Electrocatalytic hydrogen generation using tripod containing pyrazolylborate-based copper(II), nickel(II), and iron(III) complexes loaded on a glassy carbon electrode†

Mohamed M. Ibrahim,†a G. A. M. Mersal,†a Ahmed M. Fallatah,†a Khaled Althubeiti,†c Rabah Boukherroub,g Mohamed S. Attia,†g,‡c Hamdy S. El-Sheshtawy,†d Manal F. Abou Taleb,†d,e Manash R. Das,†f Rabah Boukherroub,†g Mohamed S. Attia,†h and Mohammed A. Amin,a

Three transition metal complexes (MC) namely, [TpMeMeCuCl(H2O)] (CuC), [TpMeMeNiCl] (NiC), and [TpMeMeFeCl2(H2O)] (FeC) (TpMeMe = tris(3,5-dimethylpyrazolyl)borate) were synthesized and structurally characterized. The three complexes CuC, NiC, and FeC-modified glassy carbon (GC) were examined as molecular electrocatalysts for the hydrogen evolution reaction (HER) in alkaline solution (0.1 M KOH). Various GC-MC electrodes were prepared by loading different amounts (ca. 0.2–0.8 mg cm⁻²) of each metal complex on GC electrodes. These electrodes were used as cathodes in aqueous alkaline solutions (0.1 M KOH) to efficiently generate H₂ employing various electrochemical techniques. The three metal complexes’ HER catalytic activity was assessed using cathodic polarization studies. The charge-transfer kinetics of the HER at the (GC-MC)/OH⁻ interface at a given overpotential were also studied using the electrochemical impedance spectroscopy (EIS) technique. The electrocatalyst’s stability and long-term durability tests were performed employing cyclic voltammetry (repetitive cycling up to 5000 cycles) and 48 h of chronoamperometry measurements. The catalytic evolution of hydrogen on the three studied MC surfaces was further assessed using density functional theory (DFT) simulations. The GC-CuC catalysts revealed the highest HER electrocatalytic activity, which increased with the catalyst loading density. With a low HER onset potential (E onset) of −25 mV vs. RHE and a high exchange current density of 0.7 mA cm⁻², the best performing electrocatalyst, GC-CuC (0.8 mg cm⁻²), showed significant HER catalytic performance. Furthermore, the best performing electrocatalyst required an overpotential value of 120 mV to generate a current density of 10 mA cm⁻² and featured a Tafel slope value of −112 mV dec⁻¹. These HER electrochemical kinetic parameters were comparable to those measured here for the commercial Pt/C under the same operating conditions (−10 mV vs. RHE, 0.88 mA cm⁻², 108 mV dec⁻¹, and 110 mV to yield a current density of 10 mA cm⁻²), as well as the most active molecular electrocatalysts for H₂ generation from aqueous alkaline electrolytes. Density functional theory (DFT) simulations were used to investigate the nature of metal complex activities in relation to hydrogen adsorption. The molecular electrostatic surface potential (MESP) of the metal complexes was determined to assess the putative binding sites of the H atoms to the metal complex.
1. Introduction

In the worldwide energy demand plan, the generation of hydrogen (H₂) gas, a non-polluting multi-purpose energy carrier with a variety of properties and prospective industrial applications, has become a top priority. For efficient generation of H₂ from fossil fuels, several conventional technologies were used. Natural gas was utilized to manufacture H₂ through an industrial process called natural gas steam reforming, because it was plentiful, inexpensive, and contained the greatest H₂ content of all fossil fuels. Nonetheless, there are two key downsides to this industrial process: high energy use and CO₂ emissions. As a result, scientists around the world have been working to create safe, cost-effective (low-energy) ways for producing CO₂-free H₂ from non-fossil fuel sources.

In terms of sustainability and environmental impact, water electrolysis is the most promising technique for creating high purity H₂ from renewable energy sources, as it solely produces oxygen as a by-product and emits no greenhouse gases. The rate of the cathodic half-reaction of water splitting, viz. the hydrogen evolution reaction (HER), determines the efficiency of the water electrolysis. A highly efficient and robust heterogeneous electrocatalyst is required to accelerate the HER kinetics. The efficient electrocatalyst should be able to reduce the HER overpotential below 100 mV. The greater the activity of a catalyst, the lower its overpotential.

The efficient electrocatalyst should also have the highest possible electrochemically active surface area (ECSA), which is the active region on the catalyst’s surface that is accessible for the charge transfer process, as well as a high faradaic efficiency, which is calculated as the ratio of the experimentally determined quantity of H₂ to the calculated (theoretical) quantity of H₂ assuming 100% faradaic efficiency. In addition to the electrochemical parameters mentioned above that can be used to evaluate the electrocatalyst’s performance, the coveted electrocatalyst should have a high turnover frequency (TOF), which is defined as the amount of reactants that a catalyst can convert per catalytic site per unit time into the required product, as well as good stability.

