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Designing metal hydride complexes for water splitting reactions: A molecular electrostatic potential approach

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Abstract:

The hydridic character of octahedral metal hydride complexes of group VI, VII and VIII has been systematically studied using molecular electrostatic potential (MESP) topography. The absolute minimum of MESP at the hydride ligand \((V_{\text{min}})\) and MESP value at the hydride nucleus \((V_{\text{H}})\) are found to be very good measures of the hydridic character of the hydride ligand. The increasing/decreasing electron donating feature of the ligand environment is clearly reflected on the increasing/decreasing negative character of \(V_{\text{min}}\) and \(V_{\text{H}}\). The formation of an outer sphere metal hydride-water complex showing H...H dihydrogen interaction is supported by the location and value of \(V_{\text{min}}\) near the hydride ligand. A higher negative MESP suggested lower activation energy for H\(_2\) elimination. Thus, MESP features provided a way to fine tune the ligand environment of a metal-hydride complex to achieve high hydridicity for the hydride ligand. The applicability of MESP based hydridic descriptor in designing water splitting reactions is tested for group VI metal hydride model complexes of tungsten.

Introduction

Transition metal hydride complexes play an important catalytic role in a variety of synthetically important reactions as well as in biological reactions.\(^1\)\(^-\)\(^6\) Several transition metal hydride complexes catalyse dihydrogen formation in the presence of reducing agents.\(^7\)\(^-\)\(^13\) Recently, Milstein \textit{et al.} reported the discovery of a monomeric aromatic pincer Ru(II) hydride catalyst for water splitting reaction.\(^14\) This catalyst activates the O-H bond for water splitting through aromatization-dearomatization steps involving a PNN pincer ligand and the metal center.\(^15\)\(^,\)\(^16\) Mechanistic studies from our group have shown that the hydride ligand of the Milstein catalyst facilitates H...H dihydrogen interaction with water molecules leading to an outer sphere mechanism for H\(_2\) elimination.\(^17\) Similarly, hydride donor ability has been invoked to explain the reaction of \(W(\text{H})\)(NO)(CO)(PMe\(_3\))\(_2\) with various alcohols, under mild condition to liberate dihydrogen.\(^18\) Very recently we have shown that H...H dihydrogen bonded complexes are reactant species in the associative and migratory insertion mechanisms of MH(NO)(CO)(PH\(_3\))\(_2\), M = Cr, Mo and W complexes for H\(_2\) elimination.\(^19\) Peruzzini \textit{et al} studied the dihydrogen interaction between the metal hydride complexes \([\text{CpW(H)(CO)}_2][\text{BuPCP}=2,6-\text{C}_6\text{H}_3(\text{CH}_3\text{PPh}_2)_2]\) and reported the evolution of H\(_2\) upon the increase in reaction temperature.\(^20\) The H\(_2\) elimination is also reported for four-, five- and seven-coordinated transition metal hydride complexes.\(^21\)\(^-\)\(^23\) In the case of four-coordinated complex of \(\text{CpRu(dppe)H}\) (dppe = Ph\(_2\)PCH\(_2\)CH\(_2\)PH\(_2\)) with HA (A = acceptor), an equilibrium between dihydrogen bonded complex and cationic dihydrogen complex has been monitored at low temperature by NMR spectroscopy.\(^24\) It is noteworthy that such systems are precursor for (\(\eta^2\)-H\(_2\)) Kuba’s dihydrogen complexes.\(^25\) Moreover, these dihydrogen complexes are potential candidates for hydrogen storage applications, crystal engineering, catalyst designing and many other applications.\(^26\)\(^,\)\(^27\)

The present study mainly focuses on the analysis of molecular electrostatic potential (MESP) and its topography in metal hydride complexes. MESP features have been successfully employed to understand many important fundamental problems in chemistry.\(^28\)\(^-\)\(^31\) For example, MESP is used for the study of the site for electrophilic or nucleophilic attack, intermolecular reactivity, non covalent interaction, electronegativity of elements, substituent effects and various chemical phenomena.\(^32\)\(^-\)\(^37\) Gadre \textit{et al.} derived MESP topographical features for probing the intermolecular distance in hydrogen bonded complexes of the type D-H...A (D = N, O, B, C, F, S, Cl; A = F).\(^38\) Galabov \textit{et al.} used MESP at hydride nucleus for characterizing the reactivity of substituted aromatic molecules.\(^39\)\(^-\)\(^41\) Previously proton affinity, thermodynamic and various spectroscopic methods have been used to characterize electron donating character of a hydride ligand of metal hydride complexes.\(^42\)\(^-\)\(^45\) In the present study a
MESP based approach will be presented to characterize the hydridic character of a hydride ligand in a metal complex and the formation of electron rich hydride ligands in some metal hydride complexes will be revealed for the facile liberation of dihydrogen from water.

