Zeolite/Cellulose Acetate (ZCA) in Blend Fiber for Adsorption of Erythromycin Residue From Pharmaceutical Wastewater: Experimental and Theoretical Study

Shehdeh Jodeh1*, Israa Erman1, Othman Hamed1*, Younes Massad1, Ghadir Hanbali1, Subhi Samhan2, Omar Dagdag3, Savaş Kaya4 and Goncagül Serdaroğlu5

1Department of Chemistry, Faculty of Science, An-Najah National University, Nablus, Palestine, 2Palestinian Water Authority, Ramallah, Palestine, 3Laboratory of Agroresources, Polymers and Process Engineering (LAPPE), Department of Chemistry, Faculty of Science, Ibn Tofail University, Kenitra, Morocco, 4Department of Pharmacy, Health Services Vocational School, Sivas Cumhuriyet University, Sivas, Turkey, 5Mathematics and Science Education, Sivas Cumhuriyet University, Sivas, Turkey

The expanding amount of remaining drug substances in wastewater adversely affects both the climate and human well-being. In the current investigation, we developed new cellulose acetic acid derivation/zeolite fiber as an effective technique to eliminate erythromycin (ERY) from wastewater. The number of interchangeable sites in the adsorbent structures and the ratio of ERY to the three adsorbents were identified as the main reasons for the reduction in adsorption as the initial ERY concentrations increased. Additionally, for all adsorbents, the pseudo–second-order modeling showed better fitting for the adsorption than the pseudo–first-order modeling. However, the findings obtained in the pseudo–first-order model were still enough for explaining the sorption kinetics of ERY, showing that the surface displayed all chemisorption and physi-sorption adsorption processes by both adsorbents. The R² for the second order was very close to 1 for the three adsorbents in the case of pseudo–second-order. The adsorption capacity reached 17.76 mg/g. The three adsorbents showed negative values of ΔH, and these values were −6,200, −8,500, and −9600 kJ/mol for zeolite, CA, and ZCA, respectively, and this shows that the adsorption is exothermic. The desorption analysis shows no substantial loss of adsorption site after three trials, indicating higher stability and resilience of the three adsorbents, indicating a strong repeatability of their possible use in adsorption without contaminating the environment. In addition, the chemical attitude and possible donor–acceptor interactions of ERY were assessed by the quantum chemical parameters (QCPs) and NBO analysis performed, at the HF/6-311G** calculations.

Keywords: pharmaceutical industry, adsorption capacity, wastewater, cellulose, zeolite
INTRODUCTION

One of the most important reasons for the economic growth of developing countries and the expansion of urban areas is a society’s ability to provide fresh water for sanitation and consumption to its population. However, as the population and urbanization increase, so does the release of radioactive materials into the atmosphere and surface water. There are many sources of surface and groundwater contamination, including agricultural, industrial, oil pollution, sewage, and wastewater (Al-Shaalan et al., 2019; El-Zawily et al., 2019; Khan et al., 2019; Chon et al., 2020).

Several water pollution scenarios including the chiral pollution are a serious issue for our health and environment due to the enantioselective biodegradation of the chiral pollutants. It has adverse impact on our society and science. There is a big loss of our economy due to the use of racemic agrochemicals. The most notorious chiral pollutants are pesticides, polychloro biphenyls, polyaromatic hydrocarbons, brominated flame retardants, drugs, and pharmaceuticals (Basheer, 2018a; Basheer and Ali, 2018).

Nowadays, water contamination due to the drugs and pharmaceutical residues is increasing and alarming. These contaminants are called as new emerging pollutants. The contamination due to the new emerging contaminants is of great concern due to their endocrine, hormonal, and genetic disturbance nature (Basheer, 2018b).

In environmental samples such as surface water, groundwater, seawater, soil, and drinking water, pharmaceuticals were found (Arshad et al., 2020; Kiszkiel-Taudul, 2021), so they are referred to as emerging pollutants. The estimated global consumption of pharmaceuticals such as antibiotics is 100,000 to 200,000 tons per annum (Bungau et al., 2020). Based on the chemical properties of the drug, about 5–90% of the absorbed antibiotic doses are excreted by urine or stool as a metabolite or parent compound (Bhowmick et al., 2020). These drugs end up in drainage systems, or disposal of unwanted or un

METHODS AND MATERIALS

Chemicals

The zeolite compounds are containing aluminum and silicon (M$_2$/nO.Al$_2$O$_3$.xSiO$_2$.yH$_2$O) where $M$ can be any one of a number of metals, including sodium, lithium, potassium, calcium, and magnesium. The variable “$n$” stands for the valence of the metal and other elements.

To deal with this wastewater problem, lots of conventional and advanced technologies have been developed (Ali et al., 2018a; Mery-Araya et al., 2019). The conventional water treatments such as oxidation (Ma et al., 2020), electro precipitation, membrane separation, coagulation–floculation, evaporation, floatation, and ion exchange (Yu et al., 2021) have been largely used, but these are inadequate techniques for water treatments (Tabassum, 2019).

