Superhydrophobic diving flies (*Ephydra hians*) and the hypersaline waters of Mono Lake

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The remarkable alkali fly, *Ephydra hians*, deliberately crawls into the alkaline waters of Mono Lake to feed and lay eggs. These diving flies are protected by an air bubble that forms around their superhydrophobic cuticle upon entering the lake. To study the physical mechanisms underlying this process we measured the work required for flies to enter and leave various aqueous solutions. Our measurements show that it is more difficult for the flies to escape from Mono Lake water than from fresh water, due to the high concentration of Na_2CO_3 which causes water to penetrate and thus wet their setose cuticle. Other less kosmotropic salts do not have this effect, suggesting that the phenomenon is governed by Hofmeister effects as well as specific interactions between ion pairs. These effects likely create a small negative charge at the air–water interface, generating an electric double layer that facilitates wetting. Compared with six other species of flies, alkali flies are better able to resist wetting in a 0.5 M Na_2CO_3 solution. This trait arises from a combination of factors including a denser layer of setae on their cuticle and the prevalence of smaller cuticular hydrocarbons compared with other species. Although superbly adapted to resisting wetting, alkali flies are vulnerable to getting stuck in natural and artificial oils, including dimethicone, a common ingredient in sunscreen and other cosmetics. Mono Lake’s alkali flies are a compelling example of how the evolution of picoscale physical and chemical changes can allow an animal to occupy an entirely new ecological niche.

superhydrophobicity | insects | Hofmeister series | bubbles | biomechanics

In late summer the shores of Mono Lake, California, are bustling with small flies, *Ephydra hians*, which crawl underwater to feed and lay eggs (Fig. 1A). Their unusual behavior was eloquently described by Mark Twain during his travels to Mono Lake (1):

You can hold them under water as long as you please—they do not mind it—they are only proud of it. When you let them go, they pop up to the surface as dry as a patent office report, and walk off as unconcernedly as if they had been educated especially with a view to affording instructive entertainment to man in that particular way.

Although Twain’s observations are over 150 years old, we still do not understand the chemistry and physics underlying the ability of these flies to resist wetting as they descend below the water surface. Alkali flies are found on nearly every continent and fulfill an important ecological role by transforming the physically harsh environments of alkaline lake shorelines, including the Great Salt Lake in Utah and Albert Lake in Oregon, into important wildlife habitats (2). Aside from the flies, only algae, bacteria, and brine shrimp tolerate Mono Lake’s water, which is three times saltier than the Pacific Ocean and strongly alkaline ($\text{pH} = 10$) due to the presence of sodium bicarbonate and carbonate. For the past 60,000 y, Mono Lake has had no outlet (3), driving a steady increase in the concentration of mineral salts through a yearly evaporation of 45 inches (4). Calcium from natural springs underneath the lake’s surface reacts with the carbonate-rich water, precipitating calcium carbonate in the form of underwater towers called tufa. Alkali flies crawl underwater by climbing down the surface of the tufa, which have become exposed due to falling lake levels (Fig. 1B and Movies S1 and S2).

For the alkali fly, staying dry is paramount to their survival; if they do get wet in Mono Lake, a thin film of minerals dries on their

cuticle, which makes them more likely to be wetted in subsequent encounters with the water. Like most insects, the flies are covered in a waxy cuticle festooned with tiny hairs (setae). As in water striders (5), these hydrophobic hairs trap a layer of air, so that as a fly crawls into the water an air bubble forms around its body and wings. The bubble protects the flies from the salts and alkaline compounds present in the lake and also serves as an external lung (6), allowing flies to spend up to 15 min underwater crawling to depths of 4–8 m (7). Once finished feeding or laying eggs the flies either crawl to the surface or let go of the substratum and float up. As noted by Twain (1), the bubble pops when it hits the air–water interface, depositing its inhabitant safe and dry on the water’s surface (Fig. 1C and Movies S3 and S4). In this paper we describe the physical and chemical properties that make the alkali flies uniquely able to form these protective bubbles in Mono Lake’s dense and alkaline waters.

