Citrate-Zn/Al Layered Double Hydroxide as Adsorbent of Congo Red from Aqueous Solution

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Abstract: Layered double hydroxide (LDH) of Zn/Al and citrate-Zn/Al was prepared and used as an adsorbent of Congo red from aqueous solution. LDH was characterized by X-ray, FTIR, and BET analysis. Adsorption of Congo red was studied through kinetic, isotherm, and thermodynamic analyses. Zn/Al LDH has diffraction at 10.29° (003) with interlayer distance 8.59 Å and citrate-Zn/Al LDHs have anomalous diffraction at 7.57° (003) with interlayer distance 11.68 Å. The surface area of citrate-Zn/Al (40.50 m² g⁻¹) has higher than pristine LDH (1.97 m² g⁻¹). Adsorption of Congo red was conducted at pH 6 for Zn/Al LDH and at pH 8 for citrate-Zn/Al LDH. Adsorption of Congo red on both LDHs follows the pseudo-second-order kinetic model. The isotherm parameter follows the Freundlich isotherm model with maximum adsorption capacity 166.67 mg g⁻¹ for Zn/Al and 249.99 mg g⁻¹ for citrate-Zn/Al LDH. Adsorption of Congo red on both LDHs was classified as physical adsorption with energy 4.085-4.148 kJ mol⁻¹.

Keywords: citric acid, layered double hydroxide, Zn/Al, adsorption, Congo red.

1. Introduction

The use of dyes in the industrial process such as painting, film, textile, plastic, pigment, and so on is sharply increased. It causes an environmental problem due to the undegradable properties of dyes. Various methods have been applied to remove stains from an aqueous solution such as adsorption, precipitation, filtration, coagulation, membrane, among these methods, adsorption is a suitable method to remove dyes due to the easy way, simple process, and also low-cost process. The efficiency of the adsorption process depends on the properties of adsorbent.

Wide range organic and inorganic adsorbent has been used to reduce dyes from an aqueous solution such as chitin, chitosan, cellulose, bentonite, kaolin, clay, carbon nanotube, zeolite, and also layered double hydroxide (LDH). LDH is inorganic layer material with the general formula [M²⁺/M³⁺(OH)]ₙ[A⁻]·nH₂O where M²⁺ and M³⁺ is divalent and trivalent metal ions, from stacking brucite-like layers metal oxide containing a positive charge due to partial substitution of divalent metal cation by trivalent metal cation. The negative charge is represented by anions located in the interlamellar spaces. Interlamellar space is also known as interlayer, can exchange species A⁻ to other anions. The interlayer can be expanded to an adapted wide range of interlayer anion. Therefore, A⁻ can be exchangeable with inorganic or organic anionic compounds, such as polyoxometalaate, halide, nitrate, phosphate, carbonate, and also EDTA.

The LDH showed the efficient material as adsorbent of anionic and cationic dyes such as Congo red, safranin, indigo carmin, crystal violet and malachite green, and also methylene blue. LDH M²⁺/M³⁺ (M²⁺ = Zn, M³⁺ = Al, Fe, Cr) were used as adsorbent of direct violet follow pseudo-second-order kinetic adsorption model rather than pseudo-first-order model. Zn/Cr LDH was also used as an efficient adsorbent of Congo red and direct yellow dyes, which were an anionic dyes. Based on these results, the use of LDH as an efficient adsorbent of dyes is the priority. On the other hand, modification of LDH by intercalation with anion onto interlayer distance has been conducted to increase interlayer distance and surface area properties of LDH. Zn/Cr LDH has been intercalated with [α-SiW₁₂O₄₀]⁴⁻ to...
form Zn/Cr-[α-SiW\textsubscript{12}O\textsubscript{40}] by substitution of Keggin type polyoxometalate to nitrate ion of Zn/Cr LDH resulted increasing the surface area from 31.638 m\textsuperscript{2}/g to 128.871 m\textsuperscript{2}/g after intercalation process \textsuperscript{33}. This material after intercalation was effective as inorganic adsorbent. Organic anions such as citrate and malate have been applied as intercalant of Mg/Al LDH and show efficient as an adsorbent for removal lead and copper \textsuperscript{34}.

