Catalyst displacement assay: a supramolecular approach for the design of smart latent catalysts for pollutant monitoring and removal†

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Latent catalysts can be tuned to function smartly by assigning a sensing threshold using the displacement approach for targeted analytes. Three cyano-bridged bimetallic complexes were synthesized as “smart” latent catalysts through the supramolecular assembly of different metallic donors [FeII(CN)6]4−, [FeII(Bubpy)(CN)4]2−, and FeII(Bubpy)2(CN)2 with a metallic acceptor [CuII(dien)]2+. The investigation of both their thermodynamic and kinetic properties on binding with toxic pollutants provided insight into their smart off–on catalytic capabilities, enabling us to establish a threshold-controlled catalytic system for the degradation of pollutants such as cyanide and oxalate. With these smart latent catalysts, a new catalyst displacement assay (CDA) was demonstrated and applied in a real wastewater treatment process to degrade cyanide pollutants in both domestic (level I, untreated) and industrial wastewater samples collected in Hong Kong, China. The smart system was adjusted to be able to initiate the catalytic oxidation of cyanide at a threshold concentration of 20 µM (the World Health Organization’s suggested maximum allowable level for cyanide in wastewater) to the less harmful cyanate under ambient conditions.

Introduction

The development of “smart” catalysts for process monitoring and reaction control is highly important in modern chemistry. In particular, a smart catalyst that is able to execute a specific task under certain conditions would be very attractive. Different from traditional catalysts that initiate a reaction once the reactants are present, this type of catalyst would provide extra benefits for certain industrial processes such as contaminant removal during water treatment. Traditionally, chemists have focused on catalyst use to increase rates, yields, and/or the stereo-selectivity of reactions.1 However, it is possible that catalysts could be made “smarter” with the incorporation of extra control parameters, such as introducing an analyte-selective receptor which is also an inhibitor of the catalyst, so that the catalyst can only function when a specific analyte exists under certain conditions. This sort of smart catalyst is rarely found in the literature, although it could be rationally designed to associate with an inhibitor and be temporarily deactivated, but readily re-activated in the presence of a specific trigger or an initiator to start the targeted reaction under given conditions such as concentration, temperature, or pH. Recently, several catalytic systems demonstrated a portion of this “smart off–on” function for controlling targeted reactions, which led to very interesting reaction outcomes.2 A switch-based negative feedback loop, demonstrated with zinc(n) ion coordination-coupled deprotonation of a hydrazine, is a typical example of a smart control system, in which a particular threshold of zinc(n) causes the release of a certain concentration of protons to the environment to trigger a cascade of reactions.2i Based on these ideas, we conjectured that a further advancement of the “off–on” approach would be the introduction of a threshold-controlled function into the catalysts. The advancement demonstrated in this study overcomes the limitations of simple “catalyst-inhibitor” systems, arising from their simple off–on designs: i.e., they lack a mechanism to control the initiation threshold, so that the catalytic reaction will ensue in the presence of small amounts of initiator without having reached a specific initiator threshold.

The catalyst displacement assay (CDA) is a useful protocol for developing smart latent catalytic systems, particularly in chemical or waste treatment applications, because chemical toxicity depends on dosage. For example, cyanide (CN−) is well known for its high toxicity and has been identified as one of the
most serious threats toward the environment and human life. Nevertheless, it is widely used in gold mining, electroplating, and the production of synthetic fibers. Considering its industrial importance as well as its adverse effects, the World Health Organization has suggested maximum allowable levels of cyanide in wastewater, fresh water (class III), and drinking water of 0.5, 0.2, and 0.05 mg L⁻¹, respectively. The elimination of cyanide is therefore only required when its levels exceed these thresholds. The removal of cyanide in aqueous solution by oxidation has been well studied with oxidants in the presence of various catalysts such as TiO₂, Cu(II)/Cu(I) transition metal complexes and Fe(n)–Cu(n) bimetallic complexes. Nonetheless, these systems are not “smart” enough to begin functioning under a specific set of conditions. In recent years, the molecular design and synthesis of chemosensing systems specifically for cyanide recognition or determination have been reported. Some of these molecular systems are indicator displacement assays (IDA), which show sensing properties with good selectivity and signal changes upon binding with CN⁻ in solution. IDA involves specific analyte competition, which causes the displacement of the indicator from the receptor and gives off/on signals. The advantage of an IDA system is that the selectivity and sensitivity can be adjusted by tuning the thermodynamics of the ensemble. A number of IDA sensing systems are well-established for the determination of anions, neutral organic molecules, and other molecules; however, there are no examples of IDA systems showing smart properties that can effect a catalytic reaction under pre-set conditions.

