

Wien effect and photo-accelerated interfacial water dissociation across proton permeable graphene electrodes

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The phenomenon known as the Wien effect was previously observed using high-voltage electrolysis cells that produced fields of about 10^7 V m^{-1} . The observation of the Wien effect for the common case of water dissociation has remained elusive. Here we study the dissociation of interfacial water adjacent to proton-permeable graphene electrodes and observe strong acceleration of the reaction in fields reaching above 10^8 V m^{-1} . The observed exponential increase in proton currents is in quantitative agreement with Onsager's theory. The use of graphene as a membrane allows for measuring the proton currents arising exclusively from the dissociation of interfacial water. Illumination of the graphene under visible light produced an order-of-magnitude acceleration of the interfacial water dissociation reaction. The found photo effect is attributed to the combination of graphene's perfect selectivity with respect to protons, which prevents proton-hydroxide recombination, and to proton transport acceleration by the Wien effect. Our findings provide fundamental insights into ion dynamics near atomically thin proton-selective interfaces.

References

[1] J. Cai, E. Griffin. et al, Nat Commun, 13 (2022) 5776

[2] J. Cai, E. Griffin. et al, Nano Lett, 22 (2022) 9566-9570

Figures

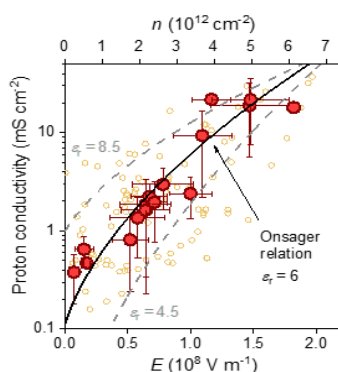


Figure 1: Proton conductivity through graphene electrode where n is the carrier density. Bottom x-axis, is Electric field strength on graphene electrode. Solid curve, best fit of Onsager model to data. Dotted curves, Onsager model for different dielectric constant.

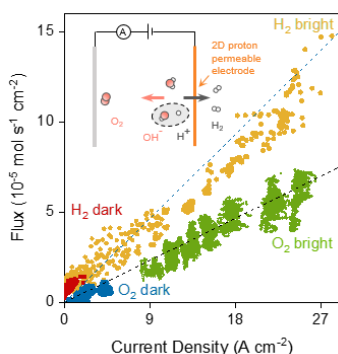


Figure 2: Faradaic efficiency measurements. Hydrogen and oxygen fluxes as a function of I under dark and bright conditions.