Many approaches have been used and reported for the removal of a variety of pesticides and drugs. Among the different methods, adsorption is the best approach because of several advantages associated with adsorption including time and cost (Ali et al., 2018a; Ali et al., 2018b; Ali et al., 2019).

Erythromycin (ERY) is a natural antibiotic used to treat a variety of bacterial infections. Antibiotics pass into the human body after consistent treatment and ultimately enter inland areas and effluents; there is even a path of environmental degradation in the poultry and livestock breeding industries. Because of the structure of their aromatic ring, ERY molecules are resistant to the environment and difficult to degrade. Several reports (Ma et al., 2020; Yu et al., 2021) have reported the presence of ERY in water and wastewater to be above the average range. As a result, removing ERY residues from wastewater is important.

Zeolite is a crystalline aluminosilicate with well-defined micropore dimensions and a strong crystal lattice form that is environmentally friendly. Zeolite structures are made up of tetrahedral SiO$_4$ and AlO$_4$ groups, and their alumina silica ratio (SAR) determines zeolite polarity (Martucci et al., 2012). Because of their three-dimensional framework, which creates nanometer-sized channels and cages, these materials have a high porosity and a large surface area. The shape of their internal pore structure can have a direct impact on their adsorption selectivity against host molecules, which is one of their distinguishing features (Zide et al., 2018).

Cellulose acetate is an excellent candidate for use as a polymer matrix because it can be easily molded into a variety of shapes and because its hydrophilic surfaces can improve the mobility of aqueous solutions to the surface of hybrid materials (Das et al., 2020). The aim of this research was to use zeolite/cellulose acetate blended fiber as a reusable, simple-to-prepare adsorbent for erythromycin adsorption. The effects of several parameters, including contact time, concentration effect, temperature effect, and equilibrium and kinetics, on erythromycin adsorption by the composite fiber were studied.

SEM, FT-IR spectroscopy, thermogravimetric analysis, and dynamic scanning calorimetry were used to characterize the zeolite/cellulose acetate fiber.

The novelty of this work is shown by using three different adsorbents which showed very high percentage of removals. Also, theoretical studies were very supportive of the experimental findings.
cation and “y” for the number of water molecules in the structure of zeolite, according to the Research Foundation at State University of New York (SUNY). Cellulose acetate (C_{10}H_{16}O_{8}) has been purchased from Al Quds Chemicals in Jerusalem. The zeolite chemical composition was included in the MSDS that has been supplied from the manufacturer. Acetone was bought from Guangzhou Chemi. Erythromycin with technical grade of 99% was purchased from Fluka (Fluka Chemie AG, Switzerland). Acetonitrile was purchased from Sigma–Aldrich, United States with analytical grade of more than 99%. The water was of the Milli-Q standard (Millipore, MA, United States).

### Preparation of ZCA Fiber

Wet spinning was used to produce the zeolite/cellulose acetate blend fiber (ZCA); cellulose acetate (6 g) was dissolved in 50 ml of acetone/water solution (6:1, w/w). The zeolite rocks were ground and sieved to achieve an average dimension of approximately 800 mesh.

1.5 g of zeolite is added to the solution and scattered by mechanical stirring. To make a solid filament, the blended solution was spun in a stainless-steel spinner and then protruded into a water coagulation tank. The fiber was taken out of the bath and washed twice with filtered water. Finally, the fiber was dried at 30°C before being cut into very small fragments (Rodchanasuripron et al., 2020).

### Characterization of ZCA Fiber

The scanning electron microscopy (SEM) manufactured by the Hitachi model (S-4700) in Japan was used to study the morphology of ZCA fiber. ZCA fiber was immersed in a liquid nitrogen atmosphere to create a very clean cross section for scanning. The Hitachi S-4700 FE-SEM is a cold nitrogen atmosphere to create a very clean cross section for ultrahigh resolution imaging of thin films and semiconductor materials on exceptionally clean specimens. It is also suitable for polymeric materials. S-4700 is configured to detect secondary and backscattered electrons as well as characteristic X-rays.

The X-Ray diffraction analysis was done using XRD-Shimadzu XD-1 with monochromatized graphite Cu-K alpha (15,418) and a scanning speed of 20°/min. The Bruker Alpha-P spectrophotometer was used to collect the Fourier transform infrared (FTIR) fiber spectrum. FT-IR spectra were reported from 400 to 4,000 cm\(^{-1}\) with 32 scans on Nicolet NEXUS-470 FT-IR (America) apparatus and a resolution of 4 cm\(^{-1}\).

The Shimadzu UV absorption spectrum of the sample was tested using an 1800 UV-Vis spectrophotometer with UV probe software. The ERY concentration was measured quantitatively using a UV–Vis spectrophotometer (SHIMADZU, UV-1201). The absorbance of the ERY solution was estimated at 481.5 nm, the wavelength at which ERY has the greatest absorbance.