All of the above-mentioned conditions for a desirable electrocatalyst were met using platinum (Pt) and Pt-based materials. As a result, among active electrocatalysts reported in the literature, such materials remain the most efficacious electrocatalysts for the HER. However, because of their high cost and rarity, their usage in industry has been severely limited. To meet the industry’s need for H₂ generation, substantial scientific research has been performed to develop and manufacture non-Pt-based cost-effective and efficient electrocatalysts. For example, according to Jaramillo et al., MoS₂ demonstrated exceptional HER catalytic performance. There has recently been a lot of interest in studies on carbon-based catalysts with low overpotentials for the HER, such as N, B, or S-doped carbons.

H₂ may also be produced efficiently using molecular electrocatalysts, such as transition-metal complexes, which are the most stable materials and are used to store energy in chemical bonds as well as generate electricity through hydrogen and oxidation production. For instance, recently, two complexes with bipyridine ligand, namely [Co(bpy)₃(SCN)₂] and [Ni(bpy)₃(SCN)₂] were synthesized and characterized by C.-L. Wang et al. Both showed promising catalytic activity for the hydrogen evolution reaction (HER). In an aqueous solution of neutral buffer (pH 7.0), at an overpotential of 837.6 mV, complexes 1 and 2 electro-catalyzed the HER with turnover frequency (TOF) values of 699.6 and 1469 mol H₂ per mol catalyst per h, respectively.

In another study, Z. Niu et al. prepared three molecular catalysts with square planar geometries based on mononuclear nickel(nn) complexes, and investigated their catalytic activity for H₂ generation in aqueous and non-aqueous solutions using cyclic voltammetry. The hydrogen production in both solutions was confirmed by controlled-potential electrolysis using gas chromatography. R. Li et al. synthesized two Co(nn)/Ni(nn) complexes based on nitrogenous heterocyclic ligand and explored them as stable electrocatalysts for the HER.

Incorporating transition metal (TM) complexes into cluster-based chalcogenide frameworks is a useful synthetic method for inducing structural diversity and controlling optoelectronic characteristics, which may boost electrocatalytic performance even further. For instance, J. Li et al. solvothermally synthesized and structurally characterized three novel isostructural neutral inorganic–organic open frameworks of gallium thioantimonate that are covalently bound with certain TM complexes. Thermal and chemical stability were good in the synthesized materials. The title compounds have tunable photocatalytic H₂ evolution activity thanks to an adjustable TM center.

The hydrogenase enzymes are depicted as naturally occurring microorganisms that efficiently catalyze the reversible interconversion of protons to hydrogen using earth-abundant metals (Ni and Fe) with high turnover rates 6000–9000 ([Fe–Fe]) and 700 ([Ni–Fe]) moles of H₂ per mole of enzyme per second for the HER. However, enzymes are difficult to produce in sufficient quantities for commercial uses, and their stability outside of their native system is generally limited. For these reasons, the structural and functional mimicking of hydrogenase’s active site has recently acquired attention. As a result, the development of highly efficient, stable, and robust HER molecular electrocatalysts for the production of hydrogen, which are inexpensive and made of earth-abundant transition metal elements and different ligand combinations, has attracted the community’s interest.
complexes. The catalytic evolution of hydrogen on the three supported complexes was further studied using density functional theory (DFT) simulations.

2. Experimental

2.1 Materials and general methods

All chemicals were of commercial grade and were used without extra purification. The ligand potassium hydrotris(3-methyl-5-phenylpyrazolyl)borate (KTpMePh) and its metal complexes MC (Scheme 1) were synthesized following a previously published method, which is fully described in Section S1 (ESI†).

2.2 Electrochemical measurements

Electrochemical measurements were conducted in 0.1 M KOH electrolyte using a standard double-jacketed three electrodes electrochemical cell. Various electrochemical techniques were applied to evaluate HER electrocatalytic activity and stability of the tested materials, as fully reported in Section S2 (ESI†).

3. Results and discussion

3.1 Characterization of the ligand and its metal complexes

The syntheses of TpMeMeCuCl(H2O)] (CuC), [TpMeMeNiCl] (NiC), and [TpMeMeFeCl2(H2O)] (FeC) (TpMeMe = tris(3,5-dimethylpyrazolyl)borate) were carried in absolute methanol by the reaction of the tripod ligand KTpMeMe with an equal amount of the corresponding MCln salt (M = Cu, Ni, and Fe). The sterically hindered methyl substituents at the pyrazolyl rings were found to have a major requirement for avoiding more chelate formation and stabilization of the formed complexes. The obtained metal complexes MC were stable in the solid state as well as in solution under atmospheric conditions. Analytical spectral features confirmed that all complexes MC have a 1 : 1 (metal : ligand) stoichiometry. The structure of FeC was octahedral whereas CuC was square pyramidal and NiC complex has a quaternary surface arrangement (Scheme 1).

