## **MARINE SCIENCE**

# Kīlauea lava fuels phytoplankton bloom in the North Pacific Ocean

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From June to August 2018, the eruption of Kīlauea volcano on the island of Hawai'i injected millions of cubic meters of molten lava into the nutrient-poor waters of the North Pacific Subtropical Gyre. The lava-impacted seawater was characterized by high concentrations of metals and nutrients that stimulated phytoplankton growth, resulting in an extensive plume of chlorophyll a that was detectable by satellite. Chemical and molecular evidence revealed that this biological response hinged on unexpectedly high concentrations of nitrate, despite the negligible quantities of nitrogen in basaltic lava. We hypothesize that the high nitrate was caused by buoyant plumes of nutrient-rich deep waters created by the substantial input of lava into the ocean. This large-scale ocean fertilization was therefore a unique perturbation event that revealed how marine ecosystems respond to exogenous inputs of nutrients.

ne of the most active volcanoes in the world, Kīlauea on the island of Hawai'i, erupted from April to August 2018. The eruption produced an extensive channel of molten lava that flowed into the ocean at an estimated rate of 50 to 100 m<sup>3</sup> s<sup>-1</sup> for 2 months, from 3 June 2018 until 6 August 2018, ultimately creating an additional 3.5 km<sup>2</sup> of coastal area (1, 2). Three days after lava first entered the ocean, high chlorophyll a concentrations were observed in the vicinity by the MODIS Aqua satellite (Fig. 1). Chlorophyll aenriched waters rapidly expanded in size and within 2 weeks had extended 150 km offshore as a highly prominent filament. To understand the trajectory of the chlorophyll a plume, a numerical simulation was run in which dye was

continuously released at the lava ocean entry site into a flow field estimated from winds and pressure gradients at the sea surface. The results of the simulation indicated that the trajectory of the chlorophyll a filament was dictated by the surface flow field associated with the edge of a cyclonic eddy located to the northeast of Hawai'i (Fig. 1). Chlorophyll a concentrations within this filament remained elevated (>0.2 µg  $chl\ a\ liter^{-1})$  throughout the eruption period and exceeded values observed in this region during the past 15 years (fig. S1). Once the lava flow subsided on 6 August 2018, the chlorophyll a filament subsequently dissipated within a week, indicating that volcanic processes were directly responsible for the observed phytoplankton bloom. Therefore, the spatial distribution of the chlorophyll a filament was used to delineate the extent of lava-impacted waters.

During peak discharge of lava into the ocean (Fig. 1), an oceanographic research expedition was conducted to determine the causes and composition of the resulting phytoplankton bloom (fig. S2). In contrast to the surrounding homogeneous oligotrophic ocean, chlorophyll a concentrations within the lava-impacted plume waters were highly variable, ranging from 0.1 to 0.6 μg chl a liter<sup>-1</sup> near the lava entry site (Fig. 1). The plume also contained elevated concentrations of trace metals, rare earth elements, inorganic nutrients, and particulate material. Within the plume, surface seawater concentrations of silicic acid and trace metals including manganese, iron, and cobalt strongly resembled the composition of Kīlauea reference basalt material (3, 4) (fig. S3). The solubility of these metals

differed markedly. For example, >99% of manganese remained in the dissolved phase (<0.2 µm size fraction) while >99% of iron had precipitated (data S1). Similar to deep-sea hydrothermal vent environments (5), the micromolar levels of iron oxyhydroxide particles promoted the scavenging of rare earth elements and phosphate, thereby causing only a slight enrichment of dissolved phosphate within the plume. The scavenging onto particles also likely contributed to the observed elevated concentrations of particulate phosphorus (table S1). Surprisingly, given the lack of fixed nitrogen species in mantle-derived magmas (6), surface plume waters contained up to  $4 \mu mol$ liter<sup>-1</sup> nitrate, significantly higher than background concentrations in the surrounding oligotrophic waters (<0.01 μmol liter<sup>-1</sup>; table S1). To examine the fate of nutrients at the broader scale, we used a biogeochemical model embedded within a high-resolution regional physical flow field (fig. S4). The modeled input of silicic acid, iron, nitrate, and phosphate along the southeast coast of Hawai'i was sufficient to simulate a phytoplankton bloom both near the lava entry site and hundreds of kilometers downstream, sustained from the recycling of lava-associated nutrients (see supplementary text).

