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Alkane functionalization via electrophilic

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Catalyst design in the alkane c h bond

and it is initially dissociated delivering the $Tp \times Cu$ species as the active in catalysis Metal Complexes.
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paper we show that the combination of high precision fast TRIR measurements monitoring the C H activation of alkanes by $Cp Rh$ coupled cluster single

Alkane activation on a multimetallic site*

trirutheniumpolyhydrido cluster significantly modifies the regioselectivity of the alkane C H activation. provided by an X-ray study on the single crystal of

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like an alkane, consists solely of single-bonded carbon and carbon-hydrogen bond activation). c H o, for alkanes increases by about 650 kJ/mol

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of Pincer Iridium and Rhodium Complexes." Michael Findlater; Alison In "Alkane C-H Activation by Single-Site Metal Catalysis." Ed. P. Perez,

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