**Theoretical methods**

The MESP, \( V(r) \) at a point \( r \) in atomic units is given by eq. 1 where \( \rho(r) \) is the electronic charge density and \( N \) is the total number of nuclei in the molecule.\(^{29}\) \( V(r) \) is a real physical quantity

\[
V(r) = \sum_{\lambda} \frac{Z_{\lambda}}{|r - R_{\lambda}|} - \int \frac{\rho(r')d^3r'}{|r - r'|}
\]

and can either be positive or negative depending upon whether the effect of nuclei (first term in eq. 1) or electron (second term in eq. 1) dominates over the other. The MESP at the nucleus A of a molecule \( V(A) \) can be obtained by dropping out the nuclear contribution due to \( Z_{A} \) from eq. 1.\(^{29}\) In MESP topography analysis, negative-valued minima appear at the electron rich regions of molecules. In this work, a MESP minimum located near a hydride ligand of the metal hydride complexes is denoted as \( V_{\text{min}} \) and the notation \( V_H \) is used to represent the MESP value at the hydride nucleus.

**Gaussian09** suite of quantum chemical programs is used for performing all the calculations.\(^{46}\) Geometry optimization of all the complexes are performed using the hybrid density functional B3LYP.\(^{47}\) The LanL2DZ basis set is chosen for transition metals which also incorporates the effective core potential (ECP) of Hay and Wadt with additional f polarization functions.\(^{49,50}\) For all other atoms, 6-31G(d,p) basis set is selected.\(^{51,52}\) Further, a triple zeta split-valence basis set 6-311++G(d,p) was used for atoms other than transition metals in single point calculation to obtain more accurate description of the activation barrier for H2 elimination step. All the optimized structures were confirmed as either minima or transition state by vibrational frequency calculation. All the reported transition states show one imaginary frequency and all other structures have no imaginary frequency. IRC calculations are conducted for confirming the connectivity of the transition states to the respective minimum energy structures.\(^{53}\) In the case of water adducts, counter poise method was used to evaluate the BSSE correction for the interaction energy.\(^{54}\) Effect of phosphine coordination is mainly modeled using PH3 ligand while several test cases have been modelled using ligands such as trimethyl-, triethyl-, and triphenyl phosphines.

**Results and Discussion**

The octahedral metal hydride complexes of Mo and W from group VI, Mn and Re from group VII and Fe and Ru from group VIII given in Fig. 1 are selected for this study. At least one phosphine ligand is selected for most of the complexes because they have high synthetic applicability in tuning steric and electronic properties of the complexes.\(^{55-58}\) In the case of Mn complexes, various possible combinations of CO and PH3 ligands are considered along with chelate 1, 2-bis(phosphino)ethane ligands and N2 ligand. Remaining complexes, consists of various types of ligands, viz. CN, Cl, F, NO, CH, N2, NCO, CO, SiH4, P- and N-based bidentate, cyclic NC3H3 and N3C3H3 ligands. The 1 and 11 contain no phosphine ligand. Most of the synthetically known complexes consist of triphenyl/trimethyl/isopropyl phosphine ligands while our calculation replaces them with unsubstituted phosphine which reduces the computational cost. For a limited number of tungsten complexes, higher derivatives have been computed using bulky triethyl-, trimethyl- and triphenyl phosphine ligands. Compounds similar to 1, 2a, 3a, 3b and 4a have been reported in the literature.\(^{19}\) The 7, 8 and 9 are made up of bidentate chelating phosphine ligand.