The shipboard observations of the transition zone between lava-impacted plume waters and the ambient oligotrophic ocean revealed steep biogeochemical gradients, with silicic acid, nitrate, metals, and chlorophyll decreasing from peak values to typical background oligotrophic ocean concentrations over a distance of approximately 2 km (Fig. 2). To further investigate the lava-impacted waters and the transition zone, particularly in regard to subsurface features, an autonomous SeaGlider conducted vertical profiles in the vicinity (fig. S5). The SeaGlider identified several maxima in chlorophyll a concentrations located just beneath the mixed layer, which varied between depths of 5 to 50 m in the water column (fig. S6B). Coincident high values of particle backscatter at these depths (fig. S6C) helped distinguish these near-surface chlorophyll maxima from the deep chlorophyll maxima that are typical of stratified oligotrophic regions, which are due to changes in the phytoplankton chlorophyll:carbon ratios rather than to phytoplankton biomass (7).

Within the lava-impacted waters, plankton carbon biomass [based on measurements of carbon per unit biovolume (8) (see supplementary text)] increased by a factor of 2 to 3 relative to surrounding oligotrophic waters, and primarily comprised large chain-forming diatoms in the genus Skeletonema (37% of estimated total carbon biomass; Fig. 3, A and B). Skeletonema sp. are commonly found in coastal waters of the Hawaiian Islands, and their strong response to lava input reflects a combination of their initial population size and intrinsic traits that allow them to capitalize on the nutrient inputs associated with lava (9). Accompanying the increases in diatom abundance was an increase in zooplankton nauplii (11% of total biomass) as well as other phytoplankton, such as haptophytes (represented

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primarily by *Phaeocustis*) and radiolarians (represented primarily by Sphaeropylolena). Although large microorganisms dominated the plankton biomass, the typical oligotrophic phytoplankton community of Prochlorococcus, Synechococcus, and picoeukaryotes remained numerically abundant and together represented 18% of estimated total carbon biomass. The coexistence of diatoms and Prochlorococcus supports the hypothesis that the increase in diatom biomass in response to nutrient inputs in the open ocean does not lessen abundant nano- and picophytoplankton populations (10). Highly elevated rates of biological production were detected within the plume waters. Rates of <sup>14</sup>C-bicarbonate assimilation ranged from 2.5 to 3.5 µmol C liter<sup>-1</sup> day<sup>-1</sup> (compared to 0.5 to 0.6 µmol C liter<sup>-1</sup> day<sup>-1</sup> in ambient oligotrophic waters), and gross primary production based on the <sup>18</sup>O-H<sub>2</sub>O method ranged from 4 to 6  $\mu mol~O_2~liter^{-1}~day^{-1}$  (compared to 0.9 to 1.1 µmol O2 liter 1 day in ambient oligotrophic waters) (fig. S7). At the most near-shore sampling station, the high rates of biological production coincided with high rates of microbial respiration that reached  $5.1 \pm 0.6 \,\mu\text{mol O}_2 \,\text{liter}^{-1}$  $day^{-1}$  (mean  $\pm$  SD; fig. S7).

