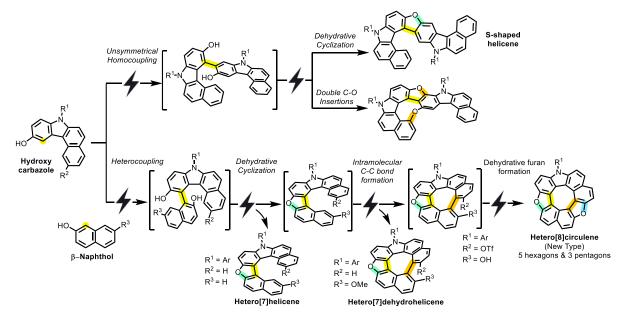
Electrochemical Cascade Synthesis of Polycyclic Heteroaromatics: A Sustainable Gateway to Functional Optoelectronic and Photocatalytic Materials

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Polycyclic heteroaromatic compounds such as helicenes, dehydrohelicenes, and circulenes are of growing interest due to their distinctive optoelectronic, redox, and chiroptical properties. However, their synthesis remains a synthetic challenge, often requiring harsh conditions and exhibiting poor functional group compatibility. In this lecture, I will introduce the first electrochemical cascade approach to access a diverse library of PHAs in a sustainable and modular fashion. This electrochemical platform leverages differential oxidation potentials to achieve precise control over sequential bondforming events, and was further streamlined into a one-pot protocol using readily available starting materials. Comprehensive structural, spectroscopic, and computational investigations reveal distinct electronic architectures and redox profiles. Notably, the synthesized dioxaza[8]circulenes function as efficient metalfree photocatalysts for C–X (X = C, B, S, P) bondforming reactions under visible-light irradiation, achieving excellent yields without the need for transition metals. Furthermore, the helicene and dehydrohelicene derivatives exhibit remarkable configurational stability and pronounced chiroptical activity, including a record-high blueregion circularly polarized luminescence (CPL) signal with a dissymmetry factor $|g_{lum}|$ of 2.5 x 10⁻³ at 433 nm.



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