On the basis of IDA systems, we attempted to establish a CDA system based on supramolecular donor–acceptor complexes which would feature a smart system that could selectively recognize a target analyte (a pollutant like CN⁻), after which a pre-set catalytic reaction would be automatically performed under given conditions. With the new CDA system design, the catalyst is controllably activated when the analyte level reaches a certain threshold to start a catalytic reaction. In this study, we demonstrate that a threshold, and the off/on control parameter, can be rationally adjusted by understanding the thermodynamic and kinetic properties of the cleavage processes in the donor–acceptor complexes. Both the thermodynamic and kinetic characteristics are crucial factors in deriving a CDA system that can generate a number of advantages for chemical processes—including the reduced or minimized use of reagents, costly catalysts, energy, and manpower—which cannot be provided by traditional chemical reactions or catalytic systems.

**Experimental section**

**Materials and general procedures**

4,4'-Di-tert-butyl-2,2'-bipyridine (tBubpy), diethylentetriamine (dien), K₂[Fe(CN)₆], potassium oxalate, and potassium cyanide were obtained from Aldrich. Complexes K₂[Fe((dien)Cl₂] (2), Fe((dien)Cl₂] (3), Cu((dien)Cl₂ (4) and K₄[FeII(CN)₆] (5) were prepared according to reported methods.

**Physical measurements and instrumentation**

Electrospray ionization mass spectrometry (ESI-MS) was performed on an AB SCIEX API 2000 LC/MS/MS system. Elemental analyses were conducted using a Vario EL CHN analyzer. Infrared spectra in the 500–4000 cm⁻¹ range using KBr pellets were recorded using a Perkin Elmer Frontier FTIR spectrometer, and UV-vis spectra were measured on a Cary 50 ultraviolet-visible spectrophotometer. Dissolved organic carbon was recorded using a Shimadzu TOC-L CSH High-Sensitivity Total organic Analyzer. Emission spectra were recorded using a Horiba FluoroMax-4 spectrophotofluorometer with a 5 nm slit width and a 0.5 s integration time. Broad-band UV irradiation was provided by a 500 W Hg(Xe) light source (Model 66485) with a digital power supply system (Model 69911) by Newport Inc.

**UV-vis spectroscopic titrations**

Analytical grade DMF used in the UV-vis spectroscopic titrations was purified by distillation. Titrations of K₄[FeII(CN)₆] (3.3 × 10⁻⁴ M) by CuII(dien)Cl₂ (0 to 2 × 10⁻⁵ M) were conducted in HEPES buffer at pH 7.4. While titrations of K₂[Fe((dien)Cl₂] (5 × 10⁻⁵ M) by CuII(dien)Cl₂ (0 to 2 × 10⁻⁴ M) and Fe((dien)Cl₂] (5 × 10⁻⁵ M) by CuII(dien)Cl₂ (0 to 1 × 10⁻⁴ M) were carried out in a mixture of aqueous HEPES buffer (1.50 mL, pH 7.4) with DMF(1.50 mL). UV-vis spectroscopic titrations of CuII(dien)Cl₂ (5 × 10⁻⁴ M) with oxalate (0 to 1 × 10⁻³ M) were carried out in aqueous phosphate buffer pH 4. All measurements were taken after standing for 12 h to ensure their equilibria were reached. Binding constants (log K) and formation energies (ΔG/kJ mol⁻¹) were analyzed according to the Benesi–Hildebrand equations from the UV-vis spectroscopic titrations.

**Kinetic measurements: cleavage of cyano-bridges between the FeII and CuII metal centers of 1–3 by cyanide**

The experimental procedure for the rate constant measurements was performed according to previous studies. Except for the reaction of complex 1, which was performed in pH 7.4 HEPES buffer, all reactions were performed in DMF/pH 7.4 HEPES buffer (1 : 1 v/v). By mixing a known amount of cyanide solution (2.5 × 10⁻¹ to 1.25 × 10⁻² M) with the test solutions containing 2.5 × 10⁻⁴ M complex at 25 °C, the changes in the absorbance due to the addition of cyanide were measured at different time intervals. All kinetics measurements were conducted under ambient conditions.