Surprisingly, the lava-impacted waters did not support an abundance of diazotrophs, nor did they lead to high rates of nitrogen fixation (fig. S7) despite the presence of sufficient amounts of iron and phosphorus expected to enhance biological nitrogen fixation (11, 12). Instead, the metabolic response to lava-impacted waters indicated that nitrate, rather than biological nitrogen fixation, was the key driver of the observed phytoplankton bloom. Detailed analysis of functional gene responses specific to diatoms in the lava-impacted waters relative to surrounding water revealed significant increases in relative transcript abundance for genes associated with growth processes such as ribosomal biogenesis, photosynthesis, and adenosine triphosphate synthesis, whereas decreases were detected in relative transcript abundance for genes involved in the assimilation of nitrogen (fig. S8F). These results are consistent with prior observations of relief from nitrogen limitation in diatoms (9, 13, 14). Shifts in relative transcript abundance specific to Skeletonema largely mirrored those observed in the whole diatom community (fig. S8G). Within the plume, Skeletonema genes involved in the assimilation of nitrogen, including those that encode nitrate and ammonium transporters, nitrate reductase, and glutamate synthase, showed significant decreases in relative transcript abundance, whereas significant increases in relative transcript abundance were observed for genes involved in photosynthesis (psbA, psbC) and carbon fixation (rbcS, rbcL) (Fig. 3C). The relative abundance of Skeletonema transcripts related to iron transport decreased in the lavaimpacted waters (Fig. 3C), consistent with the high concentrations of iron within the plume (data S3).

The composition and physiology of the phytoplankton within the plume therefore suggested that nitrate availability was a key component

of the phytoplankton bloom. We confirmed this hypothesis by determining the isotopic composition of suspended particulate nitrogen  $(\delta^{15}\text{N-PN})$  and nitrate  $(\delta^{15}\text{N-NO}_3^- \text{ and } \delta^{18}\text{O-NO}_3^-)$ within the plume. The  $\delta^{15}$ N-PN signature ranged from 4.5 to 4.8 per mil (‰), whereas a  $\delta^{15}$ N signature closer to 0% would have represented a larger contribution from biological nitrogen fixation. Moreover, the  $\delta^{15}\text{N-PN}$  values were ~5‰ lower than the  $\delta^{15}$ N-NO<sub>3</sub> within the lavaimpacted plume waters, with  $\delta^{15}\text{N-NO}_3^{-}$  values of 9.4  $\pm$  1.3‰ and  $\delta^{18}$ O-NO<sub>3</sub> values of 7.0  $\pm$ 1.6‰ (mean  $\pm$  SD, n = 16). This is consistent with the magnitude of isotopic fractionation previously observed in phytoplankton cultures (15). The nearly 1:1 correlation observed between  $\delta^{15}$ N-NO<sub>3</sub> and  $\delta^{18}$ O-NO<sub>3</sub> in lava-impacted waters (Fig. 4A) is an additional signature of biological nitrate uptake. Thus, molecular and isotopic markers of phytoplankton nitrogen utilization are consistent with nitrate supporting the elevated phytoplankton biomass in the lava-impacted

Although there was clear evidence for the biological importance of high nitrate concentrations in lava-impacted waters, the pathway of

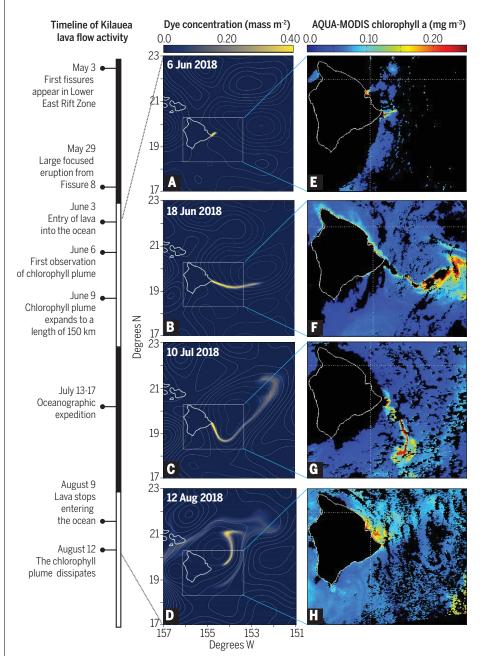


Fig. 1. Timeline of Kīlauea eruptive activity focusing on events related to lava entering the ocean. (A to D) Simulated yellow dye projections of lava-impacted waters overlain on contours of sea surface height. (E to H) Satellite observations showing the formation and subsequent disappearance of the chlorophyll a plume during June to August 2018.