DFT calculations using B3LYP/lan12dz level of theory were used to explore the structures of metal complexes MC. Fig. 1

![Scheme 1](https://example.com/scheme1.png)

**Scheme 1** The structures of ligand KTpMeMe and its metal complexes MC.

![Fig. 1](https://example.com/fig1.png)

**Fig. 1** The optimized structures of (a) CuC, (b) NiC, and (c) FeC at B3LYP/lan12dz level of theory in the gas phase.
shows the optimized structures of the metal complexes in alkaline medium, where the chloride anions (Cl\textsuperscript{−}) were substituted by hydroxyl groups (OH\textsuperscript{−}). The structures of the metal complexes showed that each metal ion is coordinated to three nitrogen atoms from the anionic ligand Tp\textsuperscript{MeMe−}, forming symmetrical structures. The stable copper(II) complex CuC binds with one more water molecule (Cu–O\textsubscript{48H\textsubscript{2}} = 2.16 Å), besides the OH\textsuperscript{−} groups (Cu–O\textsubscript{46} = 1.94 Å). While the iron(III) complex FeC binds with one water molecule (Fe–O\textsubscript{50H\textsubscript{2}} = 2.05 Å) and two hydroxyl groups (Fe–O\textsubscript{47} = 1.78 Å and Fe–O\textsubscript{46} = 1.78 Å). These results were established by the TGA data and confirmed by the stable DFT structures. On the other hand, the neutral nickel(n) complex NiC binds with only one OH\textsuperscript{−} group (Ni–O\textsubscript{46} = 1.80 Å).

The formation of metal complexes (MC) was further evidenced from the high energy shift of ν(B–H) stretching vibration near 2500 cm\textsuperscript{−1} in the IR spectra of the metal complexes compared to that of the free ligand KTp\textsuperscript{MeMe−}, which appeared at 2436 cm\textsuperscript{−1} (Fig. 2). Similar shifts were reported for other Tp complexes.\textsuperscript{32} The IR spectra also showed other characteristic stretching and bending vibration bands at ~2920, 1680, and 1150 cm\textsuperscript{−1}, assigned to ν(C–H), ν(C=N), and δ(C–H), respectively. Raman spectra of the ligand and its metal complexes (Fig. 3) revealed almost identical vibration bands with those found in their IR spectra. The non-electrolytic chloride ion in all complexes exhibits vibration bands, characteristics of the terminally-coordinated chloride ions.\textsuperscript{33} Additionally, the three metal complexes all featured weak bands at around 442, 449, and 452 cm\textsuperscript{−1}, suggesting the existence of M–N bonds.\textsuperscript{34}

The thermogram TGA-DTG-DSC of FeC (Fig. 3A) shows that the complex dehydrates one water molecule with an exothermic peak at 98 °C. It decomposes in the range of 120–900 °C (endothermic peak, DSC: 410 °C) to eliminate the organic part with a mass loss of 63.86% (calc. 63.27%), forming a mixture of ferrous(II) oxide and potassium oxide. The thermal analysis of

**Fig. 2** (A) FT-IR and (B) Raman spectra of the ligand KTp\textsuperscript{MeMe} and its metal complexes MC.

**Fig. 3** TGA-DTG-DSC curves of (A) [Tp\textsuperscript{MeMe−}FeCl\textsubscript{2}(H\textsubscript{2}O)] (FeC) and (B) [Tp\textsuperscript{MeMe−}CuCl(H\textsubscript{2}O)] (CuC).
the copper(II) complex CuC indicates that its decomposition takes place in several stages (Fig. 3B). The thermogram shows that the complex decomposes to eliminate the organic part in the range of 105–820 °C (endothermic peak, DSC: 461 K) with a mass loss of 81.98% (calc. 81.52%), generating a mixture of copper(II) and potassium oxide (found 19.64%; calc. 18.73%).

The electronic absorption spectrum of CuC in methanol (Fig. 4) exhibited weak bands in the lowest energy visible region attributable to d–d transitions of the Cu(II) ion in a square-pyramidal environment. The absorption spectrum of NiC displays three absorption bands at 330, 560, and 635 nm. An examination of these bands indicates that the complex has a tetrahedral geometry with ground state 3T1. In the regular tetrahedral nickel(II) complexes, only one d–d transition is observed in the visible region. The observation of the three transitions in the spectrum of NiC complex may be attributed to the distortion in the tetrahedral symmetry around the nickel(II) center. This spectral behavior is comparable with analogous tetrahedral nickel(II) complexes, only one d–d transition is observed in the visible region.