CuK Al radiation was used for X-ray diffraction on the Panalytical X′Pert Pro diffractometer (1.5418 Å) from 2° to 70° (2θ), with a scanning rate of 1° per minute. The water intrusion process was also used to determine membrane porosity (Wang et al., 2017; Bagaev et al., 2021).

Thermogravimetric analysis was carried out on DTG 60H equipment (Shimadzu Co., Japan). Around 3.0 mg of absorbents were heated from 25 to 700°C in the nitrogen atmosphere (50 ml/min) at a temperature of 10 0°C/min. The compounds’ decomposition temperatures were calculated using the first mass loss (percentage) vs. temperature derivative (DTGA) (Güler et al., 2013; Zhang et al., 2014).

### Adsorption Procedure (Import)

Erythromycin [C\(_{37}\)H\(_{67}\)NO\(_{13}\)] with molecular mass of 733.937 g mol\(^{-1}\) is an antibiotic used for the treatment of a variety of Fluka. The chemical structure is presented in Figure 1.

To study the adsorption equilibrium experiments, a sample of 10.0 mg of ZCA fiber was used in most of the analysis. Following that, 100 ml of aqueous solutions with varying initial ERY concentrations (10–50 mg/L) were applied and shaken at 200 rpm in an orbital incubator (Gallenkamp, model INR-250). To achieve adsorption equilibrium, the contact time was varied between 5 and 90 min. The other study was performed to see the effect of temperature on the adsorbent activity and efficiency at different temperatures and constant contact time of 30 min, and the temperatures were 25, 35, 45, and 55°C. In each study, a UV-Vis (Varian, model Cary 1E) spectrophotometer (λ\(_{\text{max}}\): 482 nm) was used to measure ERY equilibrium concentrations using a calibration curve of different concentrations (Jamshaid et al., 2020).

The effect of pH was studied from 2 to 12, and both 0.1 M NaOH and 0.1 M HCl solutions were used to change the pH as required. At 293 K, 100 ml of ERY solution containing 20 mg/L was shook with 10.0 mg of ZCA fiber.

The pH study was carried using a micro pH 2002 Crison pH meter. All equilibrium concentrations of the adsorbed ERY by ZCA were presented using different adsorption parameters; \(q_e\) (e.g., in mg/g) was calculated using the following equations (Eqs 1, 2) (Abujaber et al., 2018):

![Structure for erythromycin.](image)
\[
q_t = \frac{V(C_0 - C_t)}{W}, \quad (1)
\]
\[
\%R = \left(\frac{C_0 - C_e}{C_0}\right) \times 100\%, \quad (2)
\]

where \(q_e\) is the amount (mg g\(^{-1}\)) adsorbed, \(C_0\) and \(C_t\) are the ERY initial and equilibrium concentrations in solution (mg/L), respectively, \(W\) is the adsorbent dosage (g/L), and \(R\) percent is the adsorption efficiency coefficient. The kinetic study was done by taking several dosages of ZCA (50, 100, and 150 mg/L). This study’s tested temperatures were 293, 303, and 313 K, with a maximum contact time of 60 min.

**Adsorption Kinetics**

Pseudo–first-order and second-order models have been used to model the kinetic effects of ERY adsorption on the surface of ZCA fibers to achieve the control rate structure of adsorption including chemical reactions and mass transfer. As seen in Eq. 3, pseudo–first-order modeling is based on the premise that physical adsorption that occurs during the removal process is the rate-determining step (Azzaoui et al., 2017):

\[
\log\left(\frac{q_t}{q_e}\right) = \log q_e - \left(\frac{K_1}{2.303}\right)t, \quad (3)
\]

where \(q_t\) (mg/g) represents the equilibrium adsorbed ERY quantity, \(q_e\) (mg/g) represents the equilibrium adsorbed ERY quantity at time \(t\), and \(K_1\) (min\(^{-1}\)) represents the pseudo–first-order modeling adsorption rate constant (Azzaoui et al., 2017). The modeling of the pseudo–second-order, on the other hand, was based on the assumption that the rate-determining process, as shown by Eq. 4, is chemi-sorption:

\[
\frac{t}{q_t} = \frac{1}{q_e} t + \frac{1}{K_2 q_e^2}, \quad (4)
\]

where \(K_2\) (g/mg/min) is used as the pseudo–second-order rate constant. The slope and intercept of a plot of \(t/q_t\) vs. \(t\) are used to calculate the values of \(q_e\) and \(K_2\), respectively. ERY was controlled for attachment to the ZCA surface through chemical bond forming in the chemical adsorption process.

**Adsorption Isotherm**

Adsorption isotherms usually have data on the distribution of adsorbed molecules in equilibrium between solid and liquid phases. Most experiments used the regression coefficient (R\(^2\)) to assess the best-fitting isotherms. Adsorption equilibrium results were discovered to be more appropriate for two types of Freundlich and Langmuir isothermal models.