As a preamble to our measurements, we briefly review the physics of solid–liquid interactions. On smooth surfaces, the shape of an adhering liquid droplet may be described by the contact angle, with larger contact angles corresponding to less-wettable surfaces (Fig. 1D). The contact angle for a smooth piece of waxy insect cuticle is typically $100\text{--}120^\circ$, similar to paraffin wax, and close to the theoretical maximum (8, 9). On rough surfaces, like that of an alkali fly, a liquid drop can exist in two different states. In the Cassie–Baxter state, air pockets fill the space between roughness elements (10), resulting in “superhydrophobicity” [a.k.a. the “lotus-effect” (11)]. In the Wenzel state, the liquid replaces the air pockets (12), resulting in a fully wetted surface. Whether a liquid–surface interface exists in the Cassie–Baxter or Wenzel state is a complex function of the surface’s physical and chemical structure,

Significance

Superhydrophobic surfaces have been of key academic and commercial interest since the discovery of the so-called lotus effect in 1977. The effect of different ions on complex superhydrophobic biological systems, however, has received little attention. By bringing together ecology, biomechanics, physics, and chemistry our study provides insight into the ion-specific effects of wetting in the presence of sodium carbonate and its large-scale consequences. By comparing the surface structure and chemistry of the alkali fly—an important food source for migrating birds—to other species we show that their uniquely hydrophobic properties arise from very small physical and chemical changes, thereby connecting picoscale physics with globally important ecological impacts.

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Fig. 1. Mono Lake's alkali flies must exert up forces to 18 times their body weight to crawl underwater to feed and lay eggs. (A) Close up of an alkali fly under water. (B) Image sequence of a fly crawling into the water (Movies S1 and S2). (C) Image sequence of a fly floating upward to the surface inside its air bubble (Movies S3 and S4). (D) Illustrations of a water droplet on smooth and rough surfaces.

the chemistry of the solution, and the interactions between the surface and the liquid. In the case of an alkali fly crawling into water, the combination of hydrophobic wax and setose surface favors the Cassie–Baxter state, rendering the flies superhydrophobic (6, 9, 13), with contact angles approaching 180°. Other insects that make air–water transitions, including spiders and beetles, sport small patches with superhydrophobic properties used for plastron respiration (6).

Results

To investigate which chemical and physical properties of the flies and Mono Lake water (MLW) influence the formation of the air bubble that protects them from the mineral-rich water we built an optical force sensor (Fig. 2A), which we used in a manner similar to the Wilhelmy balance method (14) to measure the forces required for flies to enter and exit different solutions. We glued the flies to a tungsten beam (0.26-mm diameter) and slowly submerged them using a linear motor (speed 0.3 mm·s^{−1}). The average peak force required for flies to enter MLW was ~1 mN, roughly 18 times the body weight of the 5.5-mg flies (Fig. 2B and Fig. S1A). The force required to enter the water varied with body orientation, with a minimum at a vertical, headfirst orientation (Fig. S1B). This corresponds with our observations of flies at Mono Lake, which tend to enter the water by crawling down 45°–90° surfaces.

In pure water, the work required to submerge the fly is largely recovered when it is pulled out of the water—the surface tension of the bubble stores the potential energy much like a spring. Thus, we use the term “recovered work” (Fig. 2C) as a measure of how easy it is for the flies to escape the water. A positive value indicates a net upward force that pushes the fly out of the water, whereas a negative value indicates that the fly is partially wetted and trapped by surface tension at the air–water interface. With increasing concentration of MLW from 0 to 200% we found that the recovered work decreases, despite the increase in solution density (Fig. 2D).