Among the complexes reported in Fig. 1, for the higher derivatives (with bulky phosphine ligands), the literature shows the following aspects on their properties/reactivity. Through NMR measurements, they calculated that the effective charge of 23 hydride in MnH(CO)3PH3 (the higher derivative of 2a) is -0.40 e, which was higher than 1. Darenbourg et al. reported the exopeptide ring opening by H ligand in the chelated Mn complex MnH(dppp)(CO)2 where dppp is 1,3-bis(diphenylphosphino)-propane)\(^{42}\) Nietlispach et al. synthesized a series of Re-hydride complexes ReH(CO)n(PR3)6−n (n=5, 4, 3, 2, 1 and R=Ph, CH3) and showed that hydridic character depends on the increasing number of phosphine ligand and they undergo insertion reactions with CO, CO2, CS2, and ketones.\(^{50}\) Berke et al. studied the unusual insertion behaviour in the metal-H bond of chelated MoH(dmpc)NO complex (dmpc = 1,2-bis(dimethyl phosphino)-) with inactive imines and argued that the high propensity of the dmpe ligand towards imine was due to the chelating effect of cis-bidentate phosphine ligand which enhanced the hydridic character compared to MoH(NO)(CO)3(PR3)2 and MoH(NO)(CO)(PR3)3. Abramov et al. noted that by substituting PPh3 in place of CO group in system 1 gives higher hydridic character to the hydride ligand.\(^{59}\) The MoH(CO)3(PR3)6−n (n=3, 2, 0) and WH(CO)3(PR3)6−n (n=3, 2, 1) complexes have shown hydride transfer activity to CO, CO2, CS2, acetylene and ketones.\(^{44,60,61}\) Hillhouse and Haymore synthesised WH(NO)(CO)2(PR3)2 (R = C6H5 and CH3C6H4) and showed that these complexes undergo various type of insertion and elimination reactions with small molecules.\(^{62}\) Recently we have described the insertion behaviour of water with bis- and, tris-phosphine Cr, Mo and W hydride complexes for H2 elimination.\(^{19}\) The enhancement in the hydridic character of a metal hydride complex is strongly related to the weakening of the metal-H bond and suggests that a metal-H bond is susceptible for various reactions such as insertion, elimination, proton donation and proton transfer.
structural features of Mo, W, Mn, Re, Fe and Ru hydride complexes

All the complexes acquired pseudo octahedral geometry (some representative structures are listed in Fig. 2), in which most of them possess slight inclination of equatorial ligands to the hydride ligand (the average ligand-metal-H angle, $\delta$ is 85$^\circ$). The metal-H (M-H) distance is found to be in the range 1.506 to 1.897 Å which can be correlated to the position of the metal in the periodic table and the associated ligands in the complex (Table 1). A general trend of increase in M-H distance with increase in the size of the metal is observed in each group. The M-H distance increases to the maximum for Mo and W complexes and suggests easy M-H bond dissociation for group VI complexes. Minor discrepancies in the general trend of M-H distance can be observed in cases of (Mo, W) complexes such as (31, 37), (33, 38) and (35, 40). This can be attributed to variations in the ligand

Fig. 1 Schematic representations of group VI, VII and VIII octahedral metal hydride complexes.

Fig. 2 Structures of a representative set of metal-hydride complexes. Bond distance in Å and bond angles in degree.
NO oxygen. The MESP feature showing two ligand. The MESP isosurface depicted in Fig. 3b is useful to around the hydride ligand as well as around the oxygen of the NO ligands (eg. 40 in Fig. 2).

33 It is noteworthy that \( V_{\text{min}} \) of the hydride ligand (-25.6 kcal/mol) is more negative than \( V_{\text{min}} \) of NO ligand (-21.7 kcal/mol). This suggests a highly localized electron distribution around the hydride ligand where the electron concentration is higher than the NO oxygen. The MESP feature showing two \( V_{\text{min}} \) points for the hydride ligand indicates a symmetrical distribution of the electron density with respect to the P-Mo-P plane. Fig. 3c is the typical MESP painted van der Waals surface of the complex wherein the violet region around the hydride ligand appears as the most electron rich. For all the metal hydride complexes, negative-valued \( V_{\text{min}} \) is always observed for the hydride ligand. In the following section, \( V_{\text{min}} \) features will be discussed for various systems.