The most fundamental model is Langmuir, which assumes that all adsorption sites are equal and autonomous. The tendency of molecules to bind is separate from the neighboring populated sites (Radi et al., 2015). The isotherm of Langmuir can be given by the following equation:

\[
\frac{C_e}{q_e} = \frac{1}{q_m} C_e + \frac{1}{q_m K_L}, \quad (5)
\]

where \(C_e\) is the ERY equilibrium concentration (mg/L), \(q_e\) is the sum of ERY adsorbed per gram of the three equilibrium adsorbents (mg/g), and \(q_m\) is the full potential of monolayer coverage (mg/g) (Radi et al., 2015). The Langmuir isotherm (L/mg) constant is \(K_L\).

The Freundlich isotherm, on the other hand, demonstrates un-ideal and reversible adsorption. The best representation of heterogeneous structures is preferred. It is possible to approximate Freundlich isotherm by the following equation:

\[
\ln q_e = \frac{1}{n} \ln C_e + \ln K_F, \quad (6)
\]

where \(q_e\) is the capacity of adsorption, \(C_e\) is the ERY concentration at equilibrium, and \(K_F\) and \(n\) are constants. \(K_F\) reflects the capacity of adsorption, whereas \(n\) reflects the deviation from linearity of adsorption. If \(n = 1\), the process of adsorption is linear; if \(n < 1\) the process of chemical adsorption; and if \(n > 1\), the process of adsorption is favorable. The Langmuir model is limited to monolayer adsorption. The Langmuir model is limited to monolayer adsorption systems, whereas in multilayer systems, the Freundlich model can be used.

**Adsorption Thermodynamics**

From the obtained kinetic data, the reaction rate and other thermodynamics parameters can be identified. Nonetheless, the response changes that will happen during the process of adsorption require the determination of the thermodynamic parameters, including entropy \((\Delta S, \text{kJ/mol})\), enthalpy, free-energy Gibbs \((\Delta G, \text{kJ/mol})\), and adsorption changes \((\Delta H, \text{kJ/mol})\). You can calculate the thermodynamic parameters from the van’t Hoff Eq. 7:

\[
\ln K_d = \frac{-H^o}{RT} + \frac{S^o}{R}, \quad (7)
\]

where the gas constant is \(R\) (8.314 J/mol/K) and the temperature is \(T\) (K). Eq. 8 can be used to calculate the distribution coefficient (Kd) on the adsorbent surface.

\[
K_d = \frac{V(C_0 - C_e)}{W q_e}, \quad (8)
\]

Gibbs free energy can be calculated by the following equation:

\[
\Delta G^o = -RT \ln K_d. \quad (9)
\]

Both \(\Delta H\) and \(\Delta S\) can be calculated using both slope and intercept from the van’t Hoff plot of lnK vs. 1/T (Hanbali et al., 2020).

**Computational and Theoretical Study**

The geometry optimization of ERY was performed by G09W (frisch et al., 2009) with the Hartree–Fock (Roothaan, 1951; Pople and Nesbet, 1954) method and 6-311G** (Krishtan et al., 1980; McLean and Chandler, 1980) basis set in the gas phase. In theoretical predictions of the chemical reactivity, the Koopmans’ theorem (Koopmans, 1934) is the first essential step to calculate the ionization energy \((I)\) and electron affinity \((A)\) values via the FMO energies.

\[
I = -E_{HOMO}
\]
\[ A = -\text{E}_{\text{LUMO}} \]

Moreover, the quantum chemical parameters (QCP) (Parr and Pearson, 1983; Pearson, 1986; Pearson, 1988; Parr et al., 1999), which are defined as \( \chi \) “electronic chemical potential,” \( \eta \) “global hardness,” \( \omega \) “electrophilicity index,” and \( \Delta N \) “fractional number of the electrons transferred” in case of B and C systems have contacted each other, and \( \Delta N_{\text{max}} \) “maximum charge transfer capability,” have been also obtained from the \( I \) and \( A \) values using the following formula:

\[
\begin{align*}
\chi &= \frac{I + A}{2}, \\
\eta &= \frac{I - A}{2}, \\
\omega &= \frac{\mu^2}{2\eta}, \\
\Delta N &= \frac{\chi_C - \chi_B}{2(\eta_C + \eta_B)}, \\
\Delta N_{\text{max}} &= \frac{I + A}{2(I - A)}.
\end{align*}
\]

In addition, Gázquez and coworkers introduced two useful parameters to calculate the \( \omega^+ \) “the electron-donating power” and \( \omega^- \) “the electron-accepting power” parameters (Gázquez et al., 2007)

\[
\begin{align*}
\omega^+ &= (I + 3A)^2 / (16(I - A)), \\
\omega^- &= (3I + A)^2 / (16(I - A)).
\end{align*}
\]

Also, the \( \Delta\varepsilon_{\text{back-donation}} \) “back-donation energy” (Gómez et al., 2006) is a powerful value and defined as the following equation:

\[ \Delta\varepsilon_{\text{back-donation}} = -\frac{\eta}{4}. \]

In addition, the stabilization energy lowering obtained from the second-order perturbative energy analyses depending on the NBOs “Natural Bon Orbitals” (Foster and Weinhold, 1980; Reed and Weinhold, 1985; Reed et al., 1988) is defined as follows:

\[ E^{(2)} = E_{ij} = q_i \left( \frac{F_{ij}}{(e_j - e_i)} \right)^2. \]

For the molecular system, \( q_i \) states the donor orbital occupancy, \( e_i \) and \( e_j \) are diagonal elements, and \( F_{ij} \) is the off-diagonal NBO Fock matrix element where “i” and “j” are the filled and unfilled molecular orbitals.

