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# Redox potential prediction of Fe(II)/Fe(III) complexes: a density functional theory and graph neural network approach

Fakhrul H. Bhuiyan,<sup>a</sup> Hassan Harb,<sup>b</sup> Rajeev Surendran Assary<sup>b</sup> and Álvaro Vázquez-Mayagoitia<sup>\*a</sup>

This work presents an integrated computational approach that combines tight-binding density functional theory (DFT) with standard DFT calculations to accurately compute the redox potential of micro-solvated iron-based transition metal complexes. Comparison with experimental and computational reference values confirmed the reliability of the computational approach. A comprehensive redox dataset of 2267 iron complexes was generated using the computational approach for machine learning (ML) applications. Chemical analysis of the ligand space in the dataset provided detailed insights into how different ligand classes and local Fe-coordination environments can systematically influence redox potential. Finally, a graph neural network (GNN) framework featuring automated graph data generation directly from 3D Cartesian coordinates was developed to model and predict redox potentials of metal complexes. Four distinct GNN architectures (GCN, GAT, DimeNet++, and SchNet) were evaluated using the curated redox potential dataset. The best-performing model achieved a root mean squared error of  $0.24 \pm 0.01$  V, representing state-of-the-art performance for redox potential prediction of transition metal complexes. Feature attribution analysis using integrated gradients provided insights into the factors influencing the GNN model predictions. This combined DFT-ML workflow provides both predictive power and chemical insight, offering a scalable pathway to accelerate the discovery and optimization of transition metal complexes for any application.

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## Introduction

The redox potential is a useful measure of the tendency of a chemical system to gain or lose electrons and is therefore a relevant indicator of the reactivity and stability of transition metal complexes.<sup>1</sup> The redox of iron-based metal complexes is of significant interest because of its relevance in various applications, including energy storage technologies such as redox flow batteries (RFBs),<sup>2–6</sup> bioinorganic chemistry,<sup>7–9</sup> and catalysis.<sup>10–12</sup> The redox potential of iron-based metal complexes can vary significantly depending on the coordinating ligands, reflecting their tunability and potential applications.<sup>12–14</sup> The ability to tune the redox potential of iron-based metal complexes through selective ligand complexation positions them as promising components in advanced electrochemical systems. However, the vast chemical space (billions) of possible ligand–metal combinations prohibits exhaustive experimental or accurate quantum-mechanical exploration.<sup>15,16</sup> As a result, there is a critical need for fast, accurate, and scalable methods to predict the redox potentials of iron-based metal complexes.

Experimentally, the redox potential of transition metal complexes can be measured using cyclic voltammetry,<sup>17,18</sup> differential pulse voltammetry,<sup>19</sup> or square-wave voltammetry.<sup>20,21</sup> Among these techniques, cyclic voltammetry is the most commonly used electrochemical technique that measures the electric current ( $I$ ) response of a redox-active material as its potential ( $V$ ) is linearly swept in a cyclic manner. The experimental determination of redox potentials is time-consuming and costly, rendering these experimental techniques unsuitable for high-throughput screening of a large chemical space of metal complexes.

Computational chemistry methods, particularly density functional theory (DFT), can complement experimental methods and serve as a valuable tool for determining redox potentials.<sup>22–24</sup> The DFT offers a good balance between computational cost and accuracy to simulate a wide range of chemical systems and their material properties. Quantum chemical calculations based on DFT have been successfully used in numerous studies to calculate standard redox potentials for complexes in aqueous solutions using implicit solvation models, often achieving good agreement with available experimental values.<sup>25–29</sup> Despite its utility, DFT calculations can be computationally expensive for large molecular systems and for high-throughput screening efforts.

<sup>a</sup>Computational Science Division, Argonne National Laboratory, Lemont, IL 60439, USA. E-mail: vama@alcf.anl.gov<sup>b</sup>Materials Science Division, Argonne National Laboratory, Lemont, IL 60439, USA

Classical molecular dynamics (MD) simulations using force fields have also been applied, though in a limited capacity, to study redox potentials. For example, molecular mechanics force fields were developed for transition-metal oxide clusters to estimate oxidation potentials.<sup>30,31</sup> Reactive force fields (ReaxFF) have been widely employed to model diverse chemical reactions,<sup>32–34</sup> but their lack of explicit electron treatment limits their use for redox studies. To overcome this, e-ReaxFF was introduced, enabling explicit electron treatment and applied to model redox processes such as silver halide reactions<sup>35,36</sup> and solid-oxide electrocatalysis.<sup>37</sup> However, e-ReaxFF remains in early development and is not yet widely adopted for redox potential calculations.

Machine learned interatomic potentials (MLIPs) can also be used to study chemical reactions in MD simulations.<sup>38</sup> However, the use of MLIPs for calculating redox potential is rare due to the inherent difficulty in modeling long range electrostatic interactions and performing charge equilibration using currently available MLIPs.<sup>39–41</sup> Modeling redox reactions using MLIPs require complex modification of the architecture of the MLIP model. One such modification of MLIP was recently demonstrated for the high-dimensional neural network potentials to capture the redox chemistry of Fe<sup>2+</sup> and Fe<sup>3+</sup> ions in water.<sup>42</sup> Despite the recent progress, application of MLIPs for redox potential calculation in MD simulations remains challenging and not suitable for high-throughput screening purposes.

Machine learning (ML) has emerged as a powerful approach to accelerate materials discovery by providing rapid property prediction capabilities. ML models can serve as fast surrogates for computationally intensive simulations, such as DFT, predicting properties in seconds making them suitable for high-throughput screening. For instance, Gaussian Process Regression models trained using fingerprints-based graph representation of organic ligands were used to predict redox potentials of organic molecules.<sup>43</sup> Graph neural networks (GNNs), among various ML approaches, are particularly well-suited for handling non-Euclidean data such as molecular structures.<sup>44,45</sup> GNNs operate on graph-structured data, where molecules are represented as graphs with nodes for atoms and edges for bonds. This approach allows GNNs to capture complex relationships between atoms and their chemical environments, leading to improved performance over conventional descriptor-based models.<sup>46</sup>

Several studies have demonstrated the effectiveness of GNNs in predicting redox potentials. For organic compounds, GNNs have outperformed classical ML models and have been used to screen large molecular databases.<sup>47</sup> A molecular graph attention network, a type of GNN, was developed to predict the redox potential of organic compounds using molecular structures, atomic properties, and bond attributes, outperforming other GNN variants.<sup>48</sup> Graph-based methods have also been applied to transition metal complexes. For instance, a graph-based descriptor model was used to model octahedral Fe(II/III) redox couples to show that redox potential increased with increasing weight of the complex but decreased for branched structures.<sup>26</sup> Although that work successfully demonstrated *in silico*

discovery of metal complexes from a large chemical space of ligands, the redox calculations and predictions remained constrained in a small, concentrated dataset containing less than 50 metal complexes. Despite the advances, a general methodology for the high-throughput screening of a large and diverse set of metal complexes, including iron-based ones, is highly desired.

In this work, we developed a general methodology for modeling and predicting the redox potential of thousands of iron-based metal complexes from the transition metal quantum mechanics (tmQM) dataset using high-throughput screening.<sup>49</sup> Calculation of the redox potentials led to the generation of a redox dataset containing 2267 iron complexes under explicit and implicit solvent effects. Extensive chemical analysis of the iron complexes in the dataset revealed key insights into the effect of ligand chemistry and iron-ion coordination in determining the redox behavior of a complex. The redox dataset was then used for training multiple GNN models through a neural network training framework developed here and model performances were compared. We introduce two edge-aware GNN models, RexGCN and RexGAT, that showed more precise predictions and reached earlier learning curve plateau than previously reported GNN models. To our knowledge this is the largest redox database for calculated redox potential for the Fe(II)/Fe(III) pair of structures observed experimentally and reported in the Cambridge Structural database. A schematic representation of our workflow is shown in the Fig. 1. Although the main focus of this work was on iron-based complexes, the methods developed here can be readily applied to model any metal complex from any dataset, suggesting the far-reaching utility of our approach.

## Methodology

### Data collection

The iron complexes used in this study were collected from the tmQM dataset (2024 release).<sup>49</sup> Built based upon structures retrieved from the Cambridge Structural database, the tmQM dataset is currently one of the largest openly available datasets for transition metal complexes. The 2024 release of this dataset contains the geometries and quantum mechanical properties of ~108 000 metal complexes including iron complexes.