nitrate supply was not immediately clear. The absence of fixed nitrogen species in mantlederived magmas means that the lava itself is not a source of nitrate. Yet the strong covariation of nitrate with other lava-derived elements (Fig. 2) implies a mechanistic coupling between lava input and nitrate supply. Chemical fixation of N<sub>2</sub> can occur at temperatures above 1000°C with either O2 or H2 as the reactant (16, 17), enhanced by the abundance of metal catalysts derived from the lava. However, this process would result in nitrate with  $\delta^{15}$ N-NO $_3^-$  values of  $\sim 0 \pm 4\%$  (17), rather than the  $\delta^{15}$ N-NO $_3^-$  of 9.4 ± 1.3‰ measured here. Groundwater discharge is an alternative source of high concentrations of nitrate to nearshore waters of the Hawaiian Islands (18, 19). However, the isotopic composition of nitrate in local groundwater (18) does not overlap with our measurements (Fig. 4A), and salinity anomalies expected from enhanced groundwater flow were not detected within the plume.

We instead hypothesize that injection of lava below the euphotic zone created buoyant plumes of nutrient-rich deep water bringing nitrate to

the surface (Fig. 4B). Prior measurements of nitrate isotopic composition at depths of 300 m have  $\delta^{15}$ N-NO<sub>3</sub> values of 6 to 7‰ and  $\delta^{18}$ O- $NO_3$  values of 3‰ (20, 21) (fig. S9). This isotopic composition is consistent with a nitrate source to surface plume waters that subsequently underwent nitrate assimilation, leading to a 1:1 enrichment in  $\delta^{15}$ N-NO<sub>3</sub><sup>-</sup> and  $\delta^{18}$ O-NO<sub>3</sub><sup>-</sup> (Fig. 4A). Upwelling of waters from below the euphotic zone containing low O2 would also explain the under saturation of  $\mathrm{O}_2$  in lava-impacted surface waters at the head of the plume (182  $\mu mol\ kg^{-1}$ compared to  $205\,\mu\text{mol}$  kg $^{-1}$  in non-lava-impacted waters), which is otherwise difficult to account for by biological or abiotic processes alone (table S1 and fig. S10). Historically, much smaller lava ocean-entry events along the Kīlauea coastline were sufficient to create surface "roiling" when the upward motion of water heated by submarine lava intrusions entrained subsurface water (22). It is conceivable that the lava flow in summer 2018 reached the mesopelagic environment in the early stages of the phytoplankton bloom, facilitated by the high flow rate and ex-

Longitude (deg W) -155.0-154.5 9.5 Α Latitude (deg N) 19.0 8.0 C Nitrate+Nitrite (µM) Nitrate+nitrite 0.8 Chlorophyll *a* (µg L<sup>-1</sup>) 0.6 Silicic acid 0.4 0.4 0.2 Synechococcus (106 cells L-1) 0.0 0.0 4 20 160 D G Picoeukaryotes (10° cells L-1) 15 10 icoeukarvotes Synechococcus 0 0 160 40 Prochlorococcus (10<sup>6</sup> cells L<sup>-1</sup>) Copper (nM) 30 20 10 100 15:00 16:00 16:00 17:00 15:00 17:00 18:00 18:00 Local Time on 14 July Local Time on 14 July

Fig. 2. Transition from lava-impacted waters to the oligotrophic ocean. (A) Map of Hawai'i showing sampling location. (B) Cruise track depicted by chlorophyll a concentrations. The location of the transition zone (approximately 2 km in length) is marked by the two arrows; the circles indicate the beginning and end of the datasets shown in (C) to (H). (C to H) The changes in water column properties from the lava-impacted waters (solid circle) to oligotrophic conditions (open circle) are shown for chlorophyll a (C), Synechococcus and

tremely steep offshore bathymetry with water column depths of >300 m less than 1 km offshore. Subsequent remotely operated vehicle observations during September 2018 confirmed that lava intrusions ultimately reached depths of 725 m, and low-temperature venting directly associated with the fresh lava deposits was also observed (23). Upwelled waters from this depth would contain sufficient concentrations of nitrate to fuel the observed phytoplankton bloom. After the lava flow substantially decreased on 6 August, the much smaller flow that lasted for an additional month was insufficient to cause buoyant plumes of nutrient-rich deep water.