### MESP analysis of Mn Mo, W, Re, Fe and Ru hydride complexes

The MESP is very sensitive to electronic perturbations in a molecule and hence variations in the ligand environment can cause subtle changes in the MESP value around the hydride ligand. This can be easily proved by analyzing the \( V_{\text{min}} \) of all the metal hydride complexes (Table 2). For instance, in the case of Mn complexes, 1 with five CO groups shows the lowest \( V_{\text{min}} \) -2.7 kcal/mol while complex 6 having only PH3 ligands shows the most negative \( V_{\text{min}} \) -34.1 kcal/mol. The drastic change in \( V_{\text{min}} \) can be associated to the \( \sigma \) donating power of PH3 ligand compared to the electron withdrawing character of CO. The \( V_{\text{min}} \) of 1, 2, 3, and 4 are -2.7, -12.6, -22.7 and -32.9 kcal/mol, respectively which indicates that for substituting every CO by a PH3, \( V_{\text{min}} \) becomes deeper by ~10 kcal/mol. Similar conclusion is valid for isomers of 1, 2, 3, and 4. For instance, \( V_{\text{min}} \) of 2a is -12.5 kcal/mol while that of 3b is -22.7 kcal/mol. Other examples for comparison are (3a, 4a) and (4b, 5b). In the case of the three isomers of tris-phosphone systems 4a, 4b, and 4c, significantly more negative \( V_{\text{min}} \) is observed for 4c which suggests that equatorial/axial orientation of the ligands also play a role in controlling the hydridic character. As we move from 3c to 7, enhancement of the negative character of \( V_{\text{min}} \) is observed which can be attributed to the chelating effect of the bidentate phosphine ligand. The 9 and 8 complexes are very similar except for the axial ligand, the former having N2 has \( V_{\text{min}} \) ~30.7 kcal/mol while the later having CO has \( V_{\text{min}} \) ~29.9 kcal/mol. These results suggest that CO and N2 have similar electronic effect on the hydridic character.

The \( V_{\text{min}} \) values clearly suggest that negative MESP around the hydride ligand undergoes remarkable changes with respect to the nature of the metal and its ligand environment. For instance, Re complexes show more negative \( V_{\text{min}} \) than Mn complexes. In general, going down the group enhances the negative character of \( V_{\text{min}} \). Among the seven Re complexes, 17 has the maximum negative \( V_{\text{min}} \). The \( V_{\text{min}} \) of 17 is more negative than \( V_{\text{min}} \) of its isomer 15 by 5.1 kcal/mol suggesting that the influence PH3

### Table 1. Metal hydrogen distance in Å for all the complexes.

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Fig. 3 Electrostatic potential features of 31. (a) MESP plotted on a plane. (b) MESP isosurface plot, value -2.2 kcal/mol (c) MESP on van der Waals (value range from -0.02 au (violet) to 0.02 au (red)).

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ligand trans to the hydride ligand is more electron donating than the trans orienting N₂. The 11 exhibits the lowest \( V_{\text{min}} \) (-9.2 kcal/mol) which can be attributed to the electron withdrawing effect of CO ligands. Group VII complexes show a wide range of \( V_{\text{min}} \) values due to the inclusion of various electron withdrawing/donating ligands.

On the basis of \( V_{\text{min}} \), the ligand environment of metal hydride complexes can be understood either as electron withdrawing or as electron donating. The \( V_{\text{min}} \) of a representative set of Mn and Fe complexes are depicted in Fig. 4 along with a MESP isosurface of value -19.0 kcal/mol. The phosphine, CH, NO, bidentate chelating phosphate ligands and amino ligands show large negative values for \( V_{\text{min}} \) compared to other ligands. Such ligands could be treated as electron donating whereas the metal hydride complexes containing CO, Cl, F and SiH groups always show smaller negative \( V_{\text{min}} \) values. These ligands act as electron withdrawing and hence a decrease in the hydridic character is expected in the corresponding metal hydride complexes. In most cases, the donating cis ligands are more useful than trans withdrawing ligands for improving hydridic character. For instance, 20, 21 and 22 complexes have moderate \( V_{\text{min}} \) values with electron withdrawing group in the trans position.

In the case of tungsten complexes containing bulky triethyl-, trimethyl- and triphenyl phosphate ligands, the strong \( \sigma \)-donating alkyl substitution on phosphorous is found to enhance \( V_{\text{min}} \) of the hydride ligand than the \( V_{\text{min}} \) of PH₃ ligand (s.i.). Such bulky ligands improve the hydridic character. Since MESP is a one electron property, it is well established that the absolute values as well as trends in \( V_{\text{min}} \) of molecules are largely unaffected by the level of theory and basis set used. In order to verify this, \( V_{\text{min}} \) of a representative set of Ru hydride complexes is evaluated using TPSS/aug-cc-pVDZ DFT method. At this level, the magnitude of \( V_{\text{min}} \) only slightly decreased and the overall trend of \( V_{\text{min}} \) remained unaffected compared to B3LYP results (s.i).