The electronic spectrum of FeC exhibits four d–d transitions at 325, 374, 490, and 588 nm, assignable to 5A1g  4T1g (G), 6A1g  4T2g (G), 6A1g  4Eg (G) and 6A1g  4Eg (D), respectively in the octahedral stereochemistry. Hence the nickel(II) ion has an sp3 hybridization with tetrahedral geometry. The intensity of the UV band at around 330 nm is consistent with a ligand-centered π → π* transition or a charge-transfer transition.

The magnetic moment value of CuC was determined to be 1.98 BM, confirming the presence of magnetically dilute Cu(n) complex whereas the magnetic moment for FeC was close to the spin-only value of 5.63 and 5.44 μB, which is characteristic of high spin of d5 configuration.

The quantitative elemental analysis of the three complexes CuC, NiC, and FeC were also determined using energy dispersive X-ray fluorescence (EDXRF) spectroscopy at atmospheric pressure and the obtained spectra are illustrated in Fig. 5. The Ar-Kz was found in all the spectra at 2.957 keV and it has no influence on the characteristic lines of Cu, Ni and Fe. Additional impurities of Ca and Zn were also found, which were originated from the used Mylar foil. In the case of complex NiC, the characteristic radiation Ni-Kz and Ni-Kβ was measured at 7.473 keV and 8.265 keV respectively. Addition escape peak of Ni-Kz was found at 5.733. The spectrum belongs to complex CuC, the Cu-Kz and Kβ were detected successfully at 8.042 keV and 8.906 keV respectively. Whereas in the case of complex FeC, the characteristic radiation of Fe–K lines could be easily detected free from interference at 5.895 keV and 6.491 keV for Kz and Kβ respectively. Escape peak of Fe-Kz could be recognized also at 4.155 keV. The % of the coordinated TpMe,Me in all complexes were calculated and omitted from the fundamental parameter program. This is due to the limitation of EDXRF setup whereas there is not possibility to detected elements of atomic number less than 11 (Z < 11). The weight% of complexes CuC, NiC, and FeC were found to be 15.27 ± 0.18, 15.22 ± 0.18, and 12.55 ± 0.18, respectively.

3.2 HER electrocatalytic studies

3.2.1 Cathodic polarization measurements. Fig. 6(a) depicts the cathodic polarization curves measured for the studied electrocatalysts, namely GC-loaded metal complex electrodes at various loading densities (ca. 0.2–0.8 mg cm−2).

Current densities in this work are obtained by dividing currents by the electrochemical active surface area (EASA), calculated from cyclic voltammetry measurements, see later. The various HER electrochemical kinetic parameters (Table 1) associated with such polarization measurements were derived from Tafel plots, Fig. 6(b).

Such parameters, which include exchange current density (j0), cathodic Tafel slope (β), and the overpotential required to deliver a current density of 10 mA cm−2 (η100), are essential to evaluate and compare the HER catalytic activity of the studied metal complexes 1–3.
Table 1  Mean value (standard deviation) of the HER electrochemical kinetic parameters estimated for the investigated catalysts. Data derived from Fig. S1(b)

| Tested cathode   | \(E_{\text{HER}}/\text{mV (RHE)}\) | \(-\beta_\text{a}/\text{mV dec}^{-1}\) | \(j_0/\text{mA cm}^{-2}\) | \(\eta_{10}/\text{mV}\) |
|------------------|-----------------------------|-----------------|----------------|---------------|
| Bare GC electrode| —                          | 254(6)         | 6.3(0.15) \times 10^{-4} | —             |
| GC-FeC (0.2 mg cm\(^{-2}\)) | 305(5)                  | 162(3.3)  | 1.1(0.05) \times 10^{-2} | 467(8)      |
| GC-FeC (0.4 mg cm\(^{-2}\)) | 253(3.4)              | 160(3.2)  | 2.5(0.07) \times 10^{-2} | 423(7.6)    |
| GC-FeC (0.8 mg cm\(^{-2}\)) | 230(3.2)              | 158(3)    | 3.7(0.13) \times 10^{-2} | 372(5)      |
| GC-NiC (0.2 mg cm\(^{-2}\)) | 180(2.6)              | 157(2.8)  | 4.3(0.33) \times 10^{-2} | 278(4.2)    |
| GC-NiC (0.4 mg cm\(^{-2}\)) | 107(2.2)              | 160(2.6)  | 10(0.12) \times 10^{-2}  | 250(4.6)    |
| GC-NiC (0.8 mg cm\(^{-2}\)) | 80(1.7)                | 155(2.2)  | 14(0.35) \times 10^{-2}  | 213(4)      |
| GC-CuC (0.2 mg cm\(^{-2}\)) | 45(1.5)                | 118(1.8)  | 29(0.6) \times 10^{-2}   | 176(4.4)    |
| GC-CuC (0.4 mg cm\(^{-2}\)) | 30(1.1)                | 114(2.1)  | 60(0.9) \times 10^{-2}   | 140(2.6)    |
| GC-CuC (0.8 mg cm\(^{-2}\)) | 25(0.5)                | 112(1.6)  | 70(1.1) \times 10^{-2}   | 120(2.2)    |
| Pt/C             | 10(0.2)                 | 108(1.2)  | 88(1.2) \times 10^{-2}   | 110(1.4)    |
efficacious transfer of charge during the water molecules reduction process thus, boosts the HER kinetics.

