX*-HL, *aaeX*-HR, *hyaF*-HL and *hyaF*-HR. The construction was confirmed by the gene-specific primers *yefR*-192-F, *aaeX*-76-F and *hyaF*-181-F, or the universal primer GFP-R.

**Identification of transcription factors modulating VOC phenotypes.** The screening of transcription-related factors affected by VOCs was performed using a library of fluorescent transcriptional reporters of *E. coli* (Open Biosystems)<sup>46</sup>. By using the term 'transcription factor or DNA-binding transcription regulator' in both EcoCyc and microarray data, 60 available reporter genes were selected (Supplementary Table S4). Overnight cultures of *E. coli* cells containing individual transcriptional reporters were inoculated into fresh LB media (1/100 dilution) containing kanamycin (50 µg ml<sup>-1</sup>) on 96-well plates in the condition of vaporization of 2,3-BD or GA (1 µM final) for 14 h at 37 °C. The promoter activity was monitored and analyzed<sup>68</sup> with 2104 EnVision multilabel plate readers (PerkinElmer). The normalized promoter activity (fluorescence/OD<sub>600</sub>) to control vector of the reporter system was analysed for all 64 genes from triplicate experiments ( $n = 3$ ,  $P = 0.05$ ).

**Swimming and swarming motility assay.** For swimming motility assays, TSB- or LB-agar (0.3%) plates were separately prepared in either side of I-plate. *B. subtilis* or *E. coli* (~10<sup>5</sup> cells) were spread onto either TSB- or LB-agar plate, respectively, followed by incubation at 37 °C for 12 h and photography of results. The swarming motility assay was performed using a different recipe for the growth of receiver bacteria on the plate: LB agar (LB broth + 0.5% agar + 0.5% glucose) for *E. coli* and *B. glumae*; NB agar (NB broth + 0.5% agar + 0.5% glucose) for *P. aeruginosa*; TSB agar (TSB + 0.9% agar) for *B. subtilis* and *P. polymyxa*. For the assay, *B. subtilis* (~10<sup>5</sup>), cells were spread onto TSB agar, and bacteria were inoculated onto species-specific agar described above. The plates were incubated at either 30 or 37 °C for 12–48 h and photographed. To observe the effect of individual gas-phase VOCs, the specified concentration (indicated in each Figure legend) of 2,3-BD or GA solution was dropped onto the TSB-agar plate instead of emitter *B. subtilis* cells.

**Screening of bacterial antibiotic resistance by VOC treatment.** To identify bacterial antibiotic-resistant phenotypes in response to VOC treatment, *E. coli* cells were subjected to PM analysis (PM plates No. 11–20). *E. coli* MG1655 cells were treated with or without total VOCs from *B. subtilis* (~10<sup>5</sup> cells) for 16 h at 37 °C. The resulting cells were scraped, diluted to OD<sub>600</sub> = 0.3 in PM media and subjected to analysis by PM for 24 h at 37 °C (ref. 39).

**Antibiotic-resistance assays.** To validate VOC-mediated resistance to both ampicillin and tetracycline, *E. coli* cells with or without vaporization of *B. subtilis* VOCs were cultured on I-plate for 12 h. The cells were collected by scraping and diluted to OD<sub>600</sub> = 0.1 in LB broth. Either ampicillin (100 µg ml<sup>-1</sup>) or tetracycline (10 µg ml<sup>-1</sup>) was added to the culture and then grown further for 3 h at 37 °C with vigorous shaking. A fraction of the cultures was spread on LB plates, and the survival ratio was determined by counting the colonies on LB plate. The survival ratio of VOCs-treated and non-VOCs-treated were compared from triplicate experiments ( $n = 3$ ,  $P = 0.05$ ).

**Reverse transcription quantitative PCR.** Total RNA from *E. coli* was prepared using RNeasy plus mini kit (Qiagen) according to the manufacturer's protocol. cDNA synthesis was performed as follows: total RNA (0.5 µg) was used for the template for the RT reaction using ReverTraAce qPCR RT Master Mix with gDNA Remover (Toyobo) according to a previous report<sup>68</sup>. A CFX connector real-time qPCR system (Bio-Rad) was used for RT-qPCR. Reaction mixtures consisted of cDNA, iQ SYBR Green Supermix (Bio-Rad) and 1 pM of each primer set listed in Supplementary Table S6. The PCR parameters were as follows: initial polymerase activation, 3 min at 95 °C, 40 cycles of 10 s at 95 °C, 10 s at 60 °C and 30 s at 72 °C. Conditions were determined by comparing threshold values in a series of dilutions of the RT product, followed by a non-RT template control and a non-template control for each primer pair. Relative RNA levels were calibrated and normalized to the level of *rrsB* (16S rDNA) RNA.

**Statistical analysis.** The statistical analyses were performed as previously described<sup>19</sup> using JMP software. Error bars are defined as s.e.m. in all data analysis.

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### Author contributions

K.-S.K., S.L. and C.-M.R. conceived the study and designed experiments. S.L. and K.-S.K. performed experiments. K.-S.K. and C.-M.R. supervised the project and wrote the paper. All authors analysed the data and discussed the results.

### Additional information

**Supplementary Information** accompanies this paper at <http://www.nature.com/naturecommunications>

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