**Oxidation of cyanide to cyanate with complexes 1–3 as catalysts**

All experiments were conducted in a 40 mL boiling tube in the absence of light, and the test solutions (15.0 mL) were stirred during the experiments. The concentration of H₂O₂ was 6.53 mM, and the concentrations of complexes 1, 2, and 3 were 0.1, 0.1, and 0.2 mM, respectively. Different initial concentrations of cyanide (0 to 1000 μM) were used in the studies. The concentrations of CN⁻ and NCO⁻ in the test solutions were
measured at regular intervals using previously reported analytical methods.\textsuperscript{18n}

Photocatalytic degradation of oxalate to CO$_2$ by complex 1 as catalyst

All experiments were conducted in a 50.0 mL volumetric flask with a 500 W Hg(Xe) ultraviolet-visible lamp (Newport) irradiation source. The experimental setup was completely shielded from surrounding light. The distance between the lamp and test solution was about 25 cm. A water bath was placed between the lamp and test solution to absorb heat generated by the UV irradiation. Prior to irradiation, the pH of the test solution was adjusted to 1.5 with HCl/KCl buffer. Generally, during the photocatalytic experiments, a test solution (50.0 mL) containing complex 1 (6.25 $\times$ 10$^{-4}$ M) and H$_2$O$_2$ (0.4 M) was irradiated while the oxalate concentration was varied from 6.25 $\times$ 10$^{-4}$ to 3.125 $\times$ 10$^{-2}$ M. The dissolved organic carbon (DOC) in the system was determined at regular intervals to evaluate catalytic efficiency.\textsuperscript{1n}n

Synthesis and characterization

\[ \text{[Fe}^{II} \{\text{CN}\}_6 \}^2-\text{[Cu}^{II} \{\text{dien}\}^2 \}^2 \}^2 \] (1). The complex was synthesized by modification of the reported method.\textsuperscript{n}n A mixture of $K_2\text{[Fe}^{II} \{\text{CN}\}_6 \}^2$ (1.00 g, 2.37 mmol) and $\text{Cu}^{II} \{\text{dien}\}^{2}$ (0.56 g, 2.37 mmol) was stirred in a water/methanol mixture (1 : 1 v/v, 5 mL) at room temperature overnight. The brown precipitate obtained upon centrifugation was washed with DI water and then air-dried. Yield: 0.069 g (83.1%). IR (KBr): 3814, 1972, 1651, 1497, 1379, 1262, 1182 cm$^{-1}$.

\[ \text{[Fe}^{II} \{\text{Bubpy}\}_2 \{\text{CN}\}_4 \}^2-\text{[Cu}^{II} \{\text{dien}\}^2 \}^2 \}^2 \] (2). The complex was synthesized following the reported method.\textsuperscript{n}n A mixture of $K_2\text{[Fe}^{II} \{\text{Bubpy}\}_2 \{\text{CN}\}_4 \}^2$ (0.115 g, 0.2 mmol) and $\text{Cu}^{II} \{\text{dien}\}^{2}$ (0.05 g, 0.2 mmol) was stirred in a water/methanol mixture (1 : 1 v/v, 5 mL) at room temperature overnight. The brown precipitate obtained upon centrifugation was washed with DI water and acetone, and then air-dried. Yield: 0.069 g (83.1%). IR (KBr): 3205, 2059, 2084, 2103, 2114 cm$^{-1}$. ESI-MS (MeOH, +ve mode): $m/z$ 544.8 \{[Fe(CN)$_6$$]_2$\}$(C_6$H$_5$)$_2$$ \}$ (mass = 545.0 g mol$^{-1}$; charge = +1); elemental analysis calcd (%) for $C_{14}H_{26}N_{12}O$ (1): C 54.02; H 7.26; N 20.10.