We first filtered the tmQM dataset to include only iron complexes containing 60 or fewer atoms. The selected iron complexes were further filtered to retain only those containing an iron ion with an oxidation state of +2 (*i.e.*, Fe(II) complexes). We calculated the oxidation state using the following equation:

$$M_{\text{ox}} = Q_{\text{total}} - \sum_i^{N_{\text{ligands}}} q_i \quad (1)$$

where  $M_{\text{ox}}$  is the ion oxidation state,  $Q_{\text{total}}$  is the total charge of the metal complex as found in the tmQM dataset,  $N_{\text{ligands}}$  is the number of ligands in the metal complex, and  $q_i$  is the ligand charge. The number of ligands and their respective charges were calculated using the xyz2tomol\_tm method.<sup>50</sup> More details about the use of the xyz2tomol\_tm method in this work is



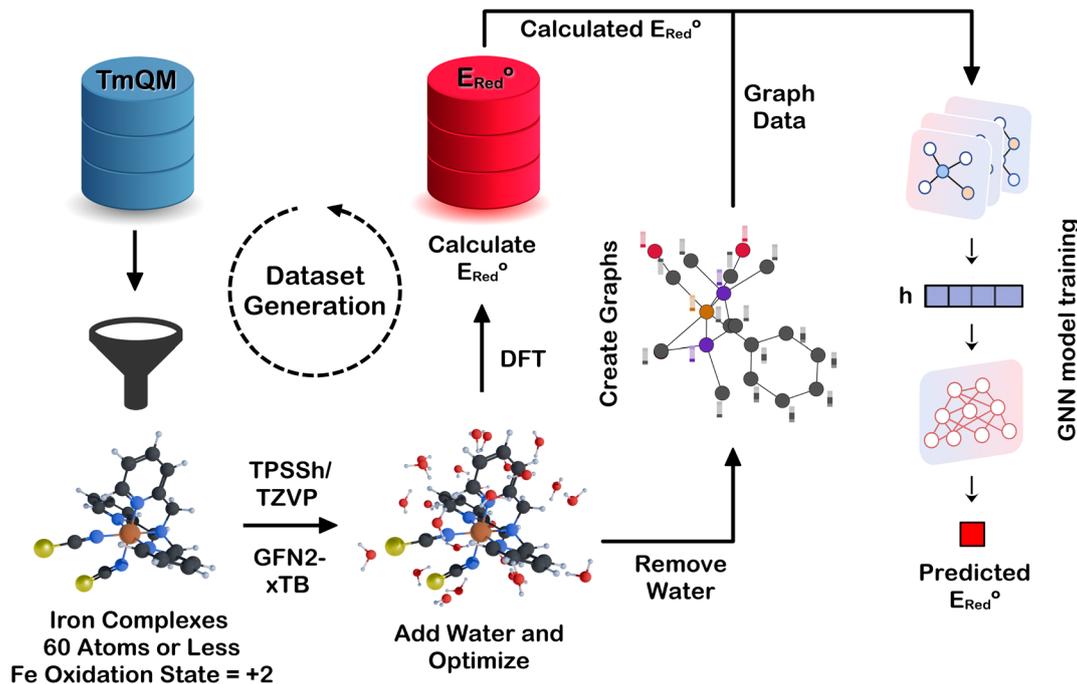


Fig. 1 Schematic diagram representing the workflow used in this work. The redox dataset generation process involved collecting iron complexes from the tmQM dataset through a series of filters, geometry optimization in a micro-solvated environment using tight-binding DFT, and standard DFT to calculate the redox potentials. The redox dataset was then used to train multiple GNN models. The graph data required to train the GNN models was generated from desolvated iron complexes that were obtained after removing the explicit solvent water molecules from the solvated and optimized iron complexes.

described later below. Following all the filtering steps, a total of 2267 iron complexes were collected and used to calculate the redox potential of Fe(II)/Fe(III) pairs. The calculation of the redox potential of the iron complexes led to the curation of a dataset that will be referred to in this text as the Redox<sub>2k</sub> dataset.

### Quantum chemical calculations

Each complex was evaluated with a hybrid aqueous solvation effect approach, where solvent effects are treated with inclusion of both explicit water molecules and an implicit continuum model. To do so, we first micro-solvated by adding 30 water molecules using the STOCASTIC algorithm in the Solvator module of ORCA 6.0 software.<sup>51,52</sup> The number of water molecules was chosen arbitrarily based on previous studies.<sup>27</sup> Next, each micro-solvated complex geometry was optimized using the GFN2-xTB semiempirical method,<sup>53</sup> with singlet spin multiplicity and the total charge reported in the tmQM dataset, and with the ALPB implicit solvation model.<sup>54</sup> We performed a vibrational frequency (Hessian) calculation to confirm that each structure corresponded to a local minimum.

To compute the redox potential, we evaluated single point energies (SPEs) of the micro-solvated Fe(III) complexes and their reduced form, Fe(II), using the TPSSH/def2-TZVP level of theory. To account for solvent effects, we used the SMD model with water dielectric constant. This level of theory was suitable for computing approximately 2000 complexes with our computational resources.

Our selection of this method was made after a comparison of the performance between B3LYP<sup>55</sup> and TPSSH<sup>56</sup> exchange-

correlation (xc) functionals, and basis sets def2-SVP, def2-TVZP, and def2-TZVPP, in the SI (Fig. S1). We found a linear correlation between both functionals, in particular with larger basis sets, this observation is consistent with previous reports comparing xc hybrid functionals.<sup>57,58</sup> Particularly, Jensen<sup>59</sup> reported that TPSSH has a good performance describing bond energies of transition metals and bioinorganic molecules. Our results show that values computed with def2-TVZP basis have a trend closer to a larger basis set, def2-TZVPP, than to the smaller def2-SVP. Semi empirical dispersion energies were not added, since we performed differences between single point energy calculations the effects of dispersion energies are canceled. Full relaxation of solvated complexes at DFT level had a prohibitive cost for the scope of this work. A systematic comparison among DFT xc functionals, basis sets and solvent models have been extensively explored in a previous studies.<sup>5,27,60</sup>

We estimated the Gibbs free energy of redox,  $G_{\text{red}}$ , as the difference of SPEs from electronic structure calculations, using the following equation:

$$\Delta G_{\text{red}} \approx \text{SPE}_{\text{III}} - \text{SPE}_{\text{II}} \quad (2)$$

The redox potential,  $E_{\text{red}}$ , was calculated using the following equation, adapted from the work of Harb<sup>27</sup>

$$E_{\text{red}}(\text{V}) = -\frac{\Delta G_{\text{red}}}{nF} - 4.3 \quad (3)$$

where  $n = 1$  (the number of electrons) and  $F$  is the Faraday constant. The factor  $-4.3$  V was used to reference the potential



**Table 1** Comparison of redox potential (V), measured with the standard hydrogen electrode as the reference, of selected Fe-ligand (L) complexes from previous results. Absolute errors (V) are shown in parenthesis

Entry	Ligand	This work		Harb <sup>27</sup>		Exp.
1	Maltolate	-0.30	(0.07)	-0.57	(0.34)	-0.23 (ref. 58)
2	Deferiprone	-0.31	(0.27)	-0.76	(0.18)	-0.58 (ref. 58)
3	Kojate	-0.33	(0.20)	-0.27	(0.14)	-0.1 (ref. 58)
4	Catecholate	-0.93	(0.10)	-0.99	(0.16)	-0.83 (ref. 58)
5	Salicylate	-0.67	(0.01)	-0.84	(0.18)	-0.66 (ref. 58)
6	Fe(CN) <sub>6</sub>	1.29	(0.93)	0.30	(0.07)	0.37 (ref. 62)
7	BIS-TRIS	-0.52	(0.39)			-0.91 (ref. 65)
8	DIPSO	-0.74	(0.10)			-0.84 (ref. 66)
9	TEA	-0.44	(0.41)			-0.85 (ref. 6)
10	TIPA	-0.63	(0.25)			-0.88 (ref. 67)
11	Phen	0.64	(0.48)			1.12 (ref. 5)
12	Terpy	0.67	(0.46)			1.13 (ref. 5)

to the standard hydrogen electrode (SHE). In the following sections, redox potentials from our results are presented in volts (V) instead of V vs. SHE unit for simplicity. Although our method to evaluate the redox potential differ from evaluations using Born-Haber cycles,<sup>61</sup> the approach showed here is suitable for high-throughput estimations of thousands of materials. Our approximation, suggesting standard conditions and fast charge transfer to use same geometry for reduced and oxidized species, is accurate enough to screen the iron complexes space.

In Table 1, we show a comparison of the redox potentials evaluated using our methodology, previous reports by Harb<sup>27</sup>, and experimentally measured values. In most cases, our methodology provided redox values comparable to experimental values. The largest error was observed for the pair Fe(CN)<sub>6</sub><sup>-4/-3</sup> with a calculated redox potential of +1.29 V compared with the experimental value of +0.37 V.<sup>62,63</sup> Nevertheless, our calculated redox potential value was positive, whereas previous studies reported negative values of the redox potential, for example -0.56 V,<sup>58</sup> -0.34 V,<sup>64</sup> and -0.33 V.<sup>60</sup> Although the discrepancies of redox estimations may be larger for other complexes that are not included in Table 1, the current evaluations suggest that our dataset could be a starting point for a further more in detail analysis, both theoretical and experimental.