3.2.2 EIS measurements. EIS measurements were also carried out at various overpotentials (ca. 0.1–0.5 V). The goal is to further evaluate HER kinetics and surface phenomena. Fig. 7 depicts the Nyquist plots recorded for the three studied catalysts as a function of the catalyst’s loading density (ca. 0.2–0.8 mg cm$^{-2}$) in deaerated 0.1 M KOH at an overpotential of 0.5 V vs. RHE.

Two distinct depressed semicircles can be seen in the Nyquist plots of all materials investigated. The electrocatalytic HER on Ni,$^{47}$ Co,$^{48}$ and some of their alloys$^{49-51}$ in alkaline solutions exhibited such impedance behavior. Similar impedance results were reported earlier in our lab for HER investigations on Co$_{50}$Ni$_{23}$Ga$_{26}$Al$_{1.0}$ magnetic shape memory alloy,$^{52}$ reduced graphene oxide nanosheets, some metallic nanoparticles-decorated titanium substrates, and cathodically activated carbon felt samples.$^{53-56}$

At high frequency values, a first small semicircle (with diameter $R_1$) was visible, whereas a large second semicircle (with diameter $R_2$) appeared at low frequencies. Each of them can be assigned to the resistance–capacitance (RC) network. Each RC network at the catalyst/KOH interface corresponds to the charge-transfer resistance ($R_1$ for the first semicircle and $R_2$ for the second, $R_{\text{total}} = R_1 + R_2$) and associated double-layer capacitance ($C_1$ for the first semicircle and $C_2$ for the second, $C_{\text{total}} = C_1 + C_2$).$^{56}$ The two semicircles appear as conspicuous humps in the related Bode graphs, Fig. S10 (ESI†).

It is obvious from Fig. 7 that the size of the two semicircles varies according to the type of the tested catalyst and its loading density. Also, referring to Fig. S10,† at a fixed loading density, both semicircles are clearly potential-dependent. These findings indicate that the two semicircles’ size is connected to the HER kinetics.$^{57}$

For comparison, EIS responses of bare GC and Pt/C electrodes with a single capacitive loop (Fig. S11, ESI†) were also recorded. The RC networks’ elements were estimated using the equivalent circuit illustrated in Fig. S12 (ESI†), allowing us to gain a better grasp of the HER kinetics. This electrical equivalent circuit was described in more detail elsewhere.$^{54}$

Based on the impedance data fitting, when the loading density of the catalyst was raised, the values of both $R_1$ and $R_2$, and hence the $R_{\text{ct}}$ value ($R_1 + R_2$), reduced considerably, indicating higher HER catalytic activity. For example, the $R_{\text{ct}}$ values recorded for the NiC catalyst decreased from 315 $\Omega$ cm$^2$ at a loading density value of 0.2 mg cm$^{-2}$ to 248 and 145 $\Omega$ cm$^2$ at loading densities 0.4 and 0.8 mg cm$^{-2}$, respectively. The CuC electrocatalyst showed the lowest $R_{\text{ct}}$ value among the other studied samples at any studied loading density thus, confirming the excellent HER catalytic activity of the CuC electrocatalyst. For instance, at a loading density value of 0.8 mg cm$^{-2}$, the CuC electrocatalyst recorded an $R_{\text{ct}}$ value of 51 $\Omega$ cm$^2$. This value is very close to that of the Pt/C (47 $\Omega$ cm$^2$), and lower than those measured for GC-NiC (145 $\Omega$ cm$^2$) and GC-FeC (450 $\Omega$ cm$^2$) at the same loading density.

Another critical electrochemical parameter, namely the Tafel slope ($\beta_b$), is also utilized to further assess the studied electrocatalysts’ HER catalytic leverage towards the HER. The $\beta_b$ value recorded here for the commercial Pt/C catalyst is found to be 108 mV dec$^{-1}$, close to that reported in the literature under the same operating conditions, 111 mV dec$^{-1}$,$^{54}$ meaning that the electrochemical measurements employed here are reliable.