**Regeneration of Adsorbent**

In the field of adsorption process applications, adsorbent regeneration is important. ZCA samples were pre-adsorbed for 12 h at 25°C with 10 ml of 50 mg/L ERY solution, then washed with methanol/acetic acid (v/v, 9:1) until no ERY was present in the eluent, and dried overnight at 50°C. Following that, regenerated materials were redistributed in 10 ml solutions containing an initial concentration of 50 mg/L. The effectiveness of ERY adsorption by regenerated materials has been studied after several adsorption–desorption processes.

**RESULTS AND DISCUSSION**

**Adsorbent Characterization Results (BET)**

Nitrogen adsorption–desorption isotherm measurements were carried at 77 K using a Quantachrome Autosorb AS-1 instrument (United States). The BET specific surface area of ZCA was measured using the data of nitrogen adsorption isotherm at low temperature (Brunauer et al., 1938) and involving the adsorption data at \( P/P_0 \) of 0.05–0.2 and with 2.47 m²/g. The BJH model was used to measure the pore volume and the average pore size as other previous study (Barrett et al., 1951). The pore volume of ZCA sample was determined as \( 2.45 \times 10^{-2} \) cm³/g, and pore diameter was 3.5 nm. The ZCA pore diameter was considered as a mesoporous material as the classification by the Pure and Applied Chemistry International Union (IUPAC) (Foster and Weinhold, 1980).
Characterization of ZCA Fiber Using SEM

SEM was used to examine the morphology of ZCA fiber. The surface morphologies and cross-sectional configurations of the ZCA filament are shown in Figure 2. The surface of the ZCA fiber is relatively smooth, as seen in Figure 1, and the diameter of the as-prepared fiber is approximately 250 nm. As seen in the cross-section, the ZCA fiber has a sponge-like appearance. The ZCA fiber is composed of a homogeneous, highly porous material. The ZCA fiber network is embedded with zeolite crystals about 100 μm in height. As seen in Figure 2, cellulose acetate serves as a matrix support, and the pore size of the fiber ranges between 5 and 10 μm. ERY could rapidly disperse into the pores for contact with the adsorptive sites of the ZCA particles.

The dispersion of zeolite attributed by the silica and aluminum shown in Figures 1B,C indicates that the zeolites were embedded in the cellulose acetate matrix. This is attributed to the interfacial interaction between zeolite and cellulose acetate.

X-Ray Diffraction Analysis

The diffractogram of the synthesized zeolite is identical to JCPDS No. PDF 0038-0241 for LTA form zeolite-A [Na96(AlO2) 96(SiO2)96.216H2O] as seen in Figure 3. Furthermore, diffractogram of CA, as shown in the figure, appropriates with a diffractogram reported by Fan et al. (2013), who stated that CA has distinctive angles at 2θ of 10° and 13.2°. These two typical angles were also recognized as the crystalline peaks of modified CTA II (Deus et al., 1991). Furthermore, Jayalakshmi et al. (2014) announced that the CA membrane diffractogram had a normal semicrystalline angle at 2θ of 9.6° and two crystalline angles at diffraction angles of 20.1° and 26.8°.

The diffractogram of CA membrane in this study was identified as a crystalline peak at 26.8°. Composite membrane also has a crystalline peak at 26.8°. Moreover, the composite membrane has also a weak peak at 10° and 13.2°, indicating the typical peak of CA in different intensities. It was caused by a decreasing crystallinity form in the membrane compared to CA solids. It was reviewed that the CA/ZA membrane has a peak at an angle of 10.3, 12.6, and 16.2, indicating the presence of zeolite-A. Based on the results of the composite membrane diffractograms, it was known that zeolite-A has better dispersity in the CA porous membrane as a filler.

FT-IR Analysis

Figure 4 demonstrates the ZCA fiber FTIR spectrum before and after ERY adsorption. As can be seen, a peak of 600–800 cm⁻¹ was observed, which is associated with T-O-T stretching and T-O zeolite bending (Armaroli et al., 2006).

A sharp peak in such regions indicates the presence of zeolite inside the membrane. In addition, the membrane showed a peak in the region of 1,000–1,200 cm⁻¹, indicating the interaction between Si-O-Si of zeolite and CA.