To perform all these atomistic simulations, we used a workflow manager based on Fireworks,<sup>68</sup> using Jobflow to create Firetasks, and the atomic simulation environment (ASE)<sup>69</sup> to drive calls of ORCA and xTB binaries. We used two MongoDB databases, one for managing the jobs, and a second for storing the data. All our computations used the Improv (CPU AMD Ryzen Rome) cluster at Argonne Laboratory Computing Resource Center.

### GNN training datasets

Following the redox potential calculations, three distinct datasets were constructed for training the GNN models. All training datasets inherited our calculated redox potentials as the target property from the full Redox<sub>2k</sub> dataset. The three datasets

differed in both the types of geometries included and the total number of structures or data points in the datasets.

The first training dataset was generated by collecting desolvated iron complexes obtained after removing all explicit solvent water molecules from the optimized, microsolvated structures used in the redox potential calculations. All resulting structures were converted from Cartesian coordinates into RDKit Mol objects using a modified version of the xyz2mol\_tm method, as described in the SI.<sup>50</sup> Several filtering and pre-processing steps were required to enable reliable structure conversion, which resulted in the exclusion of some metal complexes from the original Redox<sub>2k</sub> dataset and yielded a curated set of 1450 iron complexes. A detailed description of all filtering and processing steps applied during the collection of desolvated structures is provided in the SI. This dataset, consisting of 1450 desolvated iron complex geometries, is referred to, in this text, as the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset.

The second dataset was constructed using the same 1450 iron complexes as the first dataset, but with geometries obtained directly from the tmQM dataset instead of desolvated geometries. This dataset is referred to, in this text, as the GNN-Redox<sub>1450</sub><sup>tmQM-geo</sup> dataset. Finally, the third dataset was created by collecting iron complex geometries directly from the tmQM dataset. No filtering or preprocessing steps were required for the third dataset since the tmQM geometries were directly used. Thus, all iron complexes in the full Redox<sub>2k</sub> dataset could be utilized. All collected iron complexes were converted from Cartesian coordinates into RDKit Mol objects using our modified xyz2mol\_tm method to construct the dataset. This third dataset consisting of ~2000 iron complexes is referred to, in this text, as the GNN-Redox<sub>2k</sub><sup>tmQM-geo</sup> dataset.

### Graph data generation

All GNN-Redox training datasets contained RDKit mol objects to represent the iron complexes in the GNN model training process. However, some graph convolution model architectures cannot work directly with such mol objects or 3D coordinates, and instead, require graph data with handcrafted features. So, the RDKit mol objects of the iron complexes in our GNN-Redox training datasets were converted to graph representations with comprehensive node, edge, and graph-level features. A complete list of all features is provided in Table 2. Notably, we used a condensed version of the full 2048 bit Morgan fingerprint<sup>70</sup> with a bond diameter of six bonds (*i.e.*, ECFP6) as a graph feature. ECFP6 has been previously used in various molecular property prediction tasks<sup>71-73</sup> and was found to yield the best overall model performance in this study (data not shown) compared to fingerprints generated with larger diameters. The condensed fingerprint comprised of the most important 64 fingerprint bits, as determined by a Random Forest Regressor model that was trained to predict the redox potential using the full Morgan fingerprint as input features. Utility of such fingerprint condensation has been demonstrated before to detect structurally diverse active compounds.<sup>74</sup> Here, condensation of the Morgan fingerprint aimed to enhance model



Table 2 Description of node, edge, and graph features for the GNN Models

Item	Feature name	Description	Type	Size
Node features	Atomic number	Atomic number scaled by Uranium's atomic number: (atomic_number - 1)/91	Numerical	1
	Symbol	Chemical symbol of the element	One-hot vector	92
	Valence	Number of maximum valence electrons	One-hot vector	8
	Formal charge	Formal electrical charge of the atom	Numerical	1
	Radical electrons	Number of radical electrons	Numerical	1
	Hybridization	Hybridization as s, sp, sp <sup>2</sup> , sp <sup>3</sup> , sp <sup>3</sup> d, sp <sup>3</sup> d <sup>2</sup> , or unspecified	One-hot vector	7
	Aromaticity	Indicates if an atom is in an aromatic ring (1 or 0)	Numerical	1
	Hydrogens	Number of connected hydrogens, encoded as 0, 1, 2, 3, or 4	One-hot vector	5
	Atomic mass	Scaled atomic mass: (mass - 1.008)/237.021	Numerical	1
	Vdw radius	Scaled Van der Waals radius: (R <sub>V<sub>DW</sub></sub> - 1.2)/1.35	Numerical	1
Edge features	Covalent radius	Scaled covalent radius: (R <sub>covalent</sub> - 0.23)/1.71	Numerical	1
	Bond type	Bond type as single, double, triple, or dative	One-hot vector	4
	Conjugated bond	Indicates if the bond is conjugated (1 or 0)	Numerical	1
	Ring	Indicates if the bond is in a ring (1 or 0)	Numerical	1
Graph features	Stereo	Stereochemistry: StereoNone, StereoAny, StereoZ, StereoE	One-hot vector	4
	Morgan fingerprint	Condensed version of the Morgan fingerprint	Numerical	64

performance by highlighting key chemical features relevant for redox behavior of the complexes.

### Graph neural network models

We trained four GNN architectures to predict the redox potentials of iron complexes: graph convolutional network (GCN),<sup>75</sup> graph attention network (GAT),<sup>76</sup> DimeNet++,<sup>77</sup> and SchNet.<sup>78,79</sup> GCN and GAT are topology-centric models that learn from abstract graph connectivity and node features. The models differ, however, in their approach to neighborhood aggregation. In the message aggregation step, a GCN assigns fixed, structure-dependent weights to neighboring nodes, typically based on node degrees, without learning their relative importance. In contrast, a GAT aims to extend GCN by employing a learnable attention mechanism that dynamically assigns different weights to each neighbor based on their features. Both GCN and GAT models have been widely and successfully used for various molecular property prediction tasks, including the prediction of redox potential of organic molecules.<sup>47,76,80</sup>

### RexGCN

In this work, we modified the original GCN<sup>75</sup> implementation to incorporate edge features. Addition of edge features in message construction has been demonstrated previously in many other GNN models for molecular property prediction tasks.<sup>81-83</sup> The standard GCN formulation only considers node features during message passing, which can be defined as:

$$m_{ij}^{(l)} = \hat{a}_{ij} \mathbf{x}_j^{(l)} \Theta^{(l)}, \hat{\mathbf{A}} = \tilde{\mathbf{D}}^{-1/2} \tilde{\mathbf{A}} \tilde{\mathbf{D}}^{-1/2}, \tilde{\mathbf{A}} = \mathbf{A} + \mathbf{I} \quad (4)$$

where  $\mathbf{x}_j^{(l)} \in \mathbb{R}^{d^{(l)}}$  denotes the feature vector of node  $j$  at layer  $l$ ,  $\Theta^{(l)} \in \mathbb{R}^{d^{(l)} \times d^{(l+1)}}$  is the learnable transformation matrix,  $\tilde{\mathbf{A}}$  is the adjacency matrix with self-loops added,  $\tilde{\mathbf{D}}$  is its degree matrix, and  $\hat{a}_{ij} = \frac{\tilde{a}_{ij}}{\sqrt{\tilde{d}_i \tilde{d}_j}}$  are the normalized edge weights. The GCN we used in this work, which will be referred to as the RexGCN,

extends the original formulation by incorporating edge features  $\mathbf{e}_{ij}$  into the message construction:

$$m_{ij}^{(l)} = \hat{a}_{ij} [\mathbf{x}_j^{(l)} \Theta^{(l)} \parallel \mathbf{e}_{ij}] \quad (5)$$

where  $\parallel$  denotes concatenation and  $\mathbf{e}_{ij} \in \mathbb{R}^{d_e}$  represents the edge features between nodes  $i$  and  $j$ . Since the concatenation increases the message dimensionality from  $d^{(l+1)}$  to  $d^{(l+1)} + d_e$ , a learnable update function was introduced here to project the aggregated messages back to the desired output dimension:

$$\mathbf{h}_i^{(l+1)} = \mathbf{W}_{\text{update}}^{(l)} \left( \sum_{j \in \mathcal{N}(i) \cup \{i\}} m_{ij}^{(l)} \right) + \mathbf{b}^{(l)} \quad (6)$$

