Nanoscale



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Cite this: Nanoscale, 2023, 15, 5726

Light-powered swarming phoretic antimony chalcogenide-based microrobots with "on-the-fly" photodegradation abilities†

Anna Jancik-Prochazkova Da and Martin Pumera +a,b,c,d

Microrobots are at the forefront of research for biomedical and environmental applications. Whereas a single microrobot exhibits quite low performance in the large-scale environment, swarms of microrobots are representing a powerful tool in biomedical and environmental applications. Here, we fabricated phoretic Sb_2S_3 -based microrobots that exhibited swarming behavior under light illumination without any addition of chemical fuel. The microrobots were prepared in an environmentally friendly way by reacting the precursors with bio-originated templates in aqueous solution in a microwave reactor. The crystalline Sb_2S_3 material provided the microrobots with interesting optical and semiconductive properties. Because of the formation of reactive oxygen species (ROS) upon light illumination, the microrobots possessed photocatalytic properties. To demonstrate the photocatalytic abilities, industrially used dyes, quinoline yellow and tartrazine were degraded using microrobots in the "on-the-fly" mode. Overall, this proof-of-concept work showed that Sb_2S_3 photoactive material is suitable for designing swarming microrobots for environmental remediation applications.

Received 7th January 2023, Accepted 26th February 2023 DOI: 10.1039/d3nr00098b

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Introduction

Microrobots are autonomous micro-scaled devices that are designed to accomplish a specific mission on microscale. ¹⁻³ Microrobots are chemically programmed to collect, transport, and deliver cargo, ⁴⁻⁶ degrade pollutants ^{7,8} and bacterial contamination, ^{9,10} or sense desired substances, ^{11,12} *etc*. However, a single microrobot exhibits relatively poor efficiency when studied in the macro-scale. Inspired by morphogenesis in biology, microrobotic systems were programmed to communicate and cooperate with each other by forming swarms of microrobots. ^{13,14} The swarming microrobots promise a powerful tool for medical and environmental applications, as they can react in response to environmental stimuli, such as change in morphology, cargo loading and transport, imaging, *etc*. ^{15–17} Swarming behavior can be induced by using various external stimuli, *i.e.* magnetic field, light, acoustic or electric

crystal structure that were navigated under polarized light.

Similarly, metal dichalcogenides, such as WS₂ and MoS₂, were

field.¹⁸ Light-induced swarm formation is quite a versatile

approach considering its low-cost instrumentation, precise

control, and biocompatibility. The mechanism of light-

induced swarming lies in different mechanisms, such as elec-

trophoresis, diffusiophoresis, and thermophoresis. 19 A typical example of light-induced swarm formation are TiO2-based swarming photophoretic microrobots. 20 Upon light irradiation, TiO₂ microrobots show schooling behavior consisting of the expansion and contraction of the swarms that act as a microfirework or micropump system due to photophoretic properties.²¹ Eventually, due to negative phototaxis, the swarms can move within the sample and transport desired substances.²² L. Wang et al.²³ used phoretic interactions among TiO₂-based microrobots to remove microplastics from aqueous samples. In addition to TiO2, light-induced swarming behavior was observed in various types of semiconductors, for example organic photoactive molecules²⁴ and polymers,²⁵ silver chloride,²⁶ silver phosphate,²⁷ iron oxide,²⁸ or chalcogenide semiconductors. 29,30 The latter is also known for good thermodynamic stability, facile synthesizability and dopability, and unique optoelectrical properties that enable their application in solar cells, thin film transistors, diodes, or water splitting photoelectrocatalysts.31 Recently, chalcogenide conductors have been applied as suitable materials for the fabrication of light-driven microrobots; X. Zhan et al. 32 reported on preparation of Sb₂Se₃-based microrobots of anisotropic

^aCenter for Advanced Functional Nanorobots, Department of Inorganic Chemistry, University of Chemistry and Technology Prague, Technicka 5, 166 28 Prague, Czech Republic. E-mail: pumera.research@gmail.com