Some peaks were also detected at 1,735–1,738 cm⁻¹ assigned to carbonyl C=O stretching of CA and broad peak at about 3,400 cm⁻¹ assigned to O-H stretching. Furthermore, the absence of new peaks was observed on the membrane after the adsorption process. However, the peak was slightly shifted and the peak intensity decreased. This might be due to the presence of van der Waals force, indicating the physical adsorption between the metal ions and membrane.
Thermogravimetric Analysis
Thermogravimetric analysis for the three samples, namely, ZCA, cellulose acetate, and zeolite, is presented in Figure 5. From the TGA thermogram obtained for cellulose acetate (CA), there is initially a minor weight loss of about 3% up to 200°C, which is caused from the loss of volatile compounds and the moisture of H₂O that is bound to the hydrophilic (OH) groups that is bonded in the chain of cellulose acetate chains (Hong et al., 2020).

There are two steps of thermal decomposition: the first phase (300–400°C) which refers to the major loss with a proximate weight loss of 75%, while the second one (400 and 600°C) having a weight loss of 15% is referred to the complete degradation and composition (Hong et al., 2020).

Two levels of mass reduction have been found for zeolite. The first stage was between 30 and 230°C, with a weight loss of 40% which can be due to the loss of H₂O adsorbed to the material and to the deterioration of certain aluminum and silicate fractions which did not decompose at 400°C during the pyrolysis process. The second stage of zeolite thermal decomposition, starting at 380°C, was caused from the removal of minerals and salts from the material, which has 35% of its initial mass which is considered as its high mineral residue content.

The ZCA fiber thermogram showed three levels of thermal decomposition between 30 and 200°C, 215 and 380°C, and above 380°C. This thermogram showed an intermediate profile in comparison to the CA and zeolite thermograms; that is, for both of the temperature scales of the thermal events referred to above, their mass variations occurred roughly as the sum of the other two thermograms, because the fiber is made up of 50% of the weight of each part. The first process, with a weight loss of approximately 20%, can be attributed mainly to the release of H₂O from the material due to the presence of zeolite, with the CA mass being practically constant in this temperature range. The second stage of decomposition is probably due to the degradation of the CA chain, with the zeolite mass remaining almost unchanged. The CA mass loss at this stage was 80%. The third and final stage can be due to complete fiber degradation, and part of the fiber has thermal stability lower than CA, with maximum CA losses at 335 and 360°C, respectively.

Differential Scanning Calorimetry
The DSC thermogram obtained for the ZCA is shown in Figure 6. The peaks were shown at different temperatures (180, 211, and 225°C). ZCA melting happened at a temperature lower than that of CA melting as indicated by other studies (Gómez et al., 2006). This may have been explained by the fact, that is, the strengthening as well as a lower amount of contacts between the CA chains. Also, the melting enthalpy was 3,600 kJ/g for ZCA. The higher energy involved during the ZCA melting process may be due to water volatilization, since TGA showed large mass loss in this temperature range.

Adsorption Study
Effect of Contact Time
The effect of equilibrium adsorption time on adsorption efficiency was studied at room temperature close to 25°C. To study that, an initial concentration of ERY of 20 mgL⁻¹ and about 20 mg of ZCA adsorbent were used at different time intervals: 15, 30, 45, 60, 75, 90, and 120 min, as shown in Figure 7. The presence of large number of active sites made the adsorption of
ERY to the surface of the absorbents very easy and increased rapidly at an early stage. This process was followed by a slower rise in adsorption. This shows that the complex derivatives formed at the initial stage of adsorption are unstable, resulting in a rapid rate of adsorption. As a result of the presence of hydrogen protons emitted to the oxygen-containing solution on the adsorbent surface beside the presence of hydroxyl and carboxyl groups, this causes a slower adsorption speeds which could be due to a reduction in the driving force of the present adsorption sites. The various efficiencies of adsorption have shown that the absorbents do not show identical morphologies.

Effect of Temperature
Measurements of adsorption were carried out using an absorbent weight of 20 mg, an initial concentration of 20 mg/L, and a time interval of 60 min. The removal of ERY, controlled by CA, zeolite, and ZCA tests, increased with a rise in temperature from 20 to 45°C, initially indicating an endothermic adsorption mechanism up to 30°C (Figure 8). This could lead to an improvement in the diffusion rate of ERY in the porous structure of the ZCA derivatives, raising the temperature. Due to high temperatures, the adsorption mechanism can include both physical and chemical adsorption, resulting in increased active sites due to bond breakup. The endothermic adsorption process can therefore be attributed to increased pore diameter. Nevertheless, increases in the removal of ERY were controlled with a rise in temperature from 20 to 45°C using CA, zeolite, and ZCA samples, showing a concentration equilibrium between ERY and absorbents.

Effect of ERY Initial Concentration
Measurements of adsorption at room temperature (25°C) were carried out using separate initial ERY concentrations of 10, 20, 30, and 40 mg/L. For 60 min and 20 mg of the three absorbents, as shown in Figure 9. With an increase in the overall ERY content of up to 20 mg/L, the adsorption process improved and then started to decrease.