where,  $\mathbf{h}_i^{(l+1)} \in \mathbb{R}^{d^{(l+1)}}$  denotes the updated feature (hidden representation) of node  $i$  at the output of layer  $l$ ,  $\mathbf{W}_{\text{update}}^{(l)} \in \mathbb{R}^{(d^{(l+1)} + d_e) \times d^{(l+1)}}$  is the learnable update projection matrix, and  $\mathbf{b}^{(l)} \in \mathbb{R}^{d_{\text{out}}^{(l)}}$  is the bias term. The complete RexGCN layer can be expressed in matrix form as:

$$\mathbf{h}_i^{(l+1)} = \sigma \left( \mathbf{W}_{\text{update}}^{(l)} \sum_{j \in \mathcal{N}(i) \cup \{i\}} \hat{a}_{ij} [\mathbf{x}_j^{(l)} \Theta^{(l)} \parallel \mathbf{e}_{ij}] + \mathbf{b}^{(l)} \right). \quad (7)$$

where  $\mathcal{N}(i)$  is the set of neighbors of node  $i$  and  $\sigma(\cdot)$  is a nonlinear activation function (e.g., ReLU). These modifications aim to create a better description of the local chemical environments and, subsequently, enhance the GCN's performance for modeling redox potential of metal complexes. To confirm that incorporating edge features improves GCN performance, we trained a baseline GCN model without edge features and compared its results with our modified RexGCN implementation. The comparison results are presented in the Results and Discussion section below.

### RexGAT

We constructed the GAT model used in this work based on the MolGAT model, which was previously used to predict redox potential of small organic molecules.<sup>48</sup> We implemented several



modifications to enhance model robustness and performance. The primary changes include: (1) explicit self-loop handling with appropriate edge feature management, (2) implementation of residual connections with adaptive input projection, and (3) refined message passing that separates attention computation from message content.

The GAT layer used here, which will be referred to as the RexGAT, begins with a linear transformation of input node features:

$$\mathbf{h}_i^{(l),h} = \mathbf{x}_i^{(l-1)} \mathbf{W}^{(l),h}, \quad h = 1, \dots, H \quad (8)$$

where  $\mathbf{x}_i^{(l-1)} \in \mathbb{R}^{d_{in}^{(l)}}$  represents the input feature vector from the previous layer,  $\mathbf{W}^{(l),h} \in \mathbb{R}^{d_{in}^{(l)} \times d_{out}^{(l)}}$  is the learnable weight matrix for head  $h$ , and  $\mathbf{h}_i^{(l),h} \in \mathbb{R}^{d_{out}^{(l)}}$  is the head-specific transformed feature. Similar to the MolGAT model, the attention mechanism in this work incorporates both node and edge features to compute attention coefficients:

$$e_{ij}^{(l),h} = \text{LeakyReLU}(\mathbf{a}^{(l),h \top} [\mathbf{h}_i^{(l),h} \parallel \mathbf{h}_j^{(l),h} \parallel \mathbf{e}_{ij}]) \quad (9)$$

$$\alpha_{ij}^{(l),h} = \text{softmax}_j(e_{ij}^{(l),h}) = \frac{\exp(e_{ij}^{(l),h})}{\sum_{k \in \mathcal{N}(i) \cup \{i\}} \exp(e_{ik}^{(l),h})} \quad (10)$$

where  $\mathbf{h}_i^{(l),h} \in \mathbb{R}^{d_{out}^{(l)}}$  is the  $h$ -th head vector for node  $i$ ,  $\mathbf{e}_{ij} \in \mathbb{R}^{d_e}$  is the edge feature between nodes  $i$  and  $j$ ,  $\parallel$  denotes concatenation, and  $\mathbf{a}^{(l),h} \in \mathbb{R}^{2d_{out}^{(l)} + d_e}$  is the learnable attention vector for head  $h$ .

We then constructed messages based on the following equation:

$$\mathbf{m}_{ij}^{(l),h} = \alpha_{ij}^{(l),h} \mathbf{h}_j^{(l),h} \quad (11)$$

Here, the edge features do not directly contribute to the message content, unlike the original MolGAT where the edge features influence both attention weights and messages directly. The message construction approach used here ensures that edge features modulate the importance of neighboring nodes through the attention coefficient  $\alpha_{ij}^{(l),h}$  while maintaining clean node feature propagation. Next, the per-head aggregated message  $\mathbf{a}_i^{(l),h} \in \mathbb{R}^{d_{out}^{(l)}}$  is obtained by summing messages over neighbors (including self-loop):

$$\mathbf{a}_i^{(l),h} = \sum_{j \in \mathcal{N}(i) \cup \{i\}} \mathbf{m}_{ij}^{(l),h} \quad (12)$$

The final layer output incorporates multi-head averaging and a learnable update transformation:

$$\mathbf{x}_i^{(l)} = \mathbf{W}_{\text{update}}^{(l)} \left( \frac{1}{H} \sum_{h=1}^H \mathbf{a}_i^{(l),h} \right) + \mathbf{b}^{(l)} \quad (13)$$

where  $\mathbf{W}_{\text{update}}^{(l)} \in \mathbb{R}^{d_{out}^{(l)} \times d_{out}^{(l)}}$  is the learnable update matrix  $\mathbf{b}^{(l)} \in \mathbb{R}^{d_{out}^{(l)}}$  is the bias term. To improve training stability and gradient flow, residual connections are incorporated when feasible:

$$\mathbf{x}_i^{(l)} = \mathbf{x}_i^{(l)} + \text{Proj}(\mathbf{x}_i^{(l-1)}) \quad (14)$$

where  $\text{Proj}(\cdot)$  is an identity mapping when input and output dimensions match, or a learned linear projection otherwise.

The attention mechanism in our RexGAT implementation remains fundamentally the same as the original MolGAT formulation, ensuring that the model can effectively capture the importance of different molecular bonds and neighboring atoms. However, the modified architecture, with its explicit self-loop handling, residual connections, and refined message passing, provides a better model implementation, enhanced training stability, and improved capacity for capturing complex atomic bonding environments of coordination complexes.

### DimeNet++ and SchNet

Distinct from the topology-based GCN or GAT models, DimeNet++ and SchNet are geometry-centric models that incorporate 3D spatial information into the graph representation. SchNet uses continuous interatomic distances in its message-passing framework, while DimeNet++ further enriches this by incorporating both distances and angular relationships using radial and spherical basis functions. As a result, DimeNet++ and SchNet produce rotationally invariant representations, making them effective for tasks such as molecular property prediction where 3D geometry is critical. Although their usability for redox prediction has not been explored previously, both DimeNet++ and SchNet have been used for various molecular property prediction tasks.<sup>77,79,84–86</sup> We created the DimeNet++ and SchNet models for redox potentials based on the same architecture and hyper-parameters used to model the QM9 dataset.<sup>77,79</sup> Detailed model hyper-parameters are provided in the SI.

### General training parameters

The GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset containing desolvated geometries of 1450 complexes was used to train and evaluate all models. Average model performance was assessed by conducting five-fold cross validation (CV) for RexGCN, RexGAT, and DimeNet++ models, while the SchNet model was assessed using a three-fold CV. Finally, one representative model was trained from each model type with early stopping based on validation set performance for final evaluation. Before training any model, the dataset was divided into training, validation, and test sets using a 70 : 15 : 15 split. To avoid any data leakage, the random forest regressor used to condense the fingerprints was trained only using the train split in both the CV analysis and final model training. Training epoch varied depending on the model architecture. RexGCN, RexGAT, DimeNet++, and SchNet models were trained for a maximum of 200 epochs, 200 epochs, 400 epochs, and 2000 epochs, respectively, with a maximum batch size of 128 to balance between computational cost and model performance. For early stopping, patience was set to 50 epochs for RexGCN, RexGAT, and DimeNet++ models, while patience was set to 100 for the SchNet model.