^bFuture Energy and Innovation Laboratory, Central European Institute of Technology, Brno University of Technology, Purkynova 656/123, 621 00, Brno, Czech Republic ^cFaculty of Electrical Engineering and Computer Science, VSB – Technical University of Ostrava, 17. listopadu 2172/15, 70800 Ostrava, Czech Republic

dDepartment of Medical Research, China Medical University Hospital, China Medical University, No. 91 Hsueh-Shih Road, Taichung, Taiwan 40402
† Electronic supplementary information (ESI) available. See DOI: https://doi.org/

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applied for the fabrication of photophoretic swarming microrobots.29,30

Here, we fabricated swarming phoretic antimony(III) sulfide (Sb₂S₃)-based microrobots (further referred as "microrobots"). Sb₂S₃ is an earth-abundant semiconductor with low toxicity and good stability in air.³³ It shows remarkable optoelectrical properties suitable for solar cells³⁴ and lithium ion batteries fabrication,³⁵ photocatalytic water splitting³⁶ and photocatalytic pollutant degradation.^{37,38} Being aware of current "green chemistry" trends, 39 our objective was to apply the synthetic approach to microrobots manufacturing in an environmentally friendly way. Therefore, we employed bio-originated precursors as capping agents, that is, proteogenic α-amino acid L-cysteine and tartaric acid, which occurs naturally in many types of fruits. The synthetic route was then carried out in a microwave reactor providing great reproducibility, low-energy usage, and large-scale production. 40,41 The resulting microrobots were propelled under ultraviolet (UV) light irradiation due to the electrophoretic and diffusiophoretic behavior of the microrobots. Since the electrophoretic and diffusiophoreticinduced propulsion is caused by ions and ROS formation and redistribution, we applied the microrobots for photocatalytical degradation of organic pollutants. As model pollutants, we chose tartrazine and quinoline yellow food dyes. Given their poor biodegradability, synthetic food dyes can accumulate in wastewater and negatively influence the environment. 42-44 Anionic azo-dye tartrazine (E 102) and anionic quinophthalone dye quinoline yellow (E 104) are claimed to induce allergic and intolerance reactions, particularly in asthmatics, and cause hyperactivity, especially in children. 45-47

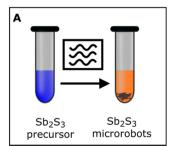
Results and discussion

Fabrication and characterization of microrobots

The Sb₂S₃ microrobots were prepared in a microwave reactor (Fig. 1) by reacting SbCl₃ (antimony(III) chloride) with biomaterial-based templates L-cysteine and tartaric acid. As suggested in a previous report, 48 Sb3+ forms a complex with tartaric acid and thus allows coupling via hydrogen bonds with L-cysteine, which acts as a sulfurizing agent to allow the formation of Sb₂S₃. To perform the synthesis in an environmentally friendly way, we proceeded the reaction in an aqueous environment. Although water is not considered the most efficient solvent to convert microwave irradiation into heat, especially in comparison to ethylene glycol, dimethyl sulfoxide, or methanol, 49 this synthetic approach showed great results and can be regarded as environmentally friendly.

To optimize the procedure, different reaction conditions were tested. Fig. S1† shows the influence of magnetic stirring, reaction time, temperature, and tartaric acid concentration on the resulting morphology of the microrobots. Briefly, stirring the reaction mixture during the synthesis enabled formation of homogeneous size and morphology distribution of microrobots. It is worth emphasizing that stirring is not typically allowed in conventional hydrothermal synthetic methods, but it can be applied and controlled in the case of using a microwave reactor. Next, the reaction time of 15 minutes caused the formation of spherical and rod-like microrobots together. After 30 minutes, only rod-shaped microrobots were obtained, suggesting homogeneous shape and size control. No further effect was observed when prolonging the reaction time to 45 minutes. The morphology was further influenced by the reaction temperature. The optimal temperature was evaluated at 180 °C as the lower temperature of 160 °C led to a reduction of microrobot size homogeneity; however, higher temperature of 200 °C caused only a slight increase in size homogeneity. Further, tartaric acid concentration was observed to have a significant effect on the resulting morphology of the microrobots as well. The optimal concentration of tartaric acid was evaluated as 8 mol. eq. with respect to SbCl3. When the concentration was reduced to 6 mol. eq. with respect to SbCl₃, sponge-like microparticles were formed, whereas increasing the concentration to 10 mol. eq. with respect to SbCl₃ caused the formation of larger rods. The protocol of the optimized microwave-assisted synthetic procedure is shown in Fig. S2.† Namely, the temperature, pressure, and power profiles are presented. It is worth noting that the precise hold of the temperature during the synthesis enables high reproducibility of the material preparation. Moreover, the power of the microwave reactor is ~20 W after reaching the set temperature suggesting low-energy consumption mode during the synthesis.