MLW contains a number of salts including NaCl, Na₂SO₄, and K₂SO₄, as well as the alkali components sodium bicarbonate and boric acid (15). To determine which of these components most influences the recovered work we made two solutions, each containing double the natural concentration of either the salts or alkali compounds. Sodium bicarbonate is an alkali buffer in which the ratios of CO₃^{2−}, HCO₃^{1−}, and H₂CO₃ are coupled to pH according to the Henderson–Hasselbalch equation. To achieve a pH equal to that of MLW (pH = 10), we used a molar ratio of NaHCO₃ to Na₂CO₃ of 0.8. Compared with MLW, recovered work was higher for the salt solution, whereas it was significantly lower for the alkali solution (Fig. 2E), implying that the alkali compounds make it more difficult for the flies to escape the water.

Fig. 2. High concentrations of sodium carbonate make it more difficult for flies to escape from MLW. (A) Diagram of optical force sensor. Forces on the fly deflect the beam, shifting the shadow cast by an LED, which is detected by a photo detector. (B) Force vs. distance traveled (normalized to fly height) for 20 CO₂ anesthetized flies dipped into MLW (bold: mean). Positive values correspond to upward forces (Movie S5). (C) One example trace from A. Shading indicates the amount of work done on the fly as it exits the water, which we term recovered work. (D) Recovered work for different concentrations of MLW, ranging from pure deionized water to double-strength MLW (produced via evaporation): 0% (n = 30), 50% (n = 20), 100% (n = 50), 200% (n = 20). Black line: expected recovered work based on the measurements in pure water and the increase in solution density. Red line: data regression (P = 0.003; r² = 0.08). (E) Recovered work for salt or alkali solutions. Salt solution (double concentration of the salts in Mono Lake): 1.3 M NaCl, 0.2 M Na₂SO₄, 0.04 M K₂SO₄, and 1.8 mM K₃PO₄. Alkali solution (double concentration of the alkali compounds in Mono Lake): 0.35 M NaHCO₃, 0.44 M Na₂CO₃, and 0.09 M boric acid. (F) Recovered work for a 0.5 M NaHCO₃ buffer solution at three different pH values. (G) Same as F, showing only the subset of 10 flies that were dipped in the order of increasing pH. (H) Recovered work for standard MLW and MLW neutralized to pH 7 with HCl. (I) Recovered work for pure water and 5 mM NaOH, at the same pH as the carbonate buffer in E and F. (J) Recovered work for HCl-neutralized 0.5 M carbonate buffer, 0.16 M carbonate buffer, and 0.5 M carbonate buffer. In this experiment all flies were dipped in the order from left to right. (D–J) Shading indicates bootstrapped 95% confidence intervals. Nonoverlapping shading generally corresponds to statistical significance of P < 0.02. Resampling test statistics are given for cases in which differences are not obvious. D–G and H–J come from two separate collections, which may explain the slightly higher water repellency in H–J than would be expected based on D. The order of solutions into which the flies were dipped was alternated for each set of experiments unless otherwise noted. E–J, n = 20. The Bond number for flies in the 0.5 M Na₂CO₃ solution is 0.46, indicating that surface tension forces are dominant (Supporting Information).

Fig. 6. Oils, notably dimethicone (a common ingredient in sunscreens and cosmetics), annihilates alkali flies' superhydrophobic properties. (A) Force traces of five flies dipped into MLW (blue), as in Fig. 2A, and MLW containing dissolved Neutrogena Ultra Sheer SPF 50 sunscreen (brown). To prepare the solution, 160 mg of sunscreen (B) was rubbed into both hands and allowed to set for 15 min. We then poured 300 mL of MLW over one hand, and used the run-off solution. (C) Work done on flies to escape MLW or pure water, with or without Neutrogena Ultra Sheer SPF 50 sunscreen (NS) run-off. (i) Sunscreen set for 5 min before rinsing with MLW. (ii) Hands thoroughly washed with soap and rinsed with warm water before rinsing. (iii) Sunscreen set for 5 min before rinsing with pure water. (iv) Sunscreen set for 15 min before rinsing with MLW. (D) Fraction of flies stuck to surface when dunked into pure water with increasing concentrations of Neutrogena sunscreen. $n = 30$ flies for each condition; mean and 95% confidence intervals are shown. (E) Fraction of flies stuck when dunked into pure water with different types of sunscreen (20 mg each). Same procedure as D. (F) To test whether dimethicone is sufficient to trap flies on the water's surface we applied 0, 2, or 10 μL to the surface and performed the same test described in D.