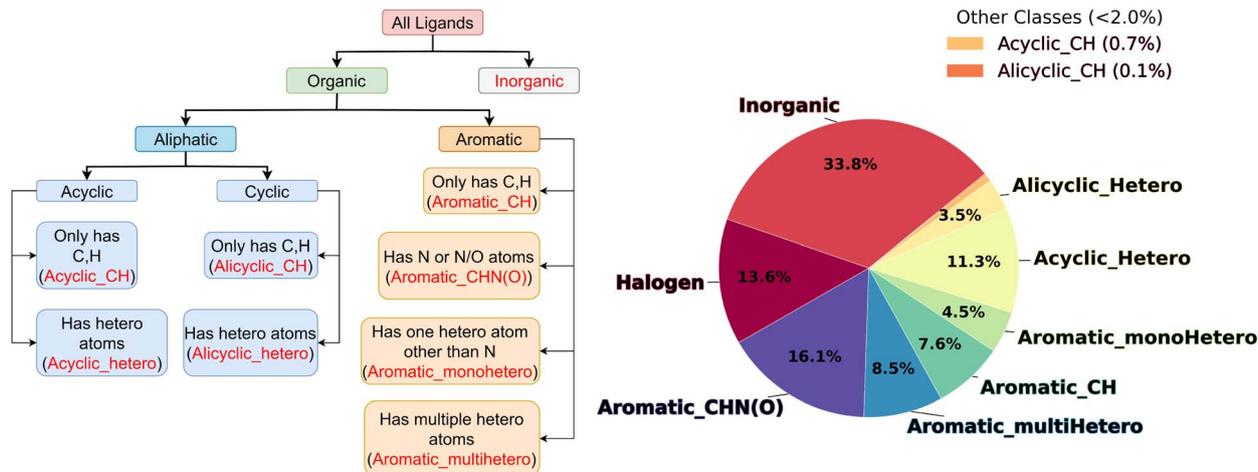


Fig. 2 Classification tree for the ligands (on the left) and frequency of the ligand classes in the Redox<sub>2k</sub> dataset (on the right). In the classification tree, the class names are identified with red texts.

## Results and discussion

### Redox<sub>2k</sub> dataset analysis

To analyze the diversity of the chemical space of the selected 2267 iron complexes in the Redox<sub>2k</sub> dataset, we systematically categorized the ligands into distinct classes. First, the ligands were classified into organic, *i.e.*, contains C–C or C–H bonds, or inorganic ligands. Next, the organic ligands were categorized as aromatic or aliphatic groups which were then further subclassified based on the presence of heteroatoms. The classification scheme and the distribution of ligands across these categories are illustrated through a classification tree and a pie chart in Fig. 2 for the Redox<sub>2k</sub> dataset. Examples of ligands from each class are presented in Fig. 3 and additional examples are shown in Fig. S2.

The iron complexes in the Redox<sub>2k</sub> dataset contained a total of 7463 ligands, of which ~37% were aromatic compounds,

~34% were inorganic, ~14% were halogens, and the remaining ~15% were aliphatic groups (Fig. 2). Among the aromatic ligands, compounds containing N or both N and O heteroatoms were found to be the most frequent (~16% of total ligands) class. In case of aliphatic ligands, with ~11% of total ligands, acyclic compounds containing heteroatoms were the most common class of ligands. Ligand class distribution in the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset (Fig. S3), which was curated from the Redox<sub>2k</sub> dataset, followed the same trend as observed in the Redox<sub>2k</sub> dataset.

Analysis of iron coordination number in the Redox<sub>2k</sub> dataset showed octahedral coordination to be the most prevalent case (Fig. S4). Analysis of ligand denticity revealed that the Redox<sub>2k</sub> dataset predominantly contained mono- and bi-dentate ligands but up to hexa-dentate ligands were observed, excluding one outlier with a higher denticity (Fig. S4). The GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset also followed these same trends in iron

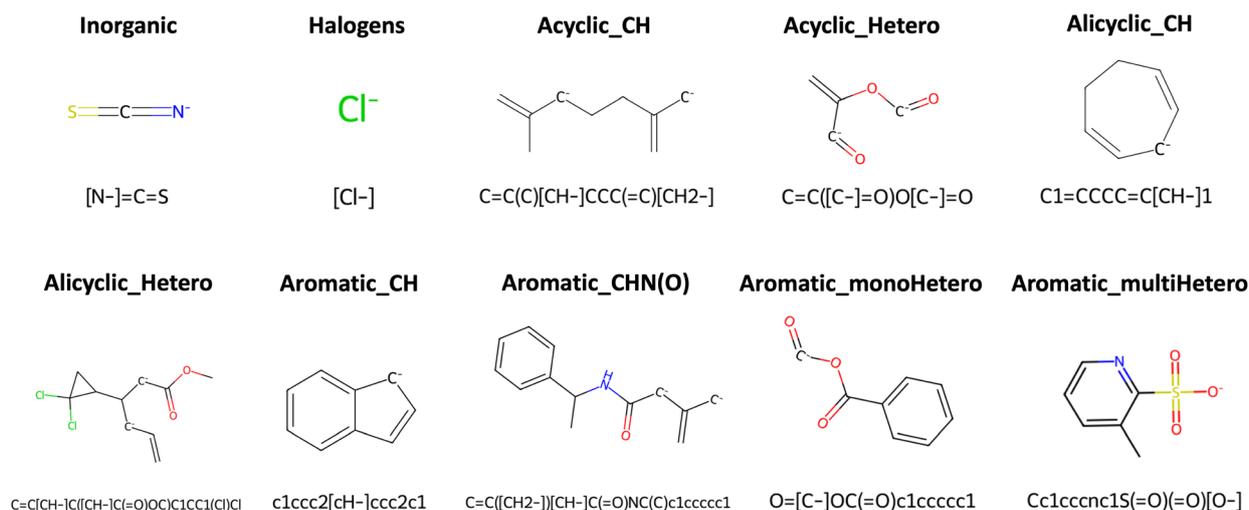


Fig. 3 Examples of representative ligands from each various ligand classes described in Fig. 2. The ligands are accompanied by their respective SMILES string.



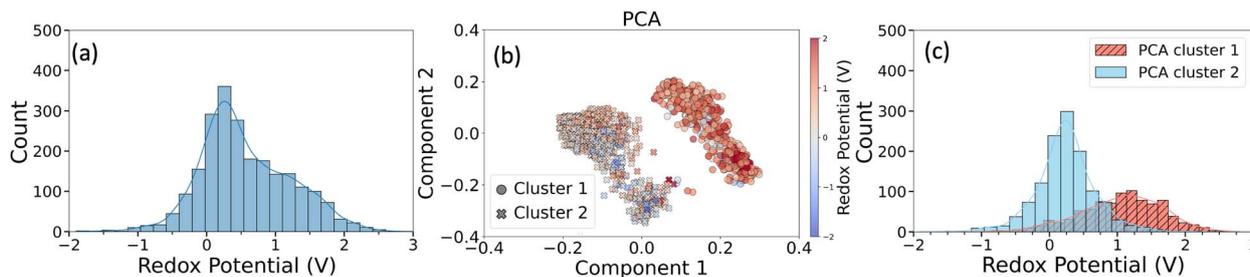


Fig. 4 (a) Distribution of the redox potentials in the Redox<sub>2k</sub> dataset. (b) The distribution of the iron complexes in the top two component space of the principal component analysis performed on the Morgan fingerprint of the complexes. The iron complexes diverged into two distinct clusters which were formally defined using *k*-means clustering algorithm, (c) distribution of the redox potentials of the two clusters detected in the PCA space in (b).

coordination number and ligand denticity as observed in the Redox<sub>2k</sub> dataset.

Fig. 4a shows the distribution of the computed redox potentials (vs. SHE) in the Redox<sub>2k</sub> dataset. The distribution ranged from  $-1.89$  V to  $7.47$  V with an average of  $0.56$  V. The distribution exhibited a slight right skew, with a tail extending towards more positive potentials. Six randomly selected iron complexes with redox potentials ranging from  $-1$  V to  $+1$  V are shown in Fig. S5 as representative structures.

The range of redox potentials observed in Fig. 4a spanned a broad spectrum ( $\sim -2$  V to  $\sim +2$  V) indicating the diverse set of electrochemical responses due to the presence of ligands-metal ion combinations, which could enable various redox-switch and flow-battery applications. More specifically, our computations showed that 377 of the iron complexes had a negative redox potential. Identification of these iron complexes with negative or lower redox potentials may provide new candidate anolytes for all-soluble, all-iron redox flow batteries. This is particularly important since current anolyte options are very limited with most notable options being iron-triethanolamine and iron-nitrilotri(methylphosphonic acid) complexes.<sup>63,87</sup>

To explore this structure-redox potential relationship, Morgan fingerprints were used to represent the chemical structure of the complexes, capturing detailed substructural information about the ligands. To reduce the dimensionality and highlight the most relevant features, a random forest regressor was first trained using the full 2048 bit Morgan fingerprints as input features to predict redox potentials. Then, 64 of the most important fingerprint bits as determined by feature importance scores were selected to generate a condensed version of the fingerprint which served as input for principal component analysis (PCA). Fig. 4b displays the PCA plot based on this condensed fingerprint, with the first two principal components capturing the largest variance in the data. This dimensionality reduction led to a clearer separation of complexes in the PCA space, offering better insight into how specific chemical features influence redox properties.