In this research, Zn/Al LDH was intercalated with citrate ion to increase both interlayer distance and surface area properties of LDH. The material was used as an adsorbent for Congo red from water. The effect of various conditions, including pH, time, initial concentration, and temperature adsorption were investigated in batch adsorption apparatus. Kinetic, isotherm, and thermodynamic adsorption of Congo red on citrate-Zn/Al LDH was discussed in this article. The structure of Congo red was shown in Figure 1.

![Figure 1. Structure of Congo Red](image)

**2. Experimental Section**

**2.1. Chemicals and Instrumentations**

Chemicals were supplied from Merck and Sigma Aldrich, such as zinc nitrate, aluminum nitrate, sodium hydroxide, hydrochloric acid, sodium carbonate, citric acid, and Congo red. These reagents were used directly after purchased without further purification. Water was obtained from the water purification Purite\textsuperscript{®} water purification system equipped with an ion-exchange system at Universitas Sriwijaya. X-ray analysis was conducted using XRD powder Rigaku Miniflex-6000. The sample was homogenized using a mortar and scanned at 2θ speed 1° min\textsuperscript{-1}. FTIR spectra were recorded using Shimadzu FTIR Prestige-21 by KBr method. The sample was examined at wavenumber 400–4000 cm\textsuperscript{-1}. Surface area analysis was measured using a Quantachrome instrument at 77K. Congo red dye was analyzed using spectrophotometer UV-Vis BioBase BK-UV 1800 PC at wavelength 497 nm.

**2.2. Preparation of Zn/Al and Intercalated Citric Acid Zn/Al Layered Double Hydroxides**

Zn/Al LDH was synthesized using the coprecipitation method at pH 10 \textsuperscript{35}. The solution of zinc nitrate (0.3 M, 100 mL) and aluminum nitrate (0.1 M, 100 mL) was mixed under constant stirring. The mixture of sodium hydroxide (1 M, 50 mL) and sodium carbonate (1 M, 50 mL) was added into the mixture, and the pH of the solution mixture was adjusted to 10 by adding sodium hydroxide. The reaction was performed overnight to form a white solid material. The solid material was washed with water and dried at 110°C for 24 hours.

Citric acid-Zn/Al LDH was prepared using the gel method under nitrogen conditions. A citric acid solution (4 g in 250 mL water) was added to a gel form of Zn/Al LDH. The reaction mixture was stirred at 60°C for 1 hour, pH was adjusted to 10 by adding sodium hydroxide solution then the reaction was kept with constant stirring for 24 hours to form citrate-Zn/Al LDH.

**2.3. pH pzc determination** \textsuperscript{36}

The pH at the point zero charge of materials was conducted using 0.05 g adsorbents added into 50 mL the solution of sodium chloride 0.1 M under various pH conditions using hydrochloric acid or sodium hydroxide solution. The solution of sodium chloride 0.1 M was adjusted to pH in the range 1-10 by addition hydrochloric acid 0.1 M or sodium hydroxide 0.1 M solutions. The material of Zn/Al and citric acid-Zn/Al LDH was added into the series of pH solution, and then the solution mixtures were stirred continuously for 24 hours. The solutions were filtered, and pH of the filtrate was determined by pH meter. The graph of pH pzc was obtained by comparison initial and final pH solution.

**2.4. Study Mixture adsorption**

Adsorption of dye using Zn/Al and citrate-Zn/Al LDHs was firstly investigated through mixtures dyes adsorption. Congo red, methyl orange and malachite green were mixed to determine the adsorption competition of dyes. The dyes mixtures were conducted at the acid and base condition. Congo red and methyl orange were anionic dyes, whereas malachite green was cationic dye. The time of adsorption was varied from 5 to 120 minutes. The filtrate was scanned using UV-Vis spectrophotometer in the range 400-700 nm.
2.5. Adsorption of Congo Red
Adsorption of Congo red with Zn/Al and citrate-
Zn/Al LDHs was carried out using a batch small
reactor system. Adsorption was studied through
the effect of adsorption time, the effect of Congo red
concentration, and adsorption temperature. Effect of
adsorption time was investigated at various
adsorption time i.e. 5, 10, 15, 20, 30, 50, 70, 90, 120,
150, 180 minutes. The effect of Congo red
coloration was varied at 40, 50, 60, 80 mg L^{-1} and
at temperature 303, 313, 323, 333 K. Concentration
of Congo red after adsorption process was analyzed
using UV-vis spectrophotometer at 497 nm.