\[ \text{[Fe}^{II} \{\text{Bubpy}\}_2 \{\text{CN}\}_4 \}^2-\text{[Cu}^{II} \{\text{dien}\}^2 \}^2 \}^2 \] (3). A mixture of $K_2\text{[Fe}^{II} \{\text{Bubpy}\}_2 \{\text{CN}\}_4 \}^2$ (0.10 g, 0.16 mmol) and $\text{Cu}^{II} \{\text{dien}\}^{2}$ (0.083 g, 0.16 mmol) was stirred in methanol (25 mL) at room temperature overnight. The reddish brown solid obtained after evaporation was washed with chloroform, DI water, and acetone, and then air-dried. Yield: 0.0745 g (61.1%). IR (KBr): 3205, 2068, 2082 cm$^{-1}$. ESI-MS (MeOH, +ve mode): $m/z$ 726.9 \{[Fe(Bubpy)$_2$(CN)$_4$$]_2$\}$(C_6$H$_5$)$_2$$ \}^2$ (mass = 1454.70 g mol$^{-1}$; charge = +2); elemental analysis calcd (%) for $C_{30}H_{40}N_{12}$ (2): C 43.27; H 6.05; N 20.19; found: C 43.22; H 6.10; N 20.10.

Results and discussion

Syntheses of bimetallic complexes 1–3

Metallic donors ($M_n$) coordinated with a bridging ligand such as the commonly used –CN, –NCS, or –NCO ligands are able to bond with a metallic acceptor ($M_A$) to form a desirable bimetallic complex of the form $M_n$–bridge–$M_A$. Through this supra-molecular assembly, bimetallic complexes can be designed to achieve desired functionalities which are not offered by the individual mono-metallic precursors ($M_n$ or $M_A$). We studied a series of bimetallic complexes containing a cyano-bridge, $M_n$–C≡N–$M_A$, where $M_D$ = Re$^{II}$, Fe$^{II}$, Ru$^{III}$, or Os$^{II}$; and $M_A$ = Ni$^{II}$, Cu$^{II}$, Pt$^{II}$, or Ln$^{III}$. Interestingly, the strength of the bridging bond between the metal centers is adjustable and depends on the combination of $M_D$ and $M_A$. Therefore, by regulating the two metal centers connected by the cyano-bridge, a bimetallic complex with specific functionality can be tailored. In this study, $M_D$, functioning as an inhibitor, was introduced into a catalyst $M_A$ to generate a CDA system with an “off-on” function in which a catalytic oxidation reaction could be triggered when a certain concentration of cyanide ions was present in the solution. The influence of the thermodynamic and kinetic properties of the cyano-bridges of the bimetallic complexes on the control of the off-on and threshold operation of the catalyst was then illustrated comprehensively.

Bimetallic complexes 1–3 shown in Scheme 1 were synthesized with the metallic acceptor Cu$^{II}$[dien]$^2$Cl$_2$ and the latent catalyst donors $K_2$[Fe(CN)$_6$]$^2$, $K_2$[Fe(Bubpy)[CN]$_4$]$^2$, and Fe(tBubpy)$_2$(C$_6$H$_5$)$_2$. All three bimetallic complexes were isolated as air-stable solids and were characterized by IR spectroscopy, electrospray ionization mass spectrometry (ESI-MS), and elementary CHN analysis. The three cyano–ferrous complexes, $K_2$[Fe(CN)$_6$]$^2$, $K_2$[Fe(Bubpy)[CN]$_4$]$^2$, and Fe(tBubpy)$_2$(C$_6$H$_5$)$_2$, were designed systematically as the MD species because they are able to participate in coordinating cyano bridging with other metal complexes to form bimetallic complexes, whereas [Cu$^{II}$[dien]$^2$Cl$_2$ was selected as the model acceptor $M_A$ because it can catalyze the oxidation of organic/inorganic pollutants such as oxalate and cyanide ions. We expected the strengths of the cyano-bridges of the three bimetallic complexes to be different due to the variation of the charge densities among the ferrous MD from $-4$, $-2$, to $0$. The cyano bridge activities of the bimetallic complexes 1–3 significantly impact their thermodynamics and kinetics, which are considered as the determinant for the threshold of a smart catalytic system.