The iron complexes formed two well-separated clusters in the PCA space, as shown in Fig. 4b. To formally define these groupings, a *k*-means clustering algorithm with  $k = 2$  was applied. When the data points were colored according to their redox potentials, a clear distinction in redox behavior emerged

between the two clusters. This is further illustrated in Fig. 4c, which presents the redox potential distributions for each cluster. Cluster 1 exhibited redox potentials ranging from  $-1.76$  V to  $2.81$  V, with an average value of  $1.05$  V, standard deviation of  $0.61$  V, and median at  $1.11$  V. Redox potentials in Cluster 2, on the other hand, ranged from  $-1.90$  V to  $2.06$  V, excluding one outlier iron complex that had  $7.48$  V redox potential. Both the average and median redox potential in cluster 2 was the same,  $0.24$  V, and the standard deviation was  $0.44$  V. The significant difference in redox potential distributions between the two clusters highlights the influence of underlying ligand chemistry and suggests that redox behavior in iron complexes can be effectively tuned by targeted ligand selection.

The ligand classes, as defined in Fig. 2, were analyzed to gain deeper insight into the chemical characteristics associated with each PCA-derived cluster. Fig. 5a highlights the five most common ligand classes within each cluster. In Cluster 1, inorganic ligands were the most dominant, accounting for approximately 50% of all ligands in Cluster 1 (blue bars in Fig. 5a). Different aromatic groups and acyclic compounds containing heteroatoms were the other notable classes in the remaining 50% of ligands in Cluster 1. Remarkably, all complexes in Cluster 1 contained at least one inorganic ligand (red bars in Fig. 5a). Aromatic groups containing only C and H atoms were the second most common class appearing in  $\sim 55\%$  of complexes. All other classes individually appeared in no more than  $\sim 35\%$  of the complexes. These trends suggest that inorganic ligands are the defining chemical feature of Cluster 1.

In contrast, the Cluster 2 includes a more diverse ligand composition. Aromatic compounds containing N and/or O as heteroatoms (aromatic\_CHN(O) class), inorganic ligands, and halogens were observed with comparable overall frequencies (blue bars in Fig. 5a). These aromatic\_CHN(O) ligands were present in approximately 60% of the complexes (red bars in Fig. 5a), making them the most prevalent ligand class in Cluster 2. Notably, the aromatic\_CHN(O) ligand class was absent from the top five classes in Cluster 1, strongly indicating that the presence of aromatic\_CHN(O) ligands drives a complex toward Cluster 2. Additionally, the presence of halogen atoms, or combinations of inorganic ligands with halogens, also appears to favor assignment to Cluster 2.



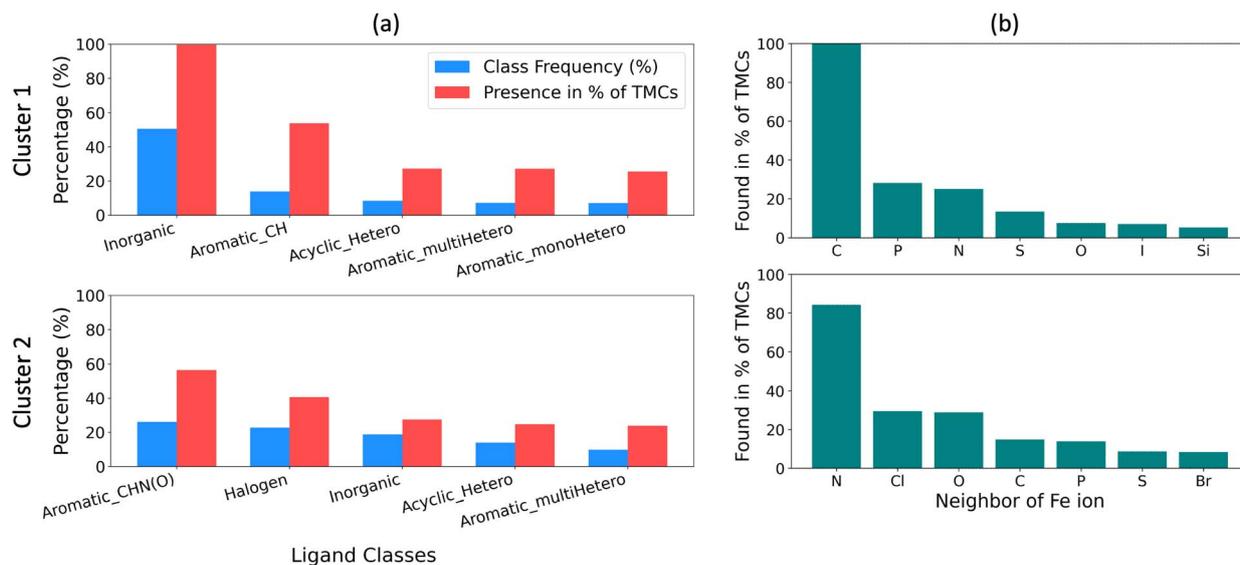


Fig. 5 Analysis of the two clusters observed in the PCA space. (a) Analysis of the ligand classes, as defined in Fig. 2, in the two clusters. The blue bars represent the frequency of ligand classes observed in all ligands of cluster 1 (top panel) and cluster 2 (bottom panel) transition metal complexes (TMCs). Only the five most frequent ligand classes are shown. The red bars represent the percentage of iron complexes that contain a ligand from a specific class. (b) Analysis of the direct neighbors of the iron ion in the two clusters. The bars represent the percentage of iron complexes that contain a specific element as a neighbor.

The chemical environment surrounding the metal ion plays a critical role in shaping the redox behavior of a metal complex. Therefore, analyzing the local coordination environment around the iron ion can offer further valuable insights into the distinct redox behaviors observed between the two PCA-derived clusters. Fig. 5b presents the distribution of neighboring chemical species directly bonded to the iron ion. In Cluster 1, every iron complex had at least one carbon atom coordinated to the iron center. Phosphorus and nitrogen were similarly frequent, appearing in approximately 30% of the complexes. In contrast, Cluster 2 was characterized by a markedly different coordination environment: nearly 80% of the complexes featured nitrogen as a neighbor to the iron ion, followed by chlorine and oxygen with each having presence in  $\sim 30\%$  of Cluster 2 complexes. Notably, carbon was observed as a neighbor in only  $\sim 20\%$  of Cluster 2 complexes. These results suggest that the presence of carbon and/or phosphorus atoms in the immediate environment of the iron ion is strongly associated with Cluster 1, while coordination with nitrogen, oxygen, and/or chlorine favors assignment to Cluster 2. This distinction in local chemical environments likely plays a significant role in driving the divergence in redox behavior between the two clusters. Further, the observed relation among redox potential of iron complexes, ligand class, and local environment can be utilized using GNN models for high-throughput screening purposes.

### Performance analysis of the GNN models

Four different GNN models (RexGCN, RexGAT, DimeNet++, and SchNet) were trained here using the GNN-Redox<sup>desolv-geo</sup><sub>1450</sub> dataset. Despite containing less structures, the training dataset remained a fair representation of the full Redox<sub>2k</sub>

dataset since the redox potential distribution (Fig. S6a), presence of the PCA clusters (Fig. S6b), and the chemical and redox properties of the clusters (Fig. S6c and S7) in the GNN-Redox<sup>desolv-geo</sup><sub>1450</sub> dataset followed the same trends as in the full Redox<sub>2k</sub> dataset. K-fold CV analysis was carried out for all the models to evaluate their average performance. Table 3 provides a summary of model performance across their respective CV folds. Among the four models, with the lowest root mean squared error (RMSE) of  $0.24 \pm 0.01$  V, the RexGCN model achieved the best average validation performance. The RMSE loss in the validation set across the CV folds are presented in Fig. S8 for all four models.