The alkaline HER step mechanism can be represented by eqn (1)–(3):$^{59}$

$$\text{H}_2\text{O} + e^- \rightarrow \text{H}_{\text{ads}} + \text{HO}^- \quad \text{(Volmer step)} \quad (1)$$

$$b = \frac{2.34RT}{aF} \equiv 120 \text{ mV dec}^{-1}$$

The Volmer step is followed by either a combination of adsorbed species to form molecular H$_2$:

$$\text{H}_{\text{ads}} + \text{H}_{\text{ads}} \rightarrow \text{H}_2 \quad \text{(Tafel step)} \quad (2)$$

$$b = \frac{2.34RT}{(1 + a)F} \equiv 30 \text{ mV dec}^{-1}$$

or the desorption Heyrovsky step:

$$\text{H}_2\text{O} + e^- + \text{H}_{\text{ads}} \rightarrow \text{H}_2 + \text{HO}^- \quad \text{(Heyrovsky)} \quad (3)$$

$$b = \frac{2.34RT}{2F} \equiv 40 \text{ mV dec}^{-1}$$
The Pt/C catalyst’s $\beta_e$ value obtained here (108 mV dec$^{-1}$), which is comparable with that reported in the literature (111 mV dec$^{-1}$) under the same operating conditions,$^{29}$ is close to the standard one (120 mV dec$^{-1}$) reported in eqn (1), meaning that the alkaline HER mechanism comprises the Volmer step as the rate-limiting.$^{60}$

The Tafel lines of FeC and NiC catalysts are parallel to each other, irrespective of the value of the catalyst's loading density, with $\beta_e$ values close to 160 mV dec$^{-1}$. Also, the three studied CuC catalysts, namely CuC (0.2 mg cm$^{-2}$), CuC (0.4 mg cm$^{-2}$), and CuC (0.8 mg cm$^{-2}$) exhibited parallel Tafel lines, but with reduced $\beta_e$ values (118, 114, and 112 mV dec$^{-1}$, respectively). These findings indicate that the kinetics of the HER on the CuC catalyst’s surface is faster than on the surfaces of FeC and NiC catalysts. This is evidenced from the marked drop in the Tafel slope values of the CuC catalyst since lower Tafel slopes often allude to a large number of catalytic active sites being available.$^{60,61}$

The CuC (0.8 mg cm$^{-2}$) catalyst’s $\beta_e$ value (112 mV dec$^{-1}$) is close to that of the Pt/C (108 mV dec$^{-1}$) thus advocating a mechanism for the HER of that catalyst controlled by the Volmer step.$^{60}$ The results show that the CuC catalyst can efficiently catalyze HER in 0.1 M KOH solution, which is similar to the most active HER electrocatalysts reported in the literature, Table S1 (ESI†).

The electrochemical active surface area (EASA) is another essential parameter used to compare different electrocatalysts’ activity, as the kinetics of the HER over electrocatalysts is directly linked to their ECSA. Being directly related to ECSA, double layer capacitance ($C_{dl}$), determined from cyclic voltammetry measurements in non-faradic region, Fig. S1 (ESI†), is used to calculate ECSA.$^{62-64}$ From the slopes of the current density versus scan rate ($v$) plots, Fig. S1 (ESI†), for any studied meal complex, $C_{dl}$ values were found to increase with increase in the loading density. At any investigated loading density, this increase in $C_{dl}$ enhances following the sequence: FeC < NiC < CuC. These findings confirm the high catalytic performance of the CuC electrocatalyst since electrocatalysts with higher $C_{dl}$ values can provide higher adsorption density of reactive sites, and therefore promote efficient charge transfer, resulting in enhanced catalytic activity.

Eqn (4) is used to calculate the ECSA from $C_{dl}$.$^{65}$

$$EASA = C_{dl}/C_s \quad (4)$$

where $C_s$ is the electrode’s specific capacitance, typically between 20 and 40 $\mu$F cm$^{-2}$. Introducing $C_s = 30 \mu$F cm$^{-2}$, the average value of the 1.0 cm$^2$ flat electrode, in eqn (4) gives EASA values of 380, 527, and 773 cm$^2$ for FeC (0.2 mg cm$^{-2}$), NiC (0.2 mg cm$^{-2}$), and CuC (0.2 mg cm$^{-2}$), respectively. These values significantly increased upon increasing the catalyst’s loading density. In all cases, the CuC electrocatalyst recorded the largest EASA value amongst.

For any studied catalyst, the value of $n$ is found to enhance with the catalyst’s loading density. At any tested loading density, the value of $n$ increases obeying the order: FeC < NiC < CuC. These findings confirm the CuC’s highest HER catalytic performance.

To further evaluate the catalytic performance of the investigated catalysts at a given overpotential, the turnover frequency (TOF) was calculated, which is the number of hydrogen molecules created per second per active site.$^{66}$ The TOF values for the tested catalysts were computed using the EASA values in Table 2 at an overpotential of 500 mV, as discussed previously.$^{52}$ Table 2 depicts the obtained TOF values, which feature a dramatic increase in the value of TOF as the loading density of the examined catalyst increases. The TOF value is always higher for the CuC at any studied loading density. To our knowledge, the TOF value recorded for the best catalyst, namely CuC, is comparable to the most efficient molecular catalysts and greater than many others described in the literature for electrochemical H$_2$ production. Table S1 (ESI†) summarizes these comparisons.