The MESP at the hydride nucleus (\( V_{\text{H}} \)) is also very sensitive to the ligand environment as well as the metal center as it is varied from -685.2 to -759.1 kcal/mol (Table 2). The most negative and the least negative \( V_{\text{H}} \) are observed for 42 and 1, respectively. In the case of Mn complexes, negative character of \( V_{\text{H}} \) is increased with increase in the number of electron donating phosphate ligands (1 – 6). Very similar observation is also found in Re complexes (11 – 17).

Electron donation of the phosphine in the equatorial position (cis to hydride ligand) is more effective than that in the axial position (trans to hydride ligand). An electron withdrawing ligand trans to the hydride ligand increases the magnitude of \( V_{\text{H}} \). For example, in Fe complexes 18 and 19 where trans ligand is Cl and F, respectively, \( V_{\text{H}} \) of 18 is -703.3 kcal/mol and that of 19 is -710.0 kcal/mol. This indicates that Cl has more electron withdrawing power than F. Similar effect is observed for Ru complex 25 which has Cl in the equatorial position and shows \( V_{\text{H}} = -710.3 \) kcal/mol while the corresponding F derivative (26) shows \( V_{\text{H}} = -716.5 \) kcal/mol. The \( V_{\text{min}} \) and \( V_{\text{H}} \) values show a good linear correlation (Fig. 5). Previously we have shown that the spatial quantity \( V_{\text{min}} \) is more sensitive to steric effect than the nuclear quantity \( V_{\text{H}} \) and hence the outliers seen in the linear trend can be attributed to the steric effect of the ligand environment on \( V_{\text{min}} \).

**Stabilization energy of dihydrogen bonded hydride complexes**

Recently, we have shown that the electron rich hydride ligand participates in dihydrogen H...H interactions with solvent water molecules and such interactions play significant role in water splitting reactions catalyzed by Ru pincer complex.21 Following this work, the H...H interaction between a water molecule and the hydride ligand in all the complexes is analyzed. An increase in the hydridic character is expected to increase the H...H interaction energy. Typical H...H distance (d_HH) in dihydrogen bonded complexes is in the range 1.6 to 2.2 Å66–68 and all the d_HH values are found in the range 1.621 to 2.140 Å (Table 2). In general, the d_HH decreases with increase in the negative character of \( V_{\text{min}} \) or \( V_{\text{H}} \) meaning that metal complexes showing strong hydridicity.

**Fig. 4** Isosurface plots of MESP (-19.0 kcal/mol) for isomeric hydride complexes of Mn and Fe. MESP minimum at the hydride ligand (\( V_{\text{min}} \)) in kcal/mol is also shown.

**Fig. 5** The correlation between MESP at the hydride nucleus (\( V_{\text{H}} \)) and MESP minimum at the hydride ligand (\( V_{\text{min}} \)) for all metal hydride complexes.
enhance outersphere binding of water to the hydride complex (s.i.). Equatorial groups such as Cl, SiH₃, CNO, heterocyclic, and CO (in cis- or trans- position), increase d_HH. Ligands trans with respect to hydride, viz. NO and CH as well as bidentate P and N ligands decrease d_HH. The orientation of water in the dihydrogen complex can be measured in terms of the M-H…H angle Θ and H…H-O angle Ψ. The Θ values are in the range 120-145° while Ψ values vary from 109 to 178° (Table 2). The Ψ is found to be more linear with more negative V_H (correlation coefficient = 0.920).

Experimental studies reveal that the stabilizing energies of H…H bonded complexes (E_HH) is in the range 2 - 7 kcal/mol. In this study, E_HH is found in the range 2.5 to 13.2 kcal/mol and several complexes have E_HH values higher than the typically observed experimental values. The highest E_HH value is observed for 29_H₂O (10.6 kcal/mol). In general, E_HH increases with increase in the negative character of V_H (Fig. 6) which means that more hydridic character of the ligand promotes H…H dihydrogen interaction. This result is important because it suggests that the electron rich character of the hydride ligand assessed in terms of V_H is useful to make a good prediction on the power of the hydride ligand to interact with a water molecule through dihydrogen bond. The prediction is only qualitative because E_HH does not fully account for the dihydrogen bond energy. In many cases, apart from dihydrogen interactions, secondary interactions involving oxygen of water contributes to E_HH. Representative two Ru hydride complexes showing this feature is given in Fig. 7.