The formation of cyano-bridges between the $M_D$ and $M_A$ units in the bimetallic complexes was confirmed using IR spectroscopy, as shown in Table 1. In general, all the $\nu_{C=O}$ stretching frequencies observed in 1–3 are red shifted with respect to their precursor cyano–ferrous complexes, and the degrees of the shift vary by 2, 29, and 60 cm$^{-1}$, respectively. These changes can be attributed to the electron density differences among the MD complexes, Fe$^{II}$[tBubpy]$_2$(CN)$_2$, Fe$^{II}$[tBubpy]$_2$(CN)$_4^2$, and Fe$^{II}$[tBubpy]$_2$(CN)$_6^3$. In the syntheses of the bimetallic complexes, 1 : 1 molar ratios of the Fe$^{II}$ and Cu$^{II}$ starting materials were used in all
Interestingly, however, the compositions of supramolecular structures 1–3 contained FeII/CuII ratios of 1:2, 1:2, and 2:1, as verified by ESI-MS, spectroscopic titration studies, and CHN elemental analysis. As shown in Fig. 1, 1–3 show peaks at m/z = 544.8, 380.1, and 726.9, representing [FeII(CN)6][CuII(dien)]2+, [FeII(tBubpy)(CN)4][CuII(dien)]22+, and [FeII(tBubpy)2(CN)2][CuII(dien)]22+, respectively. The elemental analysis results for 1–3 also matched the proposed structures. In addition, the binding isotherms (Fig. S1–S3, (ESI†) obtained by the titration of CuII(dien)Cl2 solution with K4[FeII(CN)6], K2[FeII(tBubpy)(CN)4], and FeII(tBubpy)2(CN)2, respectively, show that the bimetallic complexes 1–3 are formed in the solution in the above FeII/CuII ratios.15

**Thermodynamic and kinetic investigations of the CDA system**

In the design of a CDA, a catalyst should first be bound temporarily to an inhibitor. CuII complexes have been well studied as active catalysts for the oxidation of cyanide using H2O2 as the oxidant.17a,20 However, the catalytic activity is inhibited when the CuII-centers are bridged to FeII-centers with a ligand (C≡N) in the form of bimetallic complexes.18 The Cu-catalyst at this stage is de-activated as it is bridging with its metallic counterpart. When a competitive analyte, i.e., the reactant, is introduced into the system, it causes the displacement of the catalyst from the inhibitor, thus freeing it.18 At this stage, the Cu-catalyst is activated to execute its catalytic function under the given conditions. For the present CDA system ([L]x(CN)FeII–C≡N–CuII(dien)], Scheme 2), the

**Table 1** IR spectroscopic study of the cyano-stretching frequencies (νC≡N) of complexes 1–3 and their precursor complexes

<table>
<thead>
<tr>
<th>Entry</th>
<th>Complexes</th>
<th>νC≡N (cm⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>[FeII(CN)6][CuII(dien)]2</td>
<td>2039, 2047, 2103</td>
</tr>
<tr>
<td>2</td>
<td>[FeII(tBubpy)(CN)4][CuII(dien)]2</td>
<td>2059, 2084, 2103, 2114</td>
</tr>
<tr>
<td>3</td>
<td>[FeII(tBubpy)2(CN)2][CuII(dien)]2</td>
<td>2068, 2082</td>
</tr>
<tr>
<td>4</td>
<td>K4[FeII(CN)6]</td>
<td>2030, 2043</td>
</tr>
<tr>
<td>5</td>
<td>K2[FeII(tBubpy)(CN)4]</td>
<td>2053, 2067, 2085</td>
</tr>
<tr>
<td>6</td>
<td>FeII(tBubpy)2(CN)2</td>
<td>2080</td>
</tr>
</tbody>
</table>
study of the coordination with cyano-bridges between the Fe\textsuperscript{II} and Cu\textsuperscript{II} metal centers allows us to understand the thermodynamic and kinetic properties of these bimetallic systems. These investigations were conducted using cyanide as the model analyte.

To understand the thermodynamic properties of the CDA, the Gibbs free energy changes ($\Delta G^0$) between $M_\text{D}$ and $M_\lambda$ of 1-3 were found and calculated as $-30.4$, $-25.3$, and $-15.1$ kJ mol$^{-1}$, respectively (Table 2). The $\Delta G^0$ value of the adduct between Cu\textsuperscript{II}(dien)$^{2+}$ and CN$^-$ ($-37.9$ kJ mol$^{-1}$) is more stable relative to

Table 2  Binding constants (log $K$) and Gibbs free energy changes ($\Delta G^0$) for the complexation of cyanide anion and different Fe\textsuperscript{II} species to the [Cu\textsuperscript{II}(dien)]$^{2+}$ complex

