Mixed oxygenase-desaturase activity in carboxylic acid-directed non-heme iron catalyzed C-H oxidations

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We report that the small molecule non-heme iron catalyst Fe(PDP) is able to exploit carboxylate ligation from substrates to control reactivity, site-selectivity and even influence reaction pathways in aliphatic C—H oxidations. Moreover, carboxylic acid substrates divert the hydroxylation reactivity of Fe(PDP) toward dehydrogenations via a short-lived carbon-centered radical. Such substrate-dependent dual activity for aliphatic C—H bonds has previously only been observed in the realm of enzymatic catalysis.