Secondary interactions arise between oxygen of water and hydride of the equatorial chelating ligands or between the hydride of water and oxygen of the equatorial CO ligands. Other examples showing secondary interactions are 7_1H₂O, 8_1H₂O, 9_1H₂O, 21_1H₂O, 33_1H₂O, 35_1H₂O, 41_1H₂O and 42_1H₂O. In 1_1H₂O, 11_1H₂O and 2b_1H₂O dihydrogen bond formation is not observed as the water molecule preferred an interaction with the carbonyl than the hydride.

### Table 2. Structural parameters and binding energies (kcal/mol) of all dihydrogen bonded complexes. Dihydrogen distance, d_HH in Å and angles (Ψ(M-H-O) and Θ(M-H-H)) in degree. The V_H and V_O of the parent complexes (kcal/mol) are also given.

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### Fig. 6 Correlation between MESP at the hydride nucleus (V_H) and stabilization energies of H…H bonded complexes (E_HH).

Designing metal hydride complexes for water splitting reactions

Experimentally it is known that tris- and bis-(trimethyl phosphine) W hydride complexes and bidentate Mo, W and Fe hydride complexes are useful for liberating hydrogen from water. It is also established that increasing hydridic character favours protonation from organic molecules leading to dihydrogen complex formation. Strong hydridic character of a metal hydride can promote hydrogen release through a water splitting outer sphere mechanism. From the knowledge of V_min or V_H (Table 2) we may assess that 5b and 6 from group VII and 22 from group 8 and 35, 36, 41 and 42 from group VI have high hydridic character and hence they may show higher tendency for water splitting reactions.

The applicability of V_min or V_H for tuning the activation barrier of H₂ elimination is tested by transition state modeling for group VI complexes. Large negative MESP values and weak M-H bond
strength are main reason for selecting group VI complexes. A representative example of 42 giving the details of the outer sphere mechanism is presented in Fig. 8. The dihydrogen complex passes through a 4-center transition state to yield the hydroxide metal complex. This type of a mechanism was previously discussed for bis-, tris-, and tetrakis- phosphine complexes of W, Mo and Cr.

Fig. 7 Water adducts of Ru hydride complexes. Distance in Å.

Fig. 8 Relative free energy profile of a representative system for hydrogen elimination.

Fig. 9 Correlation between activation barrier (ΔE) and MESP minimum at the hydride ligand (V_{min}) of W hydride complexes.

Conclusions

In summary, the structural and electronic features of a large variety of metal hydride complexes wherein the metal is Mo, W, Mn, Re, Fe and Ru have been studied using DFT. In all complexes, MESP topography revealed that the hydride ligand possesses a characteristic negative minimum V_{min}. The value of V_{min} is useful to estimate the hydridic character (electron rich character) of that ligand. The hydridic nature of the ligand is also measured in terms of MESP at the hydride nucleus (V_H). The V_{min} and V_H are very sensitive to the ligand environment. Hence, by monitoring the MESP that surrounds the hydride ligand, fine tuning of the ligand environment is possible. As V_{min} of the hydride ligand becomes more negative, its interaction with a water molecule for dihydrogen bonding becomes higher leading to facile release of H_2 from the complex. A more electron rich hydride ligand showed lower activation barrier for H_2 elimination. Thus V_{min} and V_H are proposed as two useful electronic descriptors for the study of the hydridic character of a metal hydride complex. Since MESP analysis provides more insight into the interactive behaviour of the metal hydride complex with a water molecule, tuning the ligand environment for more negative V_{min} or V_H is attractive to design new water splitting metal hydride complexes.

Acknowledgements

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Reference

Designing metal hydride complexes for water splitting reactions: An molecular electrostatic potential approach

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Graphical Abstract

Molecular electrostatic potential minimum of metal-hydrides ($V_{\text{min}}$) is used as a sensitive electronic descriptor to tune $\text{H}_2$ elimination reactions.