<table>
<thead>
<tr>
<th>Acceptor</th>
<th>Donor</th>
<th>log $K$ $^a$</th>
<th>$\Delta G^0$/kJ mol$^{-1}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>[Cu\textsuperscript{II}(dien)Cl\textsubscript{2}]</td>
<td>Oxalate (HOOC-COO$^-$)</td>
<td>6.84</td>
</tr>
<tr>
<td>2</td>
<td><a href="ClO%5Ctextsubscript%7B4%7D">Cu\textsuperscript{II}(dien)(ClO\textsubscript{4})</a></td>
<td>CN$^-$</td>
<td>6.64$^b$</td>
</tr>
<tr>
<td>3</td>
<td>[Cu\textsuperscript{II}(dien)Cl\textsubscript{2}]</td>
<td>$K_\lambda[Fe\textsuperscript{II}(CN)_6]$</td>
<td>3.32</td>
</tr>
<tr>
<td>4</td>
<td>[Cu\textsuperscript{II}(dien)Cl\textsubscript{2}]</td>
<td>$K_\lambda[Fe\textsuperscript{II}(Bubpy)(CN)_4]$</td>
<td>4.44</td>
</tr>
<tr>
<td>5</td>
<td>[Cu\textsuperscript{II}(dien)Cl\textsubscript{2}]</td>
<td>Fe\textsuperscript{II}(Bubpy)$_2$(CN)$_3$</td>
<td>2.65</td>
</tr>
</tbody>
</table>

$^a$ Binding strengths were measured by UV spectroscopic titration and calculated using the Benesi–Hildebrand equations. Except entry 1, which was performed in aqueous phosphate buffer at pH 4, all reactions were performed in aqueous DMF (1 : 1 v/v, 1.50 mL aqueous HEPES buffer at pH 7.4 + 1.50 mL DMF) at 298 K. $^b$ log $K$ and $\Delta G^0$ data are cited from ref. 11b and 18a.
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Chemical Science

Threshold-controlled catalytic properties of the CDA system

For the investigation of the smart threshold-controlled properties of the CDA, the initial test solutions containing cyanide, H$_2$O$_2$, and 1, 2 or 3 in a molar ratio of 10 : 65 : 1 in buffered aqueous DMF solution were prepared for the analyses. The concentrations of CN$^-$ and NCO$^-$ in the test solutions were measured under ambient conditions. The results show that all the complexes could catalyze the oxidation of cyanide to cyanate quantitatively within 150 min with H$_2$O$_2$ as the oxidant. Control experiments in the absence of the complexes or H$_2$O$_2$ revealed the corresponding displaced Cu$^{II}$-complex [Cu$^{II}$ (dien)(CN)]$^+$ (m/z 192.0 [M$^+$]) was found and identified by ESI-MS spectroscopic analysis (Fig. S4a†).

We believe that the displacement of the Cu$^{II}$-complex (through bridge cleavage by the target analyte) from the bimetallic systems is not only controlled by the above-mentioned thermodynamic factors, but also that the different displacement kinetics of the systems could provide “smart” deactivation-activation functions which would enable us to establish a CDA system. To prove this concept, kinetic studies were conducted on the rates of cyano-bridge cleavage between the Fe$^{II}$ and Cu$^{II}$ metal centers in the bimetallic systems by cyanide. Fig. 3 shows that the rate constants for cyano-bridge cleavage in 1–3 by cyanide were determined to be 18.8, 32.0, and 58.3 M$^{-1}$ s$^{-1}$, respectively. The results indicate that complex 3 is the fastest to release the Cu-catalyst for the oxidation reaction. The rates of cyanate formation with 1–3 were found to increase with respect to their decreasing $\Delta G^0$ (1–3 = −30.4, −25.3 and −15.1 kJ mol$^{-1}$, respectively) and also in line with the rate constants for the cyano-bridge cleavage of 1–3 by cyanide ions (18.8, 32.0, and 58.3 M$^{-1}$ s$^{-1}$ Fig. 3). The rate constant of cyanate formation from the oxidation of cyanide with H$_2$O$_2$ as the oxidant under the catalytic action of Cu$^{II}$ was reported as 249.5 M$^{-1}$ s$^{-1}$ (Fig. S5†). The results reveal that the cleavage of the cyano-bridge between the metallic donor and acceptor to release the Cu$^{II}$ complex (Scheme 2; 1$^\text{st}$ step of the overall CDA process) is the rate determining step, whereas the 2$^\text{nd}$ step of the process, the catalytic oxidation of cyanide, is faster.