After optimizing the synthetic procedure, the microrobots resembled rods with average length of 16 \pm 5 μ m and width of



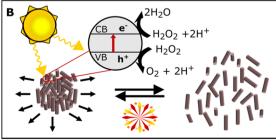


Fig. 1 Schematic illustration showing (A) the "one-pot" preparation of Sb₂S₃ microrobots using a microwave reactor and (B) their phoretic properties under light irradiation that induce swarming-like behavior. CB and VB stand for conduction and valence bands, respectively.

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 $3.0 \pm 0.9 \mu m$ (Fig. 2A). EDX images revealed a homogeneous presence of antimony and sulfur within the microrobots, suggesting that no antimony- or sulfur-rich domains were formed during the preparation (Fig. 2A). The EDX spectra are presented in Fig. S3.† The formation of the Sb₂S₃ phase was proved by recording Raman spectra. Fig. 2B demonstrates the well-resolved resulting Raman spectrum that points to the crystalline structure.50 The peaks localized at 303, 282, and 254 cm⁻¹ correspond to the Sb-S stretching vibration, and the peak at 190 cm⁻¹ corresponds to the S-Sb-S bending vibration.⁵¹ To confirm the crystalline nature of the microrobots, powder X-ray diffraction (powder XRD) analysis was performed. The resulting diffractograms (Fig. 2C) reveal a crystalline structure without signs of impurities. The diffractogram corresponded to the orthorhombic structure of stibnite with lattice parameters a = 11.33, b = 3.84, and c = 11.24 which is in agreement with reported data.⁴⁸ In the next step, we recorded absorption spectra of the powdered material to characterize its optical properties. As seen in Fig. 2D, the absorption occurs even in visible range, with the onset of absorption at around

750 nm. Assuming an indirect transition, the optical bandgap was estimated at 1.6 eV, which is in agreement with previous reports.52

It should be noted that during the typical hydrothermal preparation of Sb₂S₃ material, an amorphous phase is formed and further sintering is necessary to transform the material into a crystalline phase. 48,50 Based on the results discussed in the latter paragraph, the microwave irradiation obviously facilitated formation of the crystalline phase during the synthetic procedure. Therefore, the resulting material could be directly used for the desired application without any further energyconsuming and time-demanding thermal treatment. This observation is in agreement with previous findings in which microwave radiation assisted in the formation of high-quality crystalline structures.53

The microrobots were further characterized by measuring zeta potential in aqueous dispersions. The value of $-40 \pm$ 4 mV suggested the formation of stable colloidal solutions that do not require any further stabilization through the use of surfactants.