Table 2  Double-layer capacitance ($C_{dl}$), electrochemical active surface area (EASA), net voltammetry charge ($Q$), the number of active sites ($n$), and the turnover frequency (TOF) estimated for the investigated electrocatalysts from CV measurements Fig. S1–S9 (ESI)

| Tested cathode | $C_{dl}$/mF cm$^{-2}$ | ECSA/cm$^2$ | $Q \times 10^3$/C | $n \times 10^8$/mol | TOF @ 0.5 V |
|---------------|-------------------|-------------|-----------------|-----------------|-------------|
| GC-FeC (0.2 mg cm$^{-2}$) | 11.26 | 375.3 | 4.2 | 2.2 | 872 |
| GC-FeC (0.4 mg cm$^{-2}$) | 20.27 | 675.7 | 6.8 | 3.5 | 988 |
| GC-FeC (0.8 mg cm$^{-2}$) | 30.55 | 1018.3 | 11.2 | 5.8 | 1124 |
| GC-NiC (0.2 mg cm$^{-2}$) | 15.55 | 518.3 | 7.4 | 3.8 | 927 |
| GC-NiC (0.4 mg cm$^{-2}$) | 28.32 | 944.0 | 12.4 | 6.4 | 1192 |
| GC-NiC (0.8 mg cm$^{-2}$) | 44.43 | 1481.0 | 19.8 | 10.3 | 1338 |
| GC-CuC (0.2 mg cm$^{-2}$) | 21.2 | 706.7 | 11.6 | 6.0 | 1154 |
| GC-CuC (0.4 mg cm$^{-2}$) | 36.71 | 1223.7 | 17.8 | 9.2 | 1510 |
| GC-CuC (0.8 mg cm$^{-2}$) | 53.02 | 1767.3 | 26.9 | 13.9 | 1820 |
Origin of catalytic activity (DFT calculations). The DFT calculations were performed to explore the nature of the metal complex activities towards the hydrogen adsorption process. In order to assess the possible binding sites of the H atoms to the metal complexes, the molecular electrostatic surface potential (MESP) values of the metal complexes were calculated (Fig. 8). The calculated MESP shows the presence of electrostatic negative potential on the metal atom, which was found to be higher in Ni (−16.31 kcal mol⁻¹), while the N atoms on pyrazole ring carry negative electrostatic potential (−13.8 kcal mol⁻¹). On the other hand, the complex FeC owns a positive electrostatic potential. Accordingly, the metal complexes show two possible sites for the adsorption of the hydrogen atom to initiate the process. The DFT binding energy of the H atom (Fig. 9) to the available sites on the metal complexes reveals that the binding to the metal complexes is more stable than to the N atom by 5.1, 7.67, and 8.52 kcal mol⁻¹ for complexes CuC, NiC, and FeC, respectively.

To retain the molecular nature of the metal complex MC catalysts under the alkaline conditions, it is necessary for the ligand to be tightly bound to the metal center, these catalysts were generated by treating metal(n) chloride (MCln) with one equivalent of ligand in absolute methanol, in which one tris(-pyrazolyl)borate anion (TpMeMe⁻) is bound to the metal center with the fourth coordination site, occupied by a chloride in the case of NiC, whereas in CuC and FeC, the ligand is bound to the metal center with one and two more water molecules, respectively. Upon the addition of one equivalent of strong base, the chloride ligand in all complexes is substituted by hydroxide (OH⁻) anion, and the resulting Cu(n)-bound hydroxide complex, for example, adopts a square pyramidal structure with the OH-donor. Upon the addition of one equivalent of hydrochloric acid, the OH ligand can be protonated to regenerate the aqua complex. This reversible protonation of the ligand is an attractive feature for a water-oxidation catalyst, as it may allow for proton-coupled electron transfer (PCET), a common reaction step in water oxidation.

The actual catalyst for the current catalytic systems is generated in an alkaline medium, where the hydroxide anion must bind to the central metal ion of the catalyst. Thus, the initiation step of the present catalytic reaction involves binding of OH⁻ to the central metal ion of the catalyst. In this context, the octahedral Fe(III) complex is coordinately saturated, and thus has no vacant coordination sites, which can be utilized to bind any reacting species during the catalytic process. The ability of such coordinately saturated complexes to dissociate (ML₆ ⇔ ML₅ + L) and consequently the energy of dissociation is one of the key factors in imparting catalytic properties to these metal complexes. On the other hand, the five coordinated square pyramidal Cu(II) complex and the four coordinated tetrahedral Ni(n) complex already have a vacant coordination site for the substrate binding and thus need no energy for ligand dissociation as in the case of the octahedral Fe(III) complex.