The large-scale ocean entry of lava during the 2018 eruption of Kīlauea volcano created a large and biologically productive surface plume that was observed hundreds of kilometers offshore and persisted for the duration of the eruption. This strong biological response depended not only on the lava itself but also on an exogenous source of nitrate, which is believed to have derived from lava entering the deep ocean, creating buoyant plumes of nitrate-rich deep water into the nitrogen-starved surface ocean. It is possible that this mechanism has led to similar ocean fertilization events in the past, associated with the formation of the Hawaiian Islands and other major volcanic eruptions. Although greater by a factor of 10 than mean lava production rates at Kīlauea over the past 50 years (1), the magnitude of the 2018 eruption ( $\sim$ 0.8 km<sup>3</sup>) only represents lower-bound estimates of lava production rates during the formation of the Ontong-Java Plateau and other large igneous provinces (1 to >20 km3 per year) that occurred over thousands to millions of years (24, 25). Such major volcanic episodes coincide with the rapid release of carbon dioxide, global warming, and stratification of the deep ocean (26). Depending on their location, sustained eruption on this scale could also facilitate a large flux of nitrate from the mesopelagic ocean and perturb regional ocean circulation and biogeochemistry. Marine ecosystem responses to such substantial additions of nitrate are rarely observed or sampled in real time. The expedition in July 2018 provided a unique opportunity to see first-hand how a massive input of exogenous nutrients alters marine ecosystems attuned to oligotrophic conditions.

# REFERENCES AND NOTES

- C. A. Neal et al., Science 363, 367-374 (2019).
- U.S. Geological Survey Hawaiian Volcano Observatory, "Preliminary analysis of the ongoing Lower East Rift Zone (LERZ) eruption of Kīlauea Volcano: Fissure 8 Prognosis and Ongoing Hazards" (Cooperator Report to Hawaii County Civil Defense, 15 July 2018); https://volcanoes.usgs.gov/vsc/file\_ mngr/file-185/USGS%20Preliminary%20Analysis\_LERZ\_7-15-18\_ v1.1.pdf.
- K. Govindaraju, Geostand. Newsl. 18, 1-158 (1994).
- J. A. Resing, F. J. Sansone, Geochim. Cosmochim. Acta 66, 1925-1941 (2002)
- R. A. Feely et al., Geochim. Cosmochim. Acta 60, 2297-2323 (1996)
- Y. Li, H. Keppler, Geochim. Cosmochim. Acta 129, 13-32 (2014).
- J. Cullen, Annu. Rev. Mar. Sci. 7, 207-239 (2015).
- L. Campbell, H. A. Nolla, D. Vaulot, Limnol. Oceanogr. 39, 954-961 (1994).
- H. Alexander et al., Proc. Natl. Acad. Sci. U.S.A. 112 F5972-F5979 (2015)

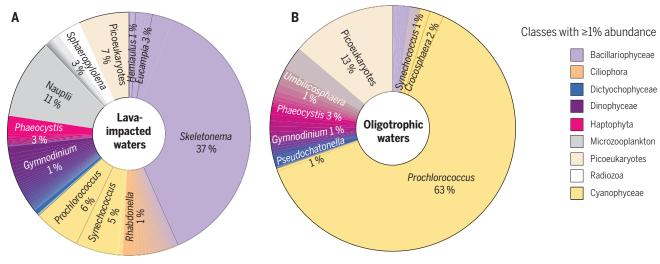
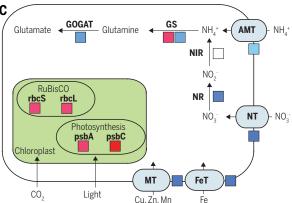