The number of interchangeable sites in the absorbent structures and the ratio of ERY to the three adsorbents were identified as the main factors for the decline in adsorption as initial ERY concentrations increased. The exchangeable sites on the absorbents are saturated after increasing the ratio of ERY, resulting in a decrease in the efficiency of adsorption. It was observed that the adsorption capacity of absorbents improved by 5% with an improvement in initial ERY concentrations from 10 to 20 mg/L. This may be the result of the substantial driving force transferred by the ERY concentration in order to defeat the resistance to mass movement between solid and liquid phases.

As seen in Table 1, with reference to the previous studies, the innovation of this study can be summarized as using zeolite/cellulose acetate blended fiber as the first example in the ERY removal literature.

**Kinetic Models and Adsorption Isotherms**
In this study, the modeling of adsorption kinetics was studied to help and describe the adsorption rate-controlling mechanism.

We studied the adsorption kinetics of ERY using the three absorbents at initial concentration of 30 mg/L and at 25°C. From

---

**TABLE 1** Previous studies on ERY removal from water.

| Adsorbent | Optimum condition | Percentage removal (%) or adsorption capacity \( (q_m) \) | Reference |
|-----------|-------------------|----------------------------------------------------------|-----------|
| Magnetic activated carbon | Initial ERY concentration of 65 mg L\(^{-1}\), sorbent weight of 1.55 g L\(^{-1}\), the contact time of 76.25 min, and at the temperature of 35°C | 95.125% | Gholamiyan et al. (2020) |
| Magnetic imprinted polymers (MIPs) from chitosan | Initial ERY concentration of 10 mg L\(^{-1}\), and at the temperature of 25°C, \( \text{pH} = 4 \) | Adsorption capacity \( (q_m) = 52.32 \text{μmol/g} \) at 15°C | 99.4% | Ou et al. (2015) |
| Multi-walled carbon nanotubes | Mixing rate of 200 rpm, amount of adsorbent up to 1 g/L, and at the temperature of 75°C | | | |
| Porous magnetic graphene (PMG) | \( \text{pH} = 3 \), contact time of 30 min, initial antibiotic concentration of 200 mg/L, and adsorbent dose of 0.35 g/L | | | |
| Fe\(_3\)O\(_4\)/activated carbon/chitosan (MACC: Magnetic activated carbon/chitosan) | 15 mg adsorbent, and at the temperature of 20°C | adsorption capacity \( (q_m) = 286 \text{ mg/g} \) | 89% | Mostafapour et al. (2018) |
| Amberlite XAD-4 | 0.002 mg adsorbent at 30°C | adsorption capacity \( (q_m) = 526.31 \text{ mg/g} \) | 98% | Fateme et al. (2020) |
| Zeolite/cellulose acetate blend fiber (our study) | Initial ERY concentration of 20 mg L\(^{-1}\), the contact time of 60 min, and at the temperature of 30°C | adsorption capacity \( (q_m) = 358 \text{ mg/g} \) | 98% | Danaioğlu et al. (2017) |

---
In this study, the obtained kinetic data were analyzed with the pseudo-first-order (Radi et al., 2015), pseudo-second-order (Hanbali et al., 2020), and intraparticle diffusion using Eqs 3, 4, 10 respectively.

As seen in Figure 10, pseudo-second-order modeling showed an improved fit for adsorption calculations relative to pseudo-first-order modeling for all adsorbents.

However, the results obtained in pseudo-first-order modeling were still adequate to define the sorption kinetics of ERY, showing that the surface showed both chemisorption and physi-sorption adsorption processes. The regression coefficient ($R^2$) of all adsorbents in the pseudo-second-order is very close to 1 more than the one for the pseudo-first-order. Also, the $q_e$ calculated for the three adsorbents in the pseudo-second-order is very close to the experimental one, as shown in Table 2. It has been shown that the pseudo-second-order modeling showed an acceptable match...
The initial stage (Hong et al., 2020).

operation, which is due to the more accessible adsorption sites at
the

frontal or micropore diffusion. A greater slope of the external transfer of mass or boundary diffusion of the layer
intraparticle diffusion model, one sees two distinct areas,
forms of experimental data. For data that match the
applicability of the intraparticle diffusion model on all three

fronts. Accordingly, the dipole moment (D), 4.421 D, 416.124 au, and

9.600, and 9,600 kcal/mol for zeolite, CA, and ZCA, respectively, and this shows that the
adsorption is exothermic. The positive values of \( \Delta S \) for ERY on the three adsorbents showed some orderliness on the surfaces
adsorbtion process occurred mainly in the monolayer system.

Thermodynamic Study

The adsorption thermodynamics for the adsorption process of
ERY onto three adsorbents are displayed in Table 4, in order to understand the nature of ERY adsorption on the three adsorbents
using Eqs. 7–9. The three adsorbents showed negative values of \( \Delta H \) and the values were \(-6,200, -8,500, \) and \(-9,600 \) kcal/mol for zeolite, CA, and ZCA, respectively, and this shows that the
adsorption is exothermic. The positive values of \( \Delta S \) for ERY on the three adsorbents showed some orderliness on the surfaces
adsorbtion process occurred mainly in the monolayer system.