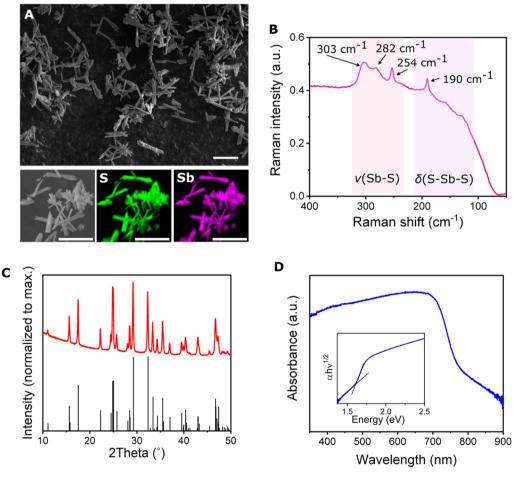


Fig. 2 Structural and optical characterization of microrobots. (A) SEM image of the microrobots and EDX elemental mapping images. Scale bar 25 μm. (B) Raman spectrum with labelled characteristic stretching (ν) and bending (δ) vibrations. (C) Powder XRD pattern of microrobots showing the characteristic fingerprint of the Sb₂S₃ material in comparison with the reference pattern (JCPDS No. 01-075-4015). (D) Absorption spectrum of microrobots with Tauc plot in the inset to determine optical bandgap.

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Light-induced locomotion of microrobots

Initially, we tested whether the photocatalytic properties of $\mathrm{Sb}_2\mathrm{S}_3$ would enable self-propulsion of microrobots under UV light irradiation. As Fig. 3A and C demonstrate, moderate self-propulsion induced by UV irradiation was indeed observed (Video S1†). The average velocity of the microrobots under UV irradiation was $0.3 \pm 0.1 \ \mu \mathrm{m \ s^{-1}}$. On the contrary, only Brownian motion was observed without any UV irradiation. To explain this observation, it should be noted that as a photoactive material, $\mathrm{Sb}_2\mathrm{S}_3$ is capable of light absorption; hence, suitable energy of irradiation can cause exciton formation where electrons (e⁻) are accumulated in the conduction band and can interact on the surface of microrobots with protons generated from the interaction of holes (h⁺) with water. The process can be summarized by the following equations:

$$2H_2O + 4h^+ \rightarrow 4H^+ + O_2$$
 (1)

$$4H^+ + 4e^- \rightarrow 2H_2 \tag{2}$$

In addition to protons, oxygen, and hydrogen, different photogenerated species, such as O_2^- , OH^+ , and OH^- , can be formed by further interactions. Due to the non-symmetrical shape of microrobots, the ion gradient around an individual

microrobot is not homogeneous, thus, self-electrophoretic motion is induced. 21,55

To investigate the origin of the phenomenon deeper, we determined the photoelectrochemical properties of microrobots by plotting Tafel curves. As the Tafel curves (Fig. 3B) suggest, there is a significant shift of about 360 mV toward positive potential after exposure of the sample to UV irradiation. This observation supports our conclusions about photochemically driven locomotion.

Self-propulsion was further enhanced by applying hydrogen peroxide as a fuel (Video S2†). The photocatalytic decomposition of hydrogen peroxide forms reactive oxygen species (ROS) by the following equations:

$$H_2O_2 + 2h^+ \to 2H^+ + O_2 \eqno(3)$$

$$H_2O_2 + 2e^- \to 2OH^-$$
 (4)

Combining the interactions described in eqn (1)–(4), the concentration of ionic species and radicals increases, therefore the self-electrophoretic phenomenon is enhanced.⁵⁵

The results of the velocity calculations are summarized in Fig. 3A. Obviously, the average velocity of the microrobots increased with increasing fuel concentration starting from the

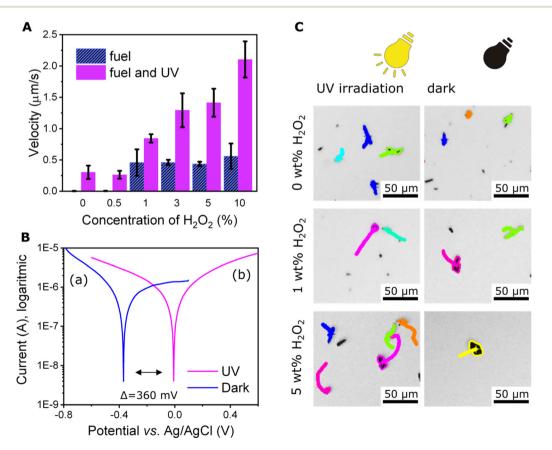


Fig. 3 Characterization of the locomotion of the microrobots. (A) Average velocity of microrobots at different fuel concentrations and the influence of UV irradiation. (B) Tafel plots of static microrobots to demonstrate photocatalytic activity. The Tafel curves were measured in pure water under dark conditions (a) and under UV irradiation (b). (C) Typical trajectories of microrobots at different conditions collected after 20 s.

