We recently described a novel bet-hedging mechanism in which the bacterium Sinorhizobium meliloti responds to starvation by forming two discrete cell types via cell division. The old-pole daughter cell retains most of the resource, polyhydroxybutyrate (PHB) and is capable of surviving long-term starvation, while the low-PHB, new-pole daughter cell is capable of quickly resuming growth when starvation ends. Here we present additional data showing that the high-PHB, old-pole cells are similar to bacterial persisters, characterized by metabolic dormancy and antibiotic tolerance. Using two independent methods, we generated clonal populations of S. meliloti that varied in the frequency of the high- and low-PHB phenotypes, and then challenged these populations with ampicillin. Populations containing more high-PHB cells were significantly more antibiotic-tolerant. In a separate experiment, we used GFP fluorescence as a marker of overall metabolic activity. After 24 hours of starvation, new-pole cells were 64% brighter than their old-pole sister cells, demonstrating that the divergence in metabolic rate is rapid.

Bacterial persistence and bet hedging in Sinorhizobium meliloti

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Abbreviations: PHB, poly-3-hydroxybutyrate; GFP, green fluorescent protein; PI, propidium iodide

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We recently demonstrated that, when starved, the bacterium Sinorhizobium meliloti uses energy and carbon stored in the polyester poly-3-hydroxybutyrate (PHB) to divide, producing one high-PHB cell and one or two low-PHB cells.1,2 High-PHB cells, but not low-PHB cells, are capable of surviving more than a year of starvation, but take longer to resume growth once starvation ends.1 The latter result suggests that the high-PHB phenotype may be dormant. In addition to conserving stored resources, dormant bacteria can gain resistance to environmental stresses, like antibiotics and heavy metals, that can kill metabolically active cells.3,4 Most clonal populations of bacteria contain a small proportion of these dormant, antibiotic-tolerant ‘persister’ cells.5 After starvation and differentiation into high and low-PHB phenotypes, are high-PHB cells entering a state of persistence?

We examined this by generating groups of genetically-uniform, starved S. meliloti with varying fractions of high-PHB-phenotype cells, and then exposed these populations to ampicillin at 500 μgmL⁻¹ for 90 minutes. Ampicillin is a broad spectrum β-lactam antibiotic that only affects actively growing bacteria; as a result it is a widely-used screen for persisters.3,5-7 We used two independent methods to generate variation in the frequency of high-PHB cells. First, density gradient centrifugation was used to fractionate five populations of starved cells (with bimodal PHB content) into low (≤1.109 gml⁻¹), medium (1.119–1.109 gml⁻¹) and high (1.138–1.119 gml⁻¹) density as previously described in reference 2; fractions with a higher buoyant density contained more high-PHB cells. Second, we starved S. meliloti at high (5 x 10⁶), medium (5 x 10⁵) and low (10⁵) density for three days; cells starved at lower densities reproduced more and formed populations with fewer high-PHB cells. Viability was assayed flow cytometrically for both experiments as previously described except that we used viability stains propidium iodide in the centrifugation experiment and YO-PRO-1 in the density-of-starvation experiment. Both viability stains are functionally equivalent ways of detecting the failure of dead cells to exclude DNA-binding stains.1 Populations with mostly...
high-PHB rhizobia survived antibiotic exposure significantly better than those with mostly low-PHB rhizobia (Fig. 1A; p < 0.0001, n = 41, r^2 = 0.46, linear regression). S. meliloti strain 1021 is normally killed by ampicillin, but the high-PHB phenotype is apparently tolerant, consistent with results for other bacterial persisters.

We next tested for persistence using an independent method. Because persisters are metabolically dormant, protein production is reduced relative to growing cells. Therefore, bacteria that are engineered to constitutively produce GFP glow less brightly when they become persisters.5

We therefore examined GFP fluorescence in S. meliloti strain pDG71, engineered to constitutively produce GFP.6,9 We followed the fate of initially high-PHB cells that were affixed to a coverslip and incubated in starvation media using methods reported previously in ref. 1. Mean GFP fluorescence per unit area was assessed for 30 dividing cells after 24 hours of starvation. Compared to their old-pole parents, new-pole cells were 65% brighter (Fig. 1B and C; t = 8.35, n = 30, p < 0.0001, matched-pair t-test).

To summarize, high-PHB cells respond more slowly to exogenous resources,1 are more antibiotic-tolerant, and are less GFP fluorescent than low-PHB cells, demonstrating that they are in a state of persistence. The low metabolic rate (persistence, including reduced competition, and C^2 uptake) by flow cytometry. Populations containing a greater fraction of high-PHB cells were less susceptible to ampicillin. There was no significant difference in linear regression slopes (t = 0.8, p = 0.43, n = 41, ANCOVA) or intercepts (t = 1, p = 0.32, n = 41, ANCOVA) for the two methods of manipulating the frequency of high-PHB cells, so a single linear regression was fit to the data. High-PHB old pole cells are less GFP fluorescent (B and C), indicating lower metabolic activity. (B) An initially high-PHB cell dividing during starvation produces a bright, GFP fluorescent new-pole cell and a dim old-pole cell. To increase brightness and lessen background fluorescence, this image was transformed with the ‘Colors; Levels’ command in GIMP v. 2.6. Both left and right panels were transformed identically. This transformation made the cells easier to see and did not qualitatively affect the image. (C) After 24 hours of starvation, GFP fluorescence intensity is 50.8% higher in new-pole cells (t = 5.29, n = 60, p < 0.0001, two-sided t-test). Gray bars represent overlap in new and old-pole distributions. Brightness units are average color depth (ranging from 0–65536) of the green channel from a 16-bit color fluorescence image, higher values indicate greater brightness.

The putative role of the high-PHB phenotype is longevity; old-pole, high-PHB cells can survive more than a year of starvation to resume growth or nodulation if conditions are slow to change. Persistence is an effective way to increase longevity if mortality from antibiotics and other stresses that kill metabolically-active cells is high. The cost of persistence, a missed opportunity to nodulate or consume exogenous resources if conditions improve suddenly, is ameliorated through individual-level bet hedging. Because individual S. meliloti produce both persistent and non-persistent cells when starved, resources that cannot be used by a dormant cell may instead be used by its nearby metabolically-active clonemate.

Bacterial persistence is widely thought of as an insurance policy taken out by a genotype (expressed by a tiny fraction of the clonal population) as a hedge against catastrophe.10 Recent work, however, suggests a more dynamic role for persistence in microbial life-history evolution. Dorr et al.11 have shown that persistence can be induced by exposure to the antibiotic ciprofloxacin. This type of phenotypic plasticity allows for much higher survival than ‘catastrophe insurance’, where only a tiny fraction of the population survives. Theoretical work has proposed alternative explanations for the evolution of persistence, including reduced competition with kin for limited resources12 or as a side-effect of senescence,13 but these hypotheses await experimental confirmation. Our findings, that persistence can be utilized in an individual-level bet hedging strategy that is triggered in response to stress, adds to the diversity of life-history strategies that employ persistence.

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