3. Results and Discussion

The XRD powder patterns of Zn/Al (a) and citrate-Zn/Al (b) LDHs are shown in Figure 2. Zn/Al LDH has
diffraction peak at 10.29°(003), 20.07°(006),
31.50°(009), 39.60°(115), 48.95°(118), and
60.20°(110) \(^{37}\). The diffraction peak at 31.50°
indicates the formation of carbonate on the interlayer
distance of Zn/Al LDH. The formation of a well-
ordered layer structure was identified at 10.29° and
60.20° with an interlayer distance of 8.59 Å at
diffraction 10.29 (003). The intercalation of Zn/Al
LDH with citric acid has slightly changed the
diffraction peaks. The crystallinity of material after
the intercalation process was decreased, as indicated
by the formation of a diffraction peak at 15°. The
diffraction of (003) was shifted to 7.56° with
interlayer distance 11.68 Å. Other peaks was detected
at 24.35°(006), 34.50°(009), 39.89°(115),
60.42°(110). The interlayer distance of Zn/Al LDH
was expanded because of the insertion of citrate ion.
The citric acid size is approximately 7.2 Å thus citrate
ion was inserted as slanted orientation as illustrated in
Figure 3.

Figure 2. XRD powder patterns of Zn/Al (a) and citrate-Zn/Al (b) LDHs

Figure 3. Structure of LDH and intercalated with citrate ion.
The FTIR spectra of Zn/Al and intercalated Zn/Al LDHs with citric acid are shown in Figure 4. As shown in Figure 4a, the board peak at 3400 cm\(^{-1}\) was assigned as the –O-H stretching vibration. These vibrations are coming from the water on interlayer distance, water on the hydroxide layer, and also adsorbed water from the air. Water also has to bend vibration at 1610 cm\(^{-1}\) as small vibration in Figure 4a. The sharp peak vibration at 1370 cm\(^{-1}\) is attributed to carbonate vibration. Intercalation of Zn/Al with citrate ion resulted in sharp vibration of citrate ion at 1600 cm\(^{-1}\) as vibration C=O. The vibration of carbonate has still remained. The –CH\(_2\) the vibration of citrate ion was detected at 2900 cm\(^{-1}\) with a small vibration peak. The vibration of water molecule appeared extremely board at 3400 cm\(^{-1}\), similar to pristine LDH.

The surface area of Zn/Al LDH and intercalated Zn/Al-citrate LDH were determined by nitrogen adsorption-desorption. The adsorption-desorption isotherm of materials is shown in Figure 5. There is a hysteresis loop with H3 type on both Zn/Al and intercalated materials. The materials are classifying as mesoporous. The BET analysis is shown in Table 1 and shows that pore diameter was slightly decreased.
after intercalation, but the significant increase of surface area and pore volume after intercalation with citric acid. The morphologies of Zn/Al and Zn/Al-citrate LDH were represented in Figure 6.

Figure 6.a shows the morphology of Zn/Al before citrate intercalation. The morphology of Zn/Al shows the agglomeration of materials caused the pH setting and temperature due to the synthesis process. Similar reported by Palapa et al. (2020), that the agglomeration of LDH indicated the heterogeneity in surface area and active site become low. According to EDX composition Figure 6.a, shows the dominant component is oxygen from water and anion; Zn forms divalent metal of LDH; N from nitrate and also Al from trivalent metal cation of LDH structure. Figure 6.b indicated the intercalation process was successfully conducted and EDX composition of the material was presented. The morphology of Zn/Al-citrate also shows the agglomeration of materials. The white particle indicated the morphology of citrate. According to Rahmanian et al. (2018), the LDH intercalated citrate anion has homogeneity plates-like shape with nano-size particles.

Table 1. BET Analysis.

| Properties          | LDHs       |
|---------------------|------------|
|                     | Zn/Al      | Citrate-Zn/Al |
| BET surface area (m² g⁻¹) | 1.968      | 40.502       |
| Pore volume (cm³ g⁻¹), BJH | 0.006      | 0.116        |
| Pore diameter (nm), BJH | 27.687     | 16.350       |

The surface charges of material were determined using pH pzc (point zero charges), as shown in Figure 7. The intersection point of Zn/Al LDH was at pH 6, whereas intercalated material at pH 8. At that point, the surface charges of materials were zero. The point of under pH pzc indicated that the materials’ surface is positively charge and vice versa. Thus the application of adsorption was conducted at pH pzc.
Figure 7. pH pzc graph of initial pH (a), Zn/Al (b) and citrate-Zn/Al (c).