To further evaluate individual model performance, a representative instance was trained for each of the four GNN model types. We trained all models using an early stopping criterion based on validation RMSE loss. The training and validation loss curves for all representative models are shown in Fig. S9. Parity plots of redox potentials for all four models on the training, validation, and test sets are shown in Fig. 6. Among the four models, the RexGCN model demonstrated the highest predictive accuracy, achieving an  $R^2 = 0.84$  and RMSE = 0.26 V on the test set. The predictive accuracy of RexGCN surpasses that of a previously developed graph-based descriptor model, which

Table 3 Average performance of GNN models in K-fold CV

Model type	Folds	Best validation RMSE (standard deviation)	
RexGCN	5	0.24 V	(0.01 V)
RexGAT	5	0.27 V	(0.01 V)
DimeNet++	5	0.28 V	(0.01 V)
SchNet	3	0.48 V	(0.02 V)



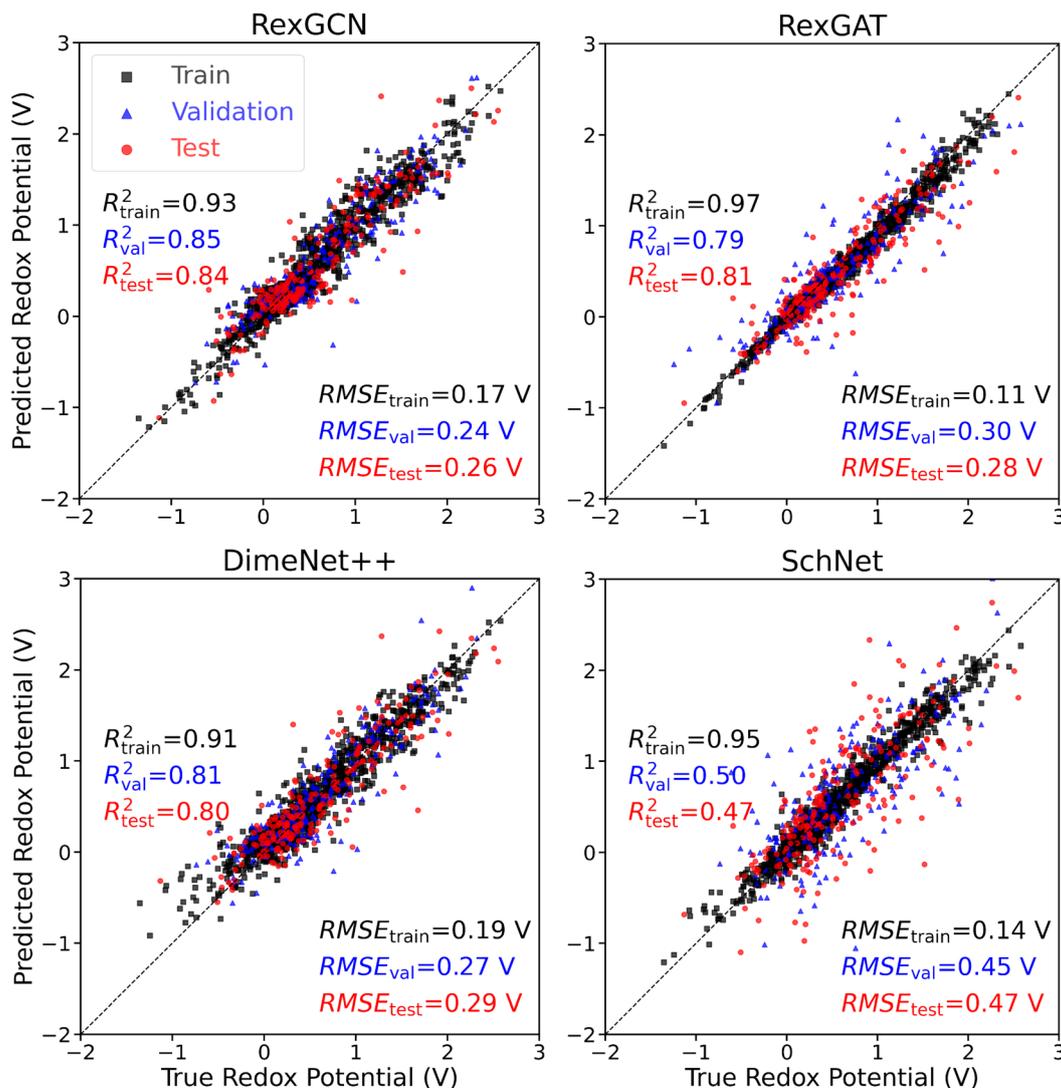


Fig. 6 Parity plots for the four representative models trained on the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset. True redox potential refers to our calculated redox potentials in the dataset. Train, validation, and test sets were created from a 70 : 15 : 15 split.

was trained to predict the redox potentials of iron complexes with nitrogen ligands and achieved an  $R^2$  score of 0.6 on the test set.<sup>26</sup> Note that the RexGCN model discussed here incorporates both node and edge features in message construction (eqn (5)). Our implementation improves upon the baseline GCN without edge features, as shown in Fig. S10. While the baseline achieved comparable average performance in the 5-fold CV analysis ( $0.27 \pm 0.01$  V), its predictive performance on the test set was poorer, with  $R^2 = 0.78$  and RMSE = 0.30 V, compared to our edge-aware RexGCN implementation.

Although RexGCN achieved the best overall performance, both RexGAT and DimeNet++ exhibited comparable predictive accuracy. As shown in Table 3, the average validation set RMSE loss of RexGAT and DimeNet++ in CV analysis differed by  $\leq 0.04$  V compared to RexGCN. In addition, representative RexGAT and DimeNet++ models achieved test set  $R^2$  scores of  $\sim 0.8$  (Fig. 6). SchNet, in contrast, exhibited the poorest performance among all evaluated models, yielding the highest

average validation RMSE of  $0.48 \pm 0.02$  V in CV analysis (Table 3). Consequently, the representative SchNet model showed the lowest predictive accuracy on the test set, characterized by the lowest  $R^2$  score and the highest RMSE loss (Fig. 6). Notably, SchNet also required the longest training time, converging after  $\sim 1750$  epochs. While its performance could potentially be improved through extensive hyperparameter optimization, such efforts would be computationally expensive, and increasing the number of model parameters would further slowdown the training process. Considering both computational cost and predictive accuracy, we found SchNet as a suboptimal choice for modeling redox potential of metal complexes.

The four models used here differ significantly in their architectural design and the way they process molecular information. GCN and GAT are both message-passing frameworks that require explicit construction of node-level, edge-level, and graph-level descriptors.<sup>75,76</sup> GAT aims to enhance the graph convolution of GCN by incorporating attention mechanisms to



weigh neighbor contributions dynamically.<sup>48,76</sup> In contrast, SchNet and DimeNet++ are continuous-filter convolution models that rely directly on atomic coordinates and atomic numbers to generate features dynamically during training, without the need for manual featurization.<sup>77,79</sup> While both models encode interatomic distances, DimeNet++ introduces greater expressivity by incorporating directional information and angular terms, and employing radial and spherical basis functions. This allows DimeNet++ to capture higher-order geometric relationships, offering a more nuanced representation of molecular structure than SchNet.

The comparative performance of the models reveals that the success of a GNN in predicting redox potentials depends less on the complexity of the learning algorithm and more on the quality and completeness of the chemical representation, whether learned or engineered. These observations underscore the critical role of feature representation in determining model performance. Models with either well-engineered graph features (as in GCN and GAT) or with architectures capable of constructing rich geometric features internally (as in DimeNet++) can learn redox behavior effectively. However, once meaningful structural and chemical information is captured either through manual featurization or geometric encoding, the choice of learning algorithm has a diminishing impact on performance. Instead, the limiting factors may be the overall quality of the dataset itself, determined by the size, accuracy, diversity, and chemical richness.

### Effect of TM geometry on model performance

The quality and chemical relevance of the training dataset have a substantial impact on model performance, as illustrated in Fig. 7 which compares the average validation RMSE obtained from 5-fold CV of the RexGCN model trained on three different datasets. These datasets are: (i) the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset, containing desolvated geometries of 1450 iron complexes; (ii) the GNN-Redox<sub>1450</sub><sup>tmQM-geo</sup> dataset, comprising of tmQM geometries of the same 1450 complexes; and (iii) the GNN-Redox<sub>2k</sub><sup>tmQM-geo</sup> dataset, which includes tmQM geometries for nearly all iron complexes in the Redox<sub>2k</sub> dataset.

As shown in Fig. 7, the RexGCN model achieved comparable average performance in validation set when trained on the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> (RMSE loss of  $0.24 \pm 0.01$  V) and GNN-Redox<sub>1450</sub><sup>tmQM-geo</sup> datasets (RMSE loss of  $0.25 \pm 0.02$  V). In contrast, with an RMSE loss of  $0.37 \pm 0.06$  V, a clear degradation in performance in validation set was observed when the model was trained on the larger GNN-Redox<sub>2k</sub><sup>tmQM-geo</sup> dataset.

The tmQM dataset consists of crystal geometries of iron complexes, which can differ substantially from solution-phase geometries in the presence of explicit solvent. In aqueous environments, water molecules may displace ligands and form solvation shells around the iron center, altering its coordination environment. However, the geometries in the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> and GNN-Redox<sub>1450</sub><sup>tmQM-geo</sup> datasets are highly similar, as complexes that exhibited significant deviations from their tmQM geometries upon solvation were excluded during the curation of the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset. As a result, these 1450 iron complexes largely retain their crystal-like coordination environments in solution, leading to comparable model performance across the two datasets. In contrast, the GNN-Redox<sub>2k</sub><sup>tmQM-geo</sup> dataset includes iron complexes that do not preserve their crystal geometries in solution and instead exhibit altered coordination environments around the iron center. The degradation in model performance when trained on the GNN-Redox<sub>2k</sub><sup>tmQM-geo</sup> dataset despite the larger dataset size suggests that the quality and consistency of structural inputs play a more critical role in model performance than simply increasing dataset size.