that the three which had a deleterious effect all contained dimethicone (Fig. 6E), which was absent from the three neutral brands (Fig. S3C). We repeated the dunk assay with pure water after applying a surface film of 0, 2, or 10 μL of dimethicone (viscosity 5 cSt; Sigma-Aldrich). As little as 45 nL of dimethicone per cm^2 of water was enough to trap 50% of the flies (Fig. 6F). Other artificial polymers such as trimethylsiloxysilicate and vp/hexadecene copolymer are likely also problematic (Fig. S3C).

Discussion

Through a series of experiments with solutions varying in salinity, pH, and charge density, we showed that a high concentration of Na_2CO_3 makes it more difficult for alkali flies to escape from the surface of water by facilitating the penetration of water into the air pockets between individual hairs. The effect of Na_2CO_3 is surprising, considering that, like most salts, Na_2CO_3 increases the surface tension (+2% for 1 M solution) (20) as well as the density of water. Both effects should theoretically increase the recovered work by making large bubbles more stable and providing a larger buoyancy force. The fact that other salts, including K_2CO_3 , have a significantly smaller effect demonstrates that this phenomenon involves interactions of specific ion pairs, and not with Debye-Hückel or Derjaguin-Landau-Verwey-Overbeek models. Instead, we offer a model based on the Hofmeister series to explain this phenomenon, which we term "ion-facilitated wetting."

In 1888, Hofmeister (21) discovered that certain ions are more likely to precipitate proteins out of egg white. The same order of compounds—now known as the Hofmeister series—explains a wide range of phenomenon including solubility and surface tension (22). The order of anions is

Ions to the right of Cl^- (kosmotropes) attract large hydration shells that structure the surrounding water, thereby increasing

surface tension and driving ions away from the air-water interface. Ions to the left of Cl^- (chaotropes) have the opposite effect; they tend to accumulate at the surface of an air-water interface (23). Cations are arranged in following order:

Again, ions on the right have larger hydration shells, although anions generally have a more pronounced effect than cations. The underlying principles that give rise to the Hofmeister series are not well understood; however, the sequence is generally correlated with the ratio of ionic charge to ionic radii (Fig. 7A) (values from ref. 24).

Our result that Na_2CO_3 , but not K_2CO_3 , has a strong effect on wetting suggests that ion-facilitated wetting is dependent on the precise combination of cations and anions. Typically, Hofmeister effects of anions and cations are considered to be independent of one another (25); however, some phenomena are known to depend on ion specific pairs, such as the inhibition of bubble coalescence (26, 27). The physical basis of ion-facilitated wetting and bubble coalescence are likely related, because both have macro-scale consequences similar to those caused by surfactants (which increase wetting and decrease bubble coalescence), yet they operate through a completely different mechanism. However, our results cannot be explained by the current models of bubble coalescence. We propose that the relative distance of the cations and anions from the air-water interface plays a crucial role. Fig. 7B illustrates how, as a rough approximation, CO_3^{2-} is on average situated 0.04 nm closer to the surface than Na^+ in Na_2CO_3 solution. This suggests that the carbonate ions will have a larger influence on the surface characteristics in the presence of Na^+ , giving the air-water interface a slight negative charge. Taking this relative distance as well as the ion charge density into account explains 77% of the variance in the correlation with the likelihood of flies' becoming trapped at the surface (Fig. 7C). In contrast, in a K_2CO_3 solution the ions are nearly equidistant from the surface. According to our theory, CaCO_3 , MgCO_3 , and Li_2CO_3 would have even stronger effects on wetting, as the hydration shells of these cations are even larger. However, these salts are only soluble