Figure 8. UV-visible spectra of a mixture of dyes (congo red, methyl orange, malachite green) with pH 4 (above) and pH 9 (below) after treating with Zn/Al and citric acid-Zn/Al LDHs.

Figure 8 shows the adsorption of mixture dyes using Zn/Al and citrate-Zn/Al LDHs. The dyes were Congo red and methyl orange as anionic dyes and malachite green as a cationic dye. The adsorption was conducted at pH acid and base. All dyes were detected at acid condition, but malachite green was not detected at base condition due to anionic dyes reacted with a hydroxyl group. At acid condition, the absorbance of Congo red was slightly decreased by increasing adsorption time.

On the other hand, the absorbance of methyl orange and malachite green was almost stable. The amount of Congo red was adsorbed dominantly than methyl orange or malachite green on both Zn/Al and intercalated materials. Thus adsorption of Congo red was further investigated through the effect of adsorption time, the concentration of Congo red, and temperature adsorption.
Figure 9. Effect of adsorption time on Zn/Al (a) and citrate-Zn/Al (b) LDHs

The effect of adsorption time of Congo red on Zn/Al and intercalated Zn/Al LDHs material was shown in Figure 9. The adsorbed Congo red was gradually increased by increasing adsorption time and reach equilibrium at 120 minutes. Adsorption of Congo red on intercalated LDH material is higher than pristine Zn/Al LDH. The data in Figure 8 was used to obtain adsorption rate $k_1$ and $k_2$ using pseudo-first-order and pseudo-second-order kinetic model as follows:

**Pseudo first-order kinetic model:**

$$\log (q_e - q_t) = \log q_e - \left(\frac{k_1}{2.303}\right) t$$ \hspace{1cm} (1)

**Pseudo second-order kinetic model:**

$$\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{1}{q_e} t$$ \hspace{1cm} (2)

where: $q_e$ is adsorption capacity at equilibrium (mg.g$^{-1}$); $q_t$ is adsorption capacity at t (mg.g$^{-1}$); t is adsorption time (minute); and $k_1$ is adsorption kinetic rate at pseudo first-order (minute$^{-1}$).

Pseudo-second-order kinetic model:

$$\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{1}{q_e} t$$ \hspace{1cm} (2)

where $q_e$ is adsorption capacity at equilibrium (mg.g$^{-1}$); $q_t$ is adsorption capacity at t (mg.g$^{-1}$); t is adsorption time (minute); and $k_2$ is adsorption kinetic rate at pseudo second-order (g.mg$^{-1}$.minute$^{-1}$). The data of kinetic adsorption is shown in Table 2.

| Kinetic Adsorption Model | Kinetic Parameter | LDH               |
|--------------------------|-------------------|-------------------|
|                          |                   | Zn/Al             | Citrate-Zn/Al |
| Pseudo First-Order       | $Q_e$ (mg g$^{-1}$) | 25.704            | 33.037        |
|                          | $Q_e$ experiment (mg g$^{-1}$) | 29.208 | 44.958 |
|                          | $R^2$             | 0.972             | 0.971         |
|                          | $k_1$ (min$^{-1}$) | 0.0299            | 0.0276        |
| Pseudo Second-Order      | $Q_e$ (mg g$^{-1}$) | 30.303            | 50.001        |
|                          | $Q_e$ experiment (mg g$^{-1}$) | 29.208 | 44.958 |
|                          | $R^2$             | 0.991             | 0.999         |
|                          | $k_2$ (g mg$^{-1}$min$^{-1}$) | 0.0014 | 0.0016 |

The pseudo second-order kinetic model is appropriate for both adsorbents due to R value close to one. The $k_2$ value for intercalated material is slightly larger than Zn/Al without intercalation. The reactivity of intercalated material is probably due to the opening of interlayer distance after intercalation. According to Sepehr et al (2012), the kinetic models determined by coefficient correlation and the value of $q_e$ experiment are close to $q_e$ calculation. In Table 2, the $q_e$ experiment use pseudo-second order, both LDH close with $q_e$ calculation.
Figure 10 shows the effect of initial concentration and temperature adsorption of congo red on Zn/Al and citrate-Zn/Al adsorbents. The amount of congo red adsorbed increased by increasing temperature adsorption. The adsorption profile shows that physical adsorption has occurred in this research. The data of isotherm adsorption was obtained using Langmuir and Freundlich isotherms equation as follows:

**Langmuir equation:**

\[
C_m = \frac{1}{bK} + \frac{C}{b}
\]  

where \( C \) is a saturated concentration of adsorbate; \( m \) is the amount of adsorbate; \( b \) is the maximum adsorption capacity (mg g\(^{-1}\)), and \( K_{ML} \) is the Langmuir constant (L mg\(^{-1}\)).