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concentration of 1 wt% hydrogen peroxide. Propulsion was also observed in the presence of >1 wt% hydrogen peroxide under dark conditions. However, in this case the average velocity did not depend on the concentration of hydrogen per-

oxide and remained constant in the range $0.4-0.5 \, \mu \text{m s}^{-1}$.

Observing microrobots in their aqueous dispersion, a strong influence of UV irradiation on microrobots organization within the sample was observed. As Fig. 4 suggests, UV irradiation induces collective behavior and the swarms of microrobots expand (original micrographs without any labels are shown in Fig. S4†). Conversely, when UV light was switched off, the swarms started shrinking, suggesting a reversible process (Video S3†). Same phenomenon was previously observed in the case of titanium dioxide-based microrobots.⁵⁵ As mentioned above, Sb₂S₃ acts as a semiconductive photocatalyst and upon UV irradiation, ions and ROS are formed according to eqn (1)-(4). In addition to electrophoresis, other phenomena, such as osmotic propulsion and surface charge interaction, occur. Ions, radicals, and ROS are generated under UV irradiation, which leads to the accumulation of surface charge and electrostatic repulsions take place causing expansion of the swarms. Under dark conditions, the accumulated charged species induce electro-osmosis during their dissociation and diffusion, which obviously causes clustering of microrobots' swarms. 21,22,55 As expected, the addition of hydrogen peroxide increased the collective behavior due to more intensive ions and ROS generation. Comparison of lightinduced swarming behavior of fueled and non-fueled microrobots is also demonstrated in Video S3.† To exclude photothermal related effects, such as thermal convection that is induced when the sample is locally heated with UV light, 28 non-photocatalytic polymer microparticles were studied at the same setting while applying 5% hydrogen peroxide as a fuel and irradiating the sample with UV light. As a result, no propulsion was induced in that experiment (Video S4†) suggesting that the swarming behavior of the microrobots was not caused by temperature gradients.

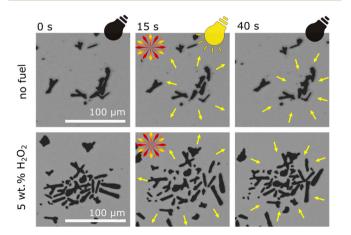


Fig. 4 Collective swarming behavior of microrobots as a function of UV irradiation. Comparison of fueled and non-fueled microrobots.

Photocatalytic degradation of food dyes

Taking into account the remarkable photocatalytic properties of Sb₂S₃, we employed microrobots for photocatalytic degradation of food dyes, i.e., quinoline yellow and tartrazine. It was assumed that the active radicals, ions, and ROS produced under UV irradiation would be able to degrade the food dves.

Quinoline yellow is an acid dye with a characteristic absorption spectrum due to the aromatic structure, with its maximum at 412 nm. On the other hand, tartrazine, as a member of azo-dye family, shows a characteristic absorption spectrum, with its maximum at 428 nm (Fig. S5†). In both cases, degradation is expected to cause a decrease in absorption bands due to the disruption of the structure. 56 Therefore, the degradation process was monitored by recording UV-Vis spectra of the samples. To perform the degradation experiments, stock solutions of food dyes were prepared and exposed to UV irradiation to first monitor stability without the presence of microrobots. After two hours of exposure to UV irradiation, the decrease in absorbance was ~7 and ~1% in the case of quinoline yellow and tartrazine, respectively, suggesting that both food dyes exhibit great stability when exposed to UV irradiation.