The importance of the immediate iron coordination environment is further highlighted by a feature attribution analysis of the RexGCN model on the test set using integrated gradients which is a gradient-based attribution method. The integrated gradients was employed to quantify the sensitivity of the predicted redox potential to perturbations in the input features of individual atoms, which in turn, elucidated the chemical rationale underlying the GNN predictions. The resulting importance scores reflect each atom's contribution to the final prediction. Atoms with high scores exert a strong influence on the predicted potential, whereas near-zero scores indicate negligible impact.

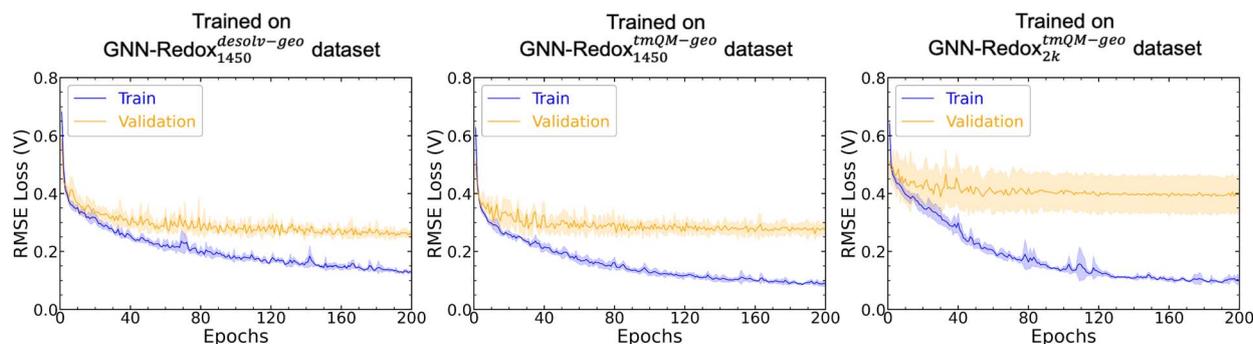


Fig. 7 5-fold CV performance comparison of RexGCN models trained on, (left panel) GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset, (middle panel) GNN-Redox<sub>1450</sub><sup>tmQM-geo</sup> dataset, and (right panel) GNN-Redox<sub>2k</sub><sup>tmQM-geo</sup> dataset. The solid lines represent the average RMSE loss of the 5 folds while the shaded region shows one standard deviation from the average.



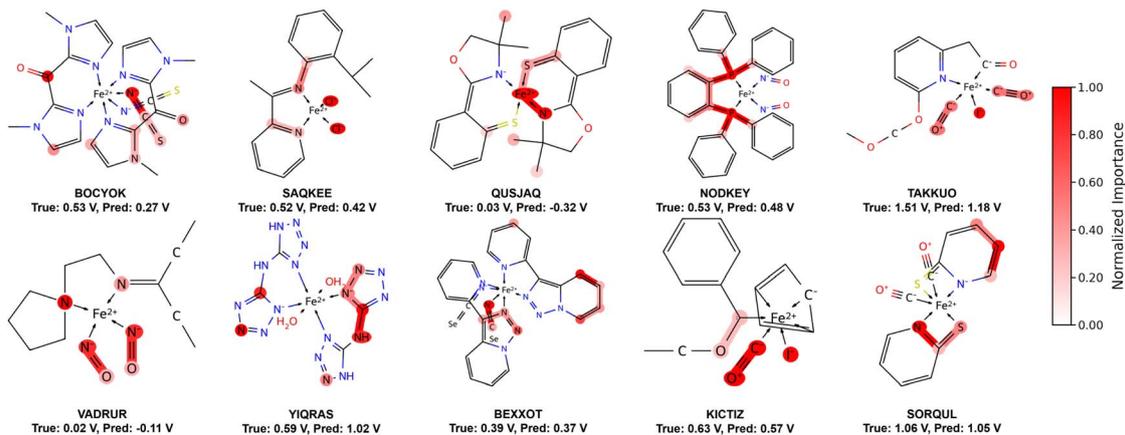


Fig. 8 Feature attribution analysis of the RexGCN model using Integrated Gradients. Atoms are colored according to their normalized importance scores for predicting redox potential, where a darker red color indicates a greater contribution to the model's output. For visual clarity, only the top 25% of atoms with the highest importance scores are highlighted. Each complex is annotated with its CSD reference code, the calculated (True) redox potential, and the model-predicted ( $P_{red}$ ) value.

As illustrated in Fig. 8, the analysis reveals a clear and consistent pattern where the highest importance scores are assigned to one or more atoms in the immediate coordination sphere of the central iron ion, *i.e.*, the directly bonded ligand donor atoms. More specifically, about 80% of the atoms with a high important score (top 25% importance) were within 3.5 Å distance of the central iron ion, evident from the distribution of iron-neighbor distances shown in Fig. S11. This strong localization of attribution indicates that the GNN has learned to prioritize the immediate environment of the redox-active metal center, in agreement with fundamental coordination chemistry principles, rather than relying on spurious correlations from distal atoms within the ligand framework.

## Conclusions

A scalable and efficient computational approach was developed here by utilizing tight-binding DFT and all-electron DFT methods to calculate the redox potential of transition metal complexes. To accurately capture the solvent effects in the redox calculations, our computational approach included both explicit water molecules and implicit continuum in the solvation structure. Redox potentials computed using our approach closely matched previous experimental values for several iron complexes, with an average absolute error of 0.13 V, demonstrating the reliability of our approach. This computational approach was then used to calculate the redox potential of 2267 iron complexes curated from the tmQM dataset, subsequently forming the Redox<sub>2k</sub> dataset.

Analysis of the Redox<sub>2k</sub> dataset revealed that ligand chemistry and the local coordination environment around the iron center are key determinants of the redox potential. Specifically, two distinct clusters with contrasting redox behavior were identified in the top two component space of the PCA analysis performed on the Morgan fingerprints of the iron complexes in the redox dataset. A closer examination of the clusters revealed that complexes with inorganic ligands tended to exhibit high

and positive redox potentials, while complexes with aromatic ligands containing nitrogen and/or oxygen as hetero atoms showed lower, possibly negative, redox potentials.

Next, a GNN framework was developed here to enable high-throughput screening of the vast chemical space of possible iron complexes. The GNN framework employed the xyz2mol method with new modifications to automatically generate fully featured graph data using only the 3D coordinates of the iron complex molecules. The GNN framework was then used to train four GNN models, namely RexGCN, RexGAT, DimeNet++, and SchNet, to learn and predict the redox potential of iron complexes from the GNN-Redox<sub>1450</sub><sup>desolv-geo</sup> dataset which was curated from the full Redox<sub>2k</sub> dataset through a series of filtering steps. The RexGCN model with engineered features provided the highest prediction accuracy, achieving an RMSE of  $0.24 \text{ V} \pm 0.01 \text{ V}$  and an  $R^2$  of 0.84 on the test set. The RexGAT and DimeNet++ models also achieved similar predictive performance compared to the RexGCN model whereas SchNet exhibited inferior performance.

Results here indicated that the quality of input features, whether engineered or learned, was more critical to model's performance than the complexity of the GNN architecture itself. Additionally, the predictive performance of the models dropped slightly when the full redox dataset was used for training by utilizing structures taken directly from the tmQM dataset. This observation highlighted the importance of the quality and consistency of the structures in the dataset rather than the dataset size to improve model performance. Finally, feature attribution analysis using integrated gradients revealed that atoms in the immediate coordination environment of the iron center were assigned greater importance by the GNN models in redox potential prediction. The methodology presented here provide a robust framework for exploring and predicting the redox properties of any transition metal complexes. This framework can accelerate the discovery of new materials for applications in redox flow batteries and catalysis.



## Author contributions

F. H. B curated the dataset for GNN model training; developed all the GNN models; prepared the figures and tables; prepared the codes and data repositories; and contributed with most of the manuscript writing. H. H. and R. S. A. calibrated the DFT protocol and validated results. A. V. M. conceived this study; performed all the simulations and data collection; outlined the manuscript. All the authors reviewed the manuscript and discussed the results.

## Conflicts of interest

The authors declare no competing interests.

## Data availability

Data and models are available at <https://doi.org/10.5281/zenodo.18419091>, and the code is available at <https://github.com/argonne-lcf/gnnredox>.

Supplementary information (SI) is available. See DOI: <https://doi.org/10.1039/d5dd00431d>.

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