**Freundlich equation:**

\[
\log q_e = \log K_F + \frac{1}{n} \log C_e
\]  

where \( q_e \) is adsorption capacity at equilibrium (mg g\(^{-1}\)); \( C_e \) is a concentration of adsorbate at equilibrium (mg L\(^{-1}\)), and \( K_F \) is Freundlich constant. The data of isotherm adsorption of congo red on LDHs are shown in Table 3.

**Table 3. Isotherm adsorption of congo red.**

| Isotherm | Isotherm Parameter | Zn/Al | Citrate-Zn/Al |
|----------|--------------------|-------|---------------|
| Langmuir | \( q_{max} \) (mg g\(^{-1}\)) | 47.619 | 249.990 |
|          | \( K_{ML} \) (L mg\(^{-1}\)) | 0.011 | 0.009 |
|          | \( R^2 \)           | 0.580 | 0.171 |
| Langmuir | \( q_{max} \) (mg g\(^{-1}\)) | 166.666 | 12.882 |
|          | \( K_{ML} \) (L mg\(^{-1}\)) | 0.007 | 0.018 |
|          | \( R^2 \)           | 0.433 | 0.426 |
| Freundlich| \( K_F \) (mg g\(^{-1}\)) | 166.666 | 161.11 |
|          | \( R^2 \)           | 0.774 | 0.746 |
| Freundlich| \( K_F \) (mg g\(^{-1}\)) | 111.11 | 112.82 |
|          | \( R^2 \)           | 1.315 | 1.283 |

Table 3 showed that adsorption capacity for citrate-Zn/Al is higher than pristine Zn/Al LDH. The adsorption capacity is up to 249.99 mg g\(^{-1}\) for citrate-Zn/Al LDH and 166.67 mg g\(^{-1}\) for Zn/Al LDH without intercalation. The adsorption capacities are related to the increased surface area properties after intercalation. The decreasing of adsorption capacity of intercalated material by increasing temperature is probably due to decreasing pore diameter of intercalated material, as shown in Table 1. The compared materials as adsorbent of congo red were shown in Table 4.
Table 4. Compared materials as an adsorbent for Congo red dye.

| Adsorbent                  | Adsorption Capacity (mg/g) | Ref |
|----------------------------|----------------------------|-----|
| Ni/Co LDH                  | 909.2                      | 45  |
| Mg/Fe-CO₃                  | 104.6                      | 46  |
| Fe₃O₄-MgAl LDH             | 813                        | 47  |
| CNT-MgAl-O LDH             | 1250                       | 48  |
| sodium dodecyl sulfate SDS-LDH | 714                       | 49  |
| Borates-Mg/Al LDH          | 138.6                      | 50  |
| Ni/Al-S₁ LDH               | 120.5                      | 51  |
| MgO-GO microspheres        | 227.7                      | 52  |
| NiFe LDH                   | 80.2                       | 53  |
| Mg/Fe-NO₃ LDH              | 58.54                      | 53  |
| Zn/Al LDH                  | 112.04                     | This work |
| Zn/Al-citrate              | 249.9                      | This work |

The thermodynamic properties of Congo red adsorption on Zn/Al and intercalated Zn/Al LDHs is shown in Table 5 and was calculated according to the equation as follows 54:

\[
\ln K_{ML} = \frac{\Delta S}{R} - \frac{\Delta H}{RT}\]  \hspace{1cm} (5)

\[
\Delta G^\circ = -RT \ln K_{ML}\]  \hspace{1cm} (6)

Where T is the temperature (K); R is the gas constant (8.314 J mol⁻¹ K⁻¹), and \( K_{ML} \) is the modified Langmuir constant.