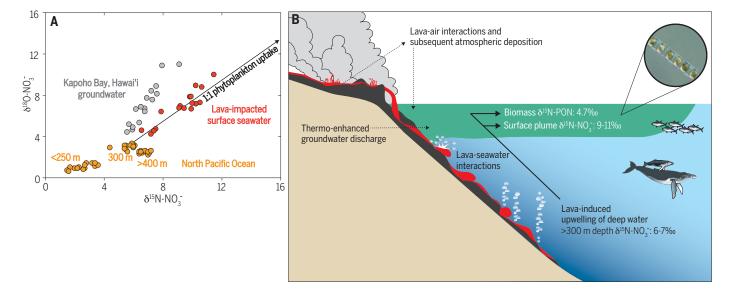
Fig. 3. Plankton community composition and biomass. (A) Estimated carbon biomass of the plankton population in lava-impacted waters. (B) Estimated carbon biomass in the surrounding oligotrophic environment. (C) Significant changes in relative transcript abundance for the diatom Skeletonema, comparing lava-impacted waters with surrounding oligotrophic waters. Note that Skeletonema has multiple glutamine synthetase sequences with distinct transcriptional responses.



AMT (ammonium transporter; K03320)
FeT (Fe complex transporter; K02013)
GOGAT (glutamate synthase; K00264,4,6)
GS (glutamine synthetase; K01915)
MT (Cu, Zn, Mn transporter; K11706)
NIR (nitrite reductase; K00366)
NR (nitrate reductase; K10534)
NT (nitrate transporter; K02575)
psbA (PSII P680 reaction center; K02703)
psbC (PSII CP43 chlorophyll apoprotein; K02705)
rbcL (RuBisCO large chain; K01601)
rbcS (RuBisCO small chain; K01602)

5.0

Log, fold change



**Fig. 4. Nitrate origins.** (**A**)  $\delta^{15}$ N-NO $_3^-$  and  $\delta^{18}$ O-NO $_3^-$  along the lava-impacted surface plume waters (red circles) compared with regional subsurface values (yellow circles) (*20*) and local groundwater values (gray circles) (*18*). The line originating from 300 m  $\delta^{15}$ N-NO $_3^-$  follows a trajectory consistent with an isotope effect of 5‰ for NO $_3^-$  assimilation, which would elevate  $\delta^{15}$ N-NO $_3^-$  and  $\delta^{18}$ O-NO $_3^-$  equally (i.e., a slope of 1).

 $(\mathbf{B})$  Schematic of potential nitrate sources during the 2018 eruption, including upwelling of  $\geq 300$  m subsurface water, near-shore groundwater discharge, and abiotic  $N_2$  fixation. Partitioning of upwelled nitrate into an isotopically lighter pool of particulate nitrogen and an isotopically heavier pool of residual surface nitrate is highlighted as the most direct explanation of the observations.