Prior the photocatalytic experiments, the samples containing microrobots alongside the food dyes were magnetically stirred for 20 minutes at dark conditions and the UV-Vis spectra were recorded to monitor the possible adsorption of dyes on the microrobots. However, no drop in the absorbance was detected during these equilibration experiments indicating that no adsorption occurred. In the next step, the samples were exposed to UV irradiation and the stability of food dyes was monitored using a spectrophotometer. Fig. 5 summarizes the degradation of food dyes as observed in the presence of microrobots. In the case of quinoline yellow, there was a 35% decrease in absorbance when microrobots were applied. In the case of tartrazine, the decrease in absorbance was by 17%, suggesting its better stability toward photocatalytic decomposition in this set up. The low degradation efficiency of tartrazine could be explained by different tendency toward photocatalytic degradation at different pH of solutions. In our study, no pH control was applied by using buffer solutions. The pH of the quinoline yellow and tartrazine solutions containing the microrobots were 6.5 and 6.4, respectively before the photocatalytic degradation experiments. As described in a previous study, photodegradation efficiency of tartrazine is higher at acidic conditions⁵⁷ whereas photodegradation of quinoline yellow showed to be the most efficient without any pH control.58

To investigate the degradation process of food dyes deeper, control experiments of the photodegradation of quinoline yellow were performed with ethylenediaminetetraacetic acid (EDTA). EDTA is a commonly used scavenger that interacts with photogenerated holes and hinders the formation of OH' radicals. 59 As clearly demonstrated in Fig. S6,† the presence of the scavenger significantly decreased the efficiency of photoNanoscale Paper

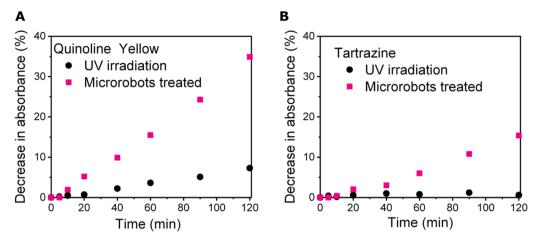


Fig. 5 Photodegradation of food dyes determined by the change in absorbance. Graphs show degradation of quinoline yellow (A) and tartrazine (B) using photocatalytically active microrobots and UV irradiation alongside the control experiment where no microrobots were applied.

catalytic degradation of quinoline yellow; the absorbance decreased by 22% after two hours This observation supports the assumption that the ROS, especially OH' radicals are generated upon exposure of the microrobots under UV illumination providing the degradation abilities.

Conclusions

We employed a microwave-assisted technique to develop a reproducible, low-cost, and "green" manufacturing process of microrobots. It is worth noting that the original synthesis of the Sb₂S₃ material was performed in an autoclave at 180 °C for 12 hours and further annealing at 250 °C for 3 hours was necessary to obtain the crystalline structure. 48 Here, we performed the reaction in a microwave reactor at 180 °C for 30 min without any further thermal treatment. Sb₂S₃ was presented as photoactive material that enables the self-propulsion of the resulting microrobots under UV irradiation. Self-propulsion was further enhanced in the presence of hydrogen peroxide as a fuel. Due to their photocatalytic properties, microrobots were applied for the degradation of food dyes, namely, tartrazine and quinoline yellow. It can be expected that such microrobots can find application in the removal of residual food dyes and beyond.

Experimental

Materials

Antimony(III) chloride (SbCl₃, ≥99.5%), L-tartaric (≥99.5%), L-cysteine (97%), tartrazine (analytical standard), quinoline yellow (mixture of the mono- and disulfonic acids), ethylenediaminetetraacetic acid (EDTA, ≥98.5%), and hydrogen peroxide (30%) were obtained from Sigma Aldrich. Ethanol absolute A.G. was purchased from Penta. All chemicals were used as received.