those of bimetallic systems 1–3. By comparing these four $\Delta G^0$ values, we predicted that when CN$^-$ is introduced into the bimetallic systems (1, 2, or 3), it would displace the Cu$^{II}$-dien$^{3+}$ unit from its ferrous partner by breaking the cyano-bridges, because the formed [CN–Cu$^{II}$-dien]$^+$ adduct is more stable than the precursor. Fig. 2a–c show evidence of breaking the cyano-bridges of complexes 1–3 by showing the restoration of the spectroscopic properties of [Fe$^{II}$(CN)$_6$]$^{3-}$, [Fe$^{II}$(Bubpy)(CN)$_4$]$^{2-}$, and Fe$^{II}$([Bubpy]$_2$(CN)$_2$) upon the addition of cyanide to 1–3, respectively. Most importantly, through cleaving the cyano-bridges upon the addition of cyanide to 1–3,

![Fig. 2](image_url)

**Fig. 2** UV-vis spectroscopic titration curves of (a) 1, (b) 2, and (c) 3 with CN$^-$. Addition of cyanide anions to 1–3 restored the characteristic spectroscopic properties of [Fe$^{II}$(CN)$_6$]$^{3-}$, [Fe$^{II}$([Bubpy](CN)$_4$]$^{2-}$, and Fe$^{II}$([Bubpy]$_2$(CN)$_2$), respectively. All of the titrations were examined in aqueous DMF (1 : 1 v/v, 1.50 mL aqueous HEPES buffer at pH 7.4 + 1.50 mL DMF) at 298 K with complex concentrations of $1 \times 10^{-4}$ M and CN$^-$ concentrations from 0 to $2.0 \times 10^{-6}$ M.

![Fig. 3](image_url)

**Fig. 3** Kinetic plots of the apparent association rate constant $k_{cat}$ (s$^{-1}$) versus cyanide concentration. The rate constant values were calculated from the slopes of the curves ($y = mx; 1 = 18.8$ M$^{-1}$ s$^{-1}$; $2 = 32.0$ M$^{-1}$ s$^{-1}$; and $3 = 58.3$ M$^{-1}$ s$^{-1}$).
that no cyanide was oxidized to cyanate throughout the period. On the other hand, in the reaction using Cu\textsuperscript{II}(dien)Cl\textsubscript{2} instead of the bimetallic complexes under the same conditions, cyanide was quantitatively oxidized to cyanate. This confirmed that the Cu\textsuperscript{II}-complex is the active catalyst in the oxidation reaction.

From the thermodynamics consideration, all of the complexes of 1–3 are able to release the inhibited Cu-catalyst in the presence of cyanide as the $\Delta G^0$ value (−37.9 kJ mol\textsuperscript{-1}) of Cu\textsuperscript{II}(dien)\textsuperscript{2+} and CN\textsuperscript{−} adduct is much lower than the formation of [(L)\textsubscript{3}(CN)Fe\textsuperscript{III}C≡N–Cu\textsuperscript{II}(dien)] (Table 2). However, Fig. 4b–d show the most interesting findings that the complexes 1–3 do not exhibit their catalytic properties, even though the reactions are considered thermodynamically feasible, if the cyanide concentration does not reach certain thresholds. However, by taking into account their kinetic properties, the initial cyanide concentration needs to reach thresholds of 0.2, 0.1, and 0.02 mM for 1, 2, and 3, respectively, to break their cyano-bridges and release [CN–Cu\textsuperscript{II}(dien)]\textsuperscript{2+} to catalyze cyanide oxidation. In addition, we observed obvious plateaus from Fig. 4b–d when the oxidation reaction of CN\textsuperscript{−} proceeds for a certain period of time so that the product of NCO\textsuperscript{−} accumulates in the reaction environment. The plateaus may indicate that the catalytic system is being slowed-down and/or an inhibition effect from the negative feedback loop of the product is generated.\textsuperscript{9} A proposed mechanism for the catalytic oxidation of CN\textsuperscript{−} to NCO\textsuperscript{−} by the latent catalyst is shown in Scheme 2.