- 10. R. T. Barber, M. R. Hiscock, Global Biogeochem. Cycles 20, GB4S03 (2006)
- 11. D. M. Karl, R. M. Letelier, Mar. Ecol. Prog. Ser. 364, 257-268 (2008).
- 12. C. L. Follett, S. Dutkiewicz, D. M. Karl, K. Inomura, M. J. Follows, ISME J. 12, 1543-1557 (2018).
- 13. S. J. Bender, C. A. Durkin, C. T. Berthiaume, R. L. Morales, E. Armbrust, Front. Mar. Sci. 1, 3 (2014).
- 14. M. J. Harke, A. R. Juhl, S. T. Haley, H. Alexander, S. T. Dyhrman, Front. Microbiol. 8, 1279 (2017).
- 15. J. Granger, D. M. Sigman, J. A. Needoba, P. J. Harrison, Limnol. Oceanogr. 49, 1763-1773 (2004).
- 16. B. Huebert et al., Biogeochemistry 47, 111-118 (1999).
- 17. J. H. Carrillo, M. G. Hastings, D. M. Sigman, B. J. Huebert, Global Biogeochem. Cycles 16, 24-1-24-16 (2002).
- 18. T. N. Wiegner, A. U. Mokiao-Lee, E. E. Johnson, Mar. Pollut. Bull. 103, 63-71 (2016).
- 19. J. M. Bishop, C. R. Glenn, D. W. Amato, H. Dulai, J. Hydrol. Reg. Stud. **11**, 194-218 (2017).
- 20. K. L. Casciotti, T. W. Trull, D. M. Glover, D. Davies, Deep-Sea Res. 55, 1661-1672 (2008).
- 21. A. N. Knapp, D. M. Sigman, F. Lipschultz, A. B. Kustka, G. Capone, Global Biogeochem. Cycles 25, GB4004 (2011).
- 22. F. J. Sansone, J. A. Resing, J. Geophys. Res. Oceans 100, 13555-13569 (1995).
- 23. C. R. German et al., Time series study of hydrothermal venting at Lō'ihi Seamount following the 2018 Kīlauea eruption. EOS Trans. Am. Geophys. Union (abstr.), V14A-07
- 24. S. E. Bryan et al., Earth Sci. Rev. 102, 207-229 (2010).
- 25. M. F. Coffin, O. Eldholm, Geology 21, 515-518 (1993).
- 26. P. B. Wignall, Earth Sci. Rev. 53, 1-33 (2001).

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A.M.H., and G.V.H. processed and curated the data. M.B., S.F., K.M.B., and S.T.W. conducted the rate measurements including <sup>18</sup>O-GPP, <sup>14</sup>C assimilation, microbial respiration, and nitrogen fixation. R.K.F. C.P.F., and K.M.B. led the nutrient analyses. T.J.B., R.T., and J.R.C. led the majority of the water column biogeochemical sampling. A.N.K. analyzed the nitrate isotopes. M.J.H. and S.T.D. measured the eukaryote metatranscriptomics, A.K.B. and A.F.I. measured the metabolites. S.D., O.J., M.J.F., and C.N.H. conducted the MITgcm, DARWIN, and OSCAR modeling. K.A.T.-K., B.A.H., and J.P.Z. measured nifH abundances. B.B. analyzed the SeaGlider data. S.T.W. and N.J.H. co-wrote the manuscript with contributions from all authors. Competing interests: The authors declare no competing interests. Data and materials availability: Data and pertinent methodologies are provided in the supplementary materials. The hydrographic and biogeochemical datasets are freely available online at http://scope.soest.hawaii. The eukaryote metatranscriptomic sequence data are deposited in the Sequence Read Archive through the National Center for Biotechnology Information under accession number PRJNA529177.

### SUPPLEMENTARY MATERIALS

science.sciencemag.org/content/365/6457/1040/suppl/DC1 Materials and Methods Supplementary Text Figs. S1 to S11 Table S1 Movie S1 Data S1 to S3 References (27-59)

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Ocean greening off Hawai'i

From June to August 2018, the eruption of Kilauea volcano triggered a diatom-dominated phytoplankton bloom. Wilson et al. set sail to sample the plume, deploying subsea gliders and using satellite monitoring to measure the dynamics of this rare event in the nutrient-poor Pacific (see the Perspective by Ducklow and Plank). They found subsurface chlorophyll maxima not visible by remote sensing, performed transcriptome and N isotope marker analysis, and measured nutrients, partitioning of biomass into different organisms, and primary production. Much of the data are corroborated by physical modeling of the ocean dynamics. The authors conclude that the plume was fed by the lava heating subsurface water and triggering upwelling of deepwater nutrients to the surface rather than by direct injection of micronutrients from lava.

Science, this issue p. 1040; see also p. 978

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