Synthesis of Sb₂S₃ microrobots

For a typical microwave-assisted synthesis of Sb₂S₃ microrobots, SbCl₃ (0.46 mmol) was dissolved in 20 mL of tartaric acid stock solution (0.17 mol dm⁻³) using a sonication bath. In the next step, 1-cysteine (0.65 mmol) was added and the solution was magnetically stirred for 10 minutes. The precursor solution was transferred to a reaction vial and placed in the microwave reactor (Monowave 400, Anton Paar). The reaction was carried out at 180 °C for 30 minutes while being magnetically stirred (600 rpm). The crude product was collected by centrifugation (4000 rpm, 3 min) and washed with ultrapure water twice and with ethanol three times. The resulting microrobots were vacuum-dried at 40 °C overnight.

Characterization

Morphology was characterized using a MIRA 3 XMU (Tescan) scanning electron microscope equipped with an EDX detector (Oxford Instruments) for elemental mapping. Powder samples were characterized on an adhesive carbon tape.

Powder XRD characterization was performed using a Rigaku SmartLab 3 kW X-ray powder diffractometer with Cu Kα radiation source. Raman spectra were measured with an IR-Raman spectrometer (Bruker RFS 100) using a 9394.4 cm⁻¹ laser for excitation. The absorption spectra of microrobots in solid state were recorded by a Jasco V-750 UV-Vis spectrometer equipped with an integrating sphere (ISV-922). The zeta potential was measured by Zetasizer Ultra (Malvern Panalytical Ltd) and an average of 5 measurements was calculated.

Tafel plots were obtained by performing linear sweep voltammetry on a PGSTAT 204 (Metrohm Autolab) equipped with UV LED (365 nm, 700 mA, radiant flux 3.3 W, LedEngin Inc.). For this purpose, microrobots' aqueous dispersion was dropcasted over an ITO substrate that was then dried at ambient conditions. Ag/AgCl and platinum electrodes were employed as reference and counter electrodes, respectively. The measurements were made in distilled water under dark conditions and under UV irradiation.

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Locomotion analysis was performed using a Nicon Eclipse Ts2R inverted microscope equipped with a Basler ace acA1920-155uc digital camera. The microscope was also equipped with a UV-LED source (pE-100, 365 nm, 1670 W cm⁻²). The sample for the locomotion analysis was prepared by placing a droplet of microrobots' solution (5 µl) on a glass slide. The concentration was adjusted according to the setting of microscope to obtain videos of at least 60 tracks. Alternatively, hydrogen peroxide was added to the microrobots in a specific amount to reach its desired concentration to act as fuel. The tracks of the microrobots were analyzed using the TrackMate plugin within ImageJ software. 60 A Python code subtracting eventual drift was used to calculate average velocities of each video. 61 The resulting velocities were obtained as an average and a standard deviation of at least three videos. Finally, the representative trajectories were visualized using NIS software.

Food dyes degradation experiments

The sample containing microrobots (0.2 mg mL $^{-1}$) and food dye (quinoline yellow or tartrazine, both 0.01 mg mL $^{-1}$) in total volume 20 mL was magnetically stirred for 20 min in a 50 mL reaction vial at dark conditions before the experiment. The sample was exposed to UV irradiation (365 nm, 700 mA, radiant flux 4.1 W, LedEngin Inc.) and aliquots were taken at given time intervals to monitor the degradation process. For this purpose, the sample was centrifuged (4000 rpm, 3 min) to remove microrobots and the absorption spectrum of the supernatant was recorded in quartz cuvettes (1 × 1 cm) using the UV-Vis spectrometer. The absorbance of quinoline yellow and tartrazine was read at 412 and 428 nm, respectively. The control experiment with EDTA was performed at same conditions with the modification that EDTA was added to the reaction mixture (0.1 mM) before the photocatalytic reaction.

Author contributions

A.J.P. performed the experiments, treated the data, and wrote the manuscript. M.P. supervised the project and revised the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the project Advanced Functional Nanorobots (reg. no. CZ.02.1.01/0.0/0.0/15_003/0000444 financed by the EFRR). CzechNanoLab project LM2018110 funded by MEYS CR is gratefully acknowledged for the financial support of the measurements at CEITEC Nano Research Infrastructure.

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