To further demonstrate the scope of the CDA as a general and smart catalytic system, another substrate, oxalate, was also investigated. The $\Delta G^0$ value (−39.0 kJ mol\textsuperscript{-1}) of the adduct between Cu\textsuperscript{II}(dien)\textsuperscript{2+} and HOOC–COO\textsuperscript{−} was found to be larger than those of the systems 1–3 (−30.4, −25.3, and −15.1 kJ mol\textsuperscript{-1}, respectively, Table 2) (Fig. S6f). Because of this thermodynamic trigger, an off–on control is introduced into the CDA: Fig. S7 in the ESI† shows that, in the presence of 1 under UV-vis irradiation at room temperature, the degradation of the oxalate (3.125 × 10\textsuperscript{−2} M) to CO\textsubscript{2} is boosted rapidly in the first 150 min and gradually increases afterwards. In addition, the kinetic properties of the systems impart smart properties. Fig. S7† also reveals that even though complex 1 could thermodynamically undergo cyano-bridge cleavage in the presence of oxalate to release the Cu\textsuperscript{II} unit (Fig. S4b†) for the catalytic oxidation of oxalate into CO\textsubscript{2}, it does not start the CDA reaction. For example, complex 1 requires that the initial oxalate concentration reaches 1.25 × 10\textsuperscript{−3} M (the threshold) to cleave the cyano-bridge, release the Cu\textsuperscript{II} unit, and catalyze oxalate oxidation. A proposed mechanism for the catalytic oxidation of oxalate to CO\textsubscript{2} by the latent catalyst is shown in Scheme 2. The CDA system is therefore being smart, playing two consecutive roles of sensing a target pollutant and executing a preset chemical process at a specific concentration level of pollutants.

Degradation of cyanide in real wastewater samples by catalyst displacement assay

After carefully verifying the basic and crucial factors (thermo-dynamic and kinetic properties) of the CDA, complex 3 was used to test its applicability and “smart” performance in real
wastewater treatment. Both domestic and industrial wastewater samples (level I, untreated sewage from a residential area system and sewage from an industrial zone, respectively) were collected in Hong Kong, China. The samples were filtered through 0.45 μm pore-size membrane filters (Pall Corporation) to remove insoluble substances before examination. The World Health Organization suggests a maximum allowable level of cyanide in wastewater of 20 μM (0.5 mg L⁻¹). The wastewater samples were spiked with 10, 20 or 30 μM cyanide and their degradation/oxidation was analyzed at room temperature with complex 3 in the presence of H₂O₂. (Based on the results described above, complex 3 was able to initiate the oxidation of cyanide at 20 μM.) In the presence of 3, when the cyanide content of the samples reached 20 μM, the oxidation of cyanide to cyanate was initiated (Table 3; Fig. S8†). However, it is important to note that when the cyanide content of the samples was set to 10 μM (i.e., less than the threshold concentration), no cyanide was oxidized to cyanate throughout the period. Control experiments in the absence of the complex revealed that no cyanide was oxidized to cyanate. These results indicate that the smart-functioning CDA concept is applicable, even in the presence of organic/inorganic matters in actual domestic and industrial wastewater samples.

### Conclusion

Three bimetallic complexes were synthesized using a supramolecular approach with different metallic donors, [FeII(CN)₆]³⁺, [FeII(tBubpy)(CN)₄]²⁻, and FeII(tBubpy)₂(CN)₂, to bridge with a metallic acceptor, [CuII(dien)]³², through a cyano-bridge as latent catalysts. The study of their thermodynamic and kinetic properties provided insight on how to establish a smart off-on sensing and threshold-controlled latent catalytic system for degradation of toxic pollutants. With this “smart” deactivation–activation function for controlling targeted reactions, the catalyst displacement assay (CDA) system could be used for on-site applications such as wastewater treatment with a pre-set threshold for the elimination of cyanide from reservoirs.

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### Table 3 Results of cyanide degradation by 3 in domestic (level I, untreated) and industrial wastewater samples

<table>
<thead>
<tr>
<th>Wastewater samples</th>
<th>Cyanide added (μM)</th>
<th>Cyanate generated (μM)</th>
<th>Conversion after 4 h (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Domestic (level I, untreated)</td>
<td>10</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>20</td>
<td>6.1</td>
<td>40.4</td>
</tr>
<tr>
<td></td>
<td>30</td>
<td>12.9</td>
<td>42.9</td>
</tr>
<tr>
<td>Industrial</td>
<td>10</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>20</td>
<td>6.6</td>
<td>33.1</td>
</tr>
<tr>
<td></td>
<td>30</td>
<td>14.8</td>
<td>49.4</td>
</tr>
</tbody>
</table>

### References


17 J. E. Erman, Biochemistry, 1974, 13, 39–44.