Although Cu(n) and Ni(n) complexes are both coordinatively unsaturated, Cu(n) complex exhibits higher catalytic activity. The superiority of Cu(n) can be explained by the possible mechanistic implication of the six coordinate intermediate [H₂ML₆-OH]. Based on Jahn–Teller theorem, this hexacoordinated species is highly unstable in the case of Cu(n) complex as compared to Ni(n) complex. According to the Jahn–Teller theorem, the hexa-coordinated Cu(n) complexes are usually associated with a tetragonal distortion reducing the symmetry from O₆ to D₄h.⁶⁷,⁶⁸ Thus, we propose that rapid structural reorganization of this active intermediate leads to rapid product formation and release. These structural changes will reduce the coordination number from six to five and return the catalyst to its original, stable form. As for the Ni(n) complex does not suffer from the Jahn–Teller effect, its catalytic reaction proceeds more slowly than it does in the case of the Cu(n) complex.

Copper and nickel are more labile than iron. This sequence of the lability degree could also be due to the electronic configuration of the metal center in these metal complexes and can be illustrated from the following discussion. The 3d-
electronic configuration (electron density) of the metal ion is a key factor for determining the degree of lability or inertness of transition metal complexes. According to valence bond theory (VBT), the electron density of MC complexes is increasing in the following order CuC (4 lone pairs) > NiC (three lone pairs) > FeC: (no lone pair). This is in accordance with the HER catalytic performance, in which the three tested metal complexes' activity follows the sequence: CuC > NiC > FeC.

Stability test. The best performing electrocatalyst, namely GC-CuC (0.8 mg cm$^{-2}$), was further evaluated for stability and long-term durability using CV measurements. Fig. 10 shows the first cycle of the catalyst in comparison to the 6000th cycle.

In these solutions, the catalyst is particularly stable, as indicated by the hume loss in current after 6000 continuous potential cycles. The Tafel slope obtained after the last cycle (115 mV dec$^{-1}$) is close to that recorded after the first cycle (112 mV dec$^{-1}$), suggesting that catalyst potential cycling has a negligible influence on the mechanism of the HER. Because the current remains nearly constant throughout the operation, the chronoamperometry measurements (inset of Fig. 10) give an additional corroboration for the catalyst stability.

4. Conclusion

Three tripod-containing ligand metal complexes namely, [TpMeMeCuCl(H$_2$O)] ($\text{CuC}$), [TpMeMeNiCl] ($\text{NiC}$), and [TpMeMeFeCl$_3$(H$_2$O)] ($\text{FeC}$)-modified glassy carbon (GC) electrodes were fabricated and examined as electrocatalysts for the efficient generation of H$_2$ at different loading densities (0.2, 0.4, and 0.8 mg cm$^{-2}$) in 0.1 M KOH. HER electrocatalytic activity of these materials was investigated using LSV and EIS techniques. The HER electrocatalytic activity of the investigated catalysts was found to increase with increase in their loading density. Among the tested catalysts, at any studied loading density, the GC-CuC exhibited the highest catalytic activity for the HER. At a loading density of 0.8 mg cm$^{-2}$, the GC-CuC recorded a low HER onset potential ($E_{\text{HER}}$) of $-25$ mV vs. RHE and a high exchange current density of 0.7 mA cm$^{-2}$. Further, to achieve a current density of 10 mA cm$^{-2}$, the most performing electrocatalyst required an overpotential of 120 mV and recorded a Tafel slope of $-112$ mV dec$^{-1}$. Under the same operating conditions, the best catalyst’s HER electrochemical kinetic parameters were close to those of the commercial Pt/C ($-10$ mV vs. RHE, 0.88 mA cm$^{-2}$, 108 mV dec$^{-1}$, and 110 mV to yield a current density of 10 mA cm$^{-2}$) thus, highlighting its outstanding HER electrocatalytic activity. The nature of metal complex activities in relation to hydrogen adsorption was investigated using DFT simulations. To examine the likely binding sites of the H atoms to the metal complex, the molecular electrostatic surface potential (MESP) of the metal complexes was calculated. The estimated MESP reveals the presence of electrostatic negative potential on the metal atom, which was higher in Ni ($-16.31$ kcal mol$^{-1}$). The negative electrostatic potential on the N atoms on the pyrazole ring was calculated to be ($-13.8$ kcal mol$^{-1}$). As a result, the metal complexes exhibit two viable sites for hydrogen atom adsorption to initiate the process. The cationic iron(II) complex, on the other hand, has a positive electrostatic potential which may be the reason behind its low HER catalytic activity compared with Cu and Ni complexes.

Author contributions

All authors have read and agreed to the published version of the manuscript.

Conflicts of interest

The authors declare no conflict of interest.

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