Fig. 7. Ion-specific interactions including relative hydration shell size and charge density help to explain ion-facilitated wetting. (A) Hofmeister series of anions is correlated with the ratio of ion charge to radius. Black line shows the regression ($P = 0.004$, $r^2 = 0.77$). Diagrams depict ions (red) and their hydration shells (blue), drawn to scale using the values reported in ref. 24. (B) Comparison of Na^+ and K^+ ions to CO_3^{2-} . (C) Correlation of the fraction of flies stuck in solutions from Fig. 5 (mean across concentrations) with the product of the relative size of the cation and anion hydration shells and the ratio of the charge (q) and radius (r) of the ion closest to the air-water interface. Black line shows the regression ($P = 0.024$, $r^2 = 0.77$).

at exceptionally low concentrations (0.1 mM to 0.17 M), not the ~0.5 M concentrations that are necessary, which makes Na_2CO_3 the most potent compound for ion-facilitated wetting.

The physical mechanism by which the slight negative charge at the air–water interface might increase the likelihood of wetting is not immediately clear. One possible explanation involves electrostatic attraction between the flies' surface and the negatively charged fluid layer. Recent research has shown that Cassie–Baxter-to-Wenzel-state transitions are more likely to occur in the presence of an applied voltage, which causes the formation of an electric double layer (28–30). The electric double layer increases the attraction between the interfacial water and individual roughness elements on the surface, thereby pulling the solution into the gaps and facilitating the transition to the wetted state. In experiments in which the distance between roughness elements was 4 μm (alkali flies' hairs are 3.2 μm apart), a voltage of 22 V was required to cause wetting (30). To relate these experiments with our results, we performed a rough calculation to determine the molar concentration of Na_2CO_3 needed to generate a 22-V potential between the water surface and the flies' cuticle (*SI Methods*). Our model suggests that a molarity of ~0.15 M of Na_2CO_3 is necessary to induce wetting, which is within a factor of 4 of the molarity we observed as necessary in our experiments, suggesting that this is a plausible mechanism warranting further study. A more thorough investigation would require detailed simulations of molecular dynamics that are beyond the scope of this paper.

Our theory also explains the role of the cuticular hydrocarbons in preventing wetting. The cuticle underneath the hydrocarbon layer is largely composed of chitin, which is slightly polar [static dielectric permittivity = 15 (31)]. Thus, the nonpolar hydrocarbon layer [static dielectric permittivity = ~2 (32)] helps to insulate the chitin surface from the electric double layer, reducing the likelihood of wetting. This theory is consistent with our finding that cuticular hydrocarbons do not influence wetting in pure water, as there would be no electric double layer.

Compared with the six other species we investigated, *E. hians* were the only species that resisted wetting in the presence of Na_2CO_3 , an adaptation that allows them to occupy a rare but ecologically important niche. Remarkably, the trait that allows them to forage and lay eggs in such an extreme aquatic environment arises from just a few minor changes in physical and chemical properties. These adaptations likely evolved over time in response to the slowly increasing concentration of mineral salts (such as Na_2CO_3) in alkaline lakes across the world. In recent times, the selective pressures on the alkali flies at Mono Lake have become even stronger. Between 1941 and 1982 the concentration of mineral salts in the lake doubled as a result of Los Angeles' policy of diverting water from the Eastern Sierra. Our experiments, however, suggest that this increase in ion concentration has had only a small influence on the flies' ability to dive and resurface in the lake. By comparison, the increasing salinity has had a much larger detrimental effect on the flies' larvae (33).

The most important adaptation that made the niche of underwater feeding available to the alkali fly, however, was not a physical or chemical one. Rather, it was the behavioral urge to crawl under water and forage in the first place. We suspect that their ancestors evolved this unusual behavior in lean times, when surface food was a limiting resource but underwater algae were abundant. In addition, selection against underwater foraging presumably decreased in alkaline lakes, because the caustic chemistry makes them uninhabitable for fish.

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