Table 5. Thermodynamic parameter of Congo red adsorption on LDHs.

|                  | \( \Delta G^\circ \) (kJ mol⁻¹) | \( \Delta H^0 \) (kJ mol⁻¹) | \( \Delta S^0 \) (J mol⁻¹ K⁻¹) |
|------------------|---------------------------------|----------------------------|-------------------------------|
|                  | 303 K                           | 313 K                      | 323 K                         | 333 K                         |
| Zn/Al            | -0.213                          | -0.355                     | -0.497                        | -0.638                        | 4.085                         | 0.014 |
| Citrate-Zn/Al    | -1.487                          | -1.673                     | -1.859                        | -2.045                        | 4.148                         | 0.019 |

Figure 11. XRD pattern of (a) Zn/Al, (b) Zn/Al after adsorption, (c) Zn/Al-Citrate and (d) Zn/Al-Citrate after adsorption Congo red.
The data in Table 5 showed the value of Gibbs energy, enthalpy, and entropy of Congo red adsorption on LDHs. All adsorption of methyl orange on LDHs was spontaneously occurred due to the negative value of Gibbs energy. Adsorption of Congo red on LDHs was classified as physical adsorption with energy in the range $4.085 - 4.148$ kJ mol$^{-1}$ and also there was increasing randomness of adsorption system on both LDHs as adsorbent. The adsorption of Congo red into Zn/Al-Cit LDH was more efficient than Zn/Al LDH. The study of a LDH structure after the adsorption process was conducted using XRD analysis. The pattern of XRD before and after adsorption of Congo red was shown in Figure 11. Figure 11.b. shows the LDH pattern after adsorption of Congo red was shifted to lower 2 theta from 10.29 with interlayer distance 8.59 Å to 9.725 with interlayer distance 8.54 Å. The interlayer was decreased after adsorption, but the structure of LDH has no change. However, intercalated LDH shows the adsorption capacity higher than pristine LDH. This phenom related that the decrease interlayer after adsorption Congo red and 2 theta shifted to a higher degree. Figure 11. c and d show the shifted with decrease the interlayer distance from 11.68 Å to 8.65 Å indicated that the Congo red was coated and piled up the active site so that the XRD confirm that the interlayer decreased slightly. The effectivity of adsorbent was determined by reusability. Desorption study was conducted before the regenerator process, desorption was conducted using several reagents to determine the suitable solvent, and the results were shown in Figure 12.

**Figure 12.** Desorption Zn/Al LDH and Zn/Al-Citrate LDH

**Figure 13.** Reusability Zn/Al LDH and Zn/Al-Citrate LDH
Figure 12 shows the desorption of Zn/Al and Zn/Al-Citrate LDHs were conducted using HCl, hot water, NaOH, hydroxylamine hydrochloride, water, ethyl acetate, and ethanol. Before desorption, both adsorbents were adsorbed the 150 mg/L of Congo red and adsorbed capacity was 96 mg/g and 132 mg/g at room temperature. The adsorbed materials were shaken with each reagent. The result obtained that the LDH pristine using HCl uptake 98.13 % and Zn/Al intercalated citrate using NaOH with 93.11 % desorption. Subsequently, Zn/Al and Zn/Al-citrate after desorption used to Congo red adsorption. The removal efficiency of these materials showed in Figure 13.

The removal efficiency of both materials slightly decreased. The Zn/Al LDH was decreased dramatically after a four-cycle process. According to Palapa et al. (2020), the LDH can be exfoliated in acid solution, and LDH has reduced reusability activity. Nevertheless, Zn/Al-citrate has more reactivity to reuse as adsorbent. The removal efficiency of Zn/Al-citrate is quietly decreased until the fourth cycle. The Zn/Al after intercalating citrate anion has more efficient as an adsorbent to remove Congo red in water solution.

4. Conclusion

Citrate-Zn/Al LDH and pristine LDHs were successfully synthesized and used as adsorbent of Congo red from aqueous solution. The adsorption of Congo red on LDHs follows the pseudo-second-order kinetic model. The adsorption capacity is up to 249.99 mg g⁻¹ for citric acid-Zn/Al LDH and 166.67 mg g⁻¹ for pristine LDH. The adsorption process was categorized as physical adsorption with energy adsorption 4.085-4.148 kJ mol⁻¹. The reusability study shows the Zn/Al after intercalated citrate anion is efficient and effective use as an adsorbent.

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