Quantum Chemical Studies

The optimized geometry and calculated physical and quantum
chemical quantities of ERY at HF/6-311G** level.

\[
\begin{align*}
\text{HOMO} & (-1) (eV) & -9.385 & \text{DM (debye)} & 4.421 \\
\text{LUMO} & (-A) (eV) & 0.372 & \alpha (\text{au}) & 416.124 \\
\Delta E_{\text{gap}} & (L-H) (eV) & 9.757 & \beta (\text{au}) & 169.795 \\
\mu & (eV) & -4.506 & \Delta E (\text{cal/mol}) & -2,467.184 \\
\eta & (eV) & -4.879 & \Delta H (\text{cal/mol}) & -2,467.129 \\
\omega & (eV) & 2.081 & \Delta G (\text{cal/mol}) & -2,467.269 \\
\omega' & (\text{au}) & 0.016 & \Delta E_{\text{vib.thermal}} (\text{cal/mol}) & 728.461 \\
\omega'' & (\text{au}) & 0.182 & \Delta E_{\text{vib.thermal}} (\text{cal/mol}) & 728.461 \\
\Delta N & (eV) & 0.032 & C_v (\text{cal/molK}) & 201.476 \\
\Delta N_{\text{back-donat}} & (eV) & -1.220 & S (\text{cal/molK}) & 294.519 \\
\Delta N_{\text{max}} & (eV) & 0.924 & S (\text{cal/molK}) & 209.940 \\
- & & & S (\text{cal/molK}) & 209.940 \\
\end{align*}
\]

The movement of ERY from aqueous solution to the adsorbents
surfaces might be in different steps, that is, intraparticle diffusion, film diffusion, or both, and that is the rate determining step. The
intraparticle diffusion model is shown.

\[
q_t = K pi t^{1/2} + C_i
\]  (10)

The constant \( C_i \) represents the boundary layer thickness, and \( K pi \) is a constant. A plot between \( q_t \) vs. \( t^{1/2} \) showed straight line with
an appropriate value of correlation coefficient \( R^2 \) giving the
applicability of the intraparticle diffusion model on all three forms of experimental data. For data that match the
intraparticle diffusion model, one sees two distinct areas, meaning that two stages are involved in the diffusion process:
the external transfer of mass or boundary diffusion of the layer
and the intraparticle or micropore diffusion. A greater slope of
the first step than the second step suggests a faster adsorption operation, which is due to the more accessible adsorption sites at
the initial stage (Hong et al., 2020).

Equilibrium Modeling

Both Langmuir and Freundlich isotherms are the most widely
used models for representing equilibrium data of adsorption of
ERY onto three adsorbents that were investigated at 25°C for
30 min, with an adsorbent weight of 30 mg/L (Figure 11).

The equilibrium study was carried out in order to understand
the mechanism of adsorption process, that is, Langmuir (Hanbali et al., 2020) and Freundlich (Brunauer et al., 1938), which
assumes the adsorption of adsorbate as a function of
equilibrium concentration. Langmuir isotherm best describes
the monolayer adsorption of the solute from solution onto the
adsorbent surface having a finite number of active sites present on
it. The linear form of the Langmuir isotherm model is shown
in Eq. 5.

The results of the models are shown in Table 3. A dimensionless constant \( R_l \) was calculated using Eq. 11.

\[
R_l = \frac{1}{(1 + K_l C_i)}.
\]  (11)

where Co is the original concentration of ERY (mg/L) and \( K_l \) is
the constant of Langmuir isotherm. The RL value represents
adsorption mechanisms that are unfavorable (RL > 1), linear (RL = 1), desirable (1 > RL > 0), or irreversible (RL = 0) (Hong et al., 2020). The \( R_l \) (0.106) values for ERY in the present study were <1 for the three adsorbents, which indicated favorable adsorption.

Freundlich isotherm considers the heterogeneous surface and nonuniform distribution of heat of sorption. It is most favorably
studied for description of the multilayer adsorption
process (Eq. 6).

In summary, the studied isotherms were best suited to
Langmuir models, which is believed due to the high regression
coefficient \( R^2 \) value (Table 3). It can also be observed that the
surfaces of all three adsorbents are homogeneous and that
adsorption process occurred mainly in the monolayer system.

TABLE 4 | Adsorption thermodynamics for the adsorption of ERY.

| Parameter | Temperature | Zeolite | CA | Zeolite/CA |
|-----------|-------------|--------|----|------------|
| \( \Delta S \) (J mol\(^{-1}\)) | 20 | 0.08 | 0.06 | 0.03 |
| \( \Delta H \) (kJ mol\(^{-1}\)) | 20 | -6,200 | -8,500 | -9,600 |
| \( \Delta G \) (kJ mol\(^{-1}\)) | 20 | -0.25 | -0.35 | -0.62 |
| 25 | -1.13 | -1.17 | -1.36 |
| 30 | -1.32 | -1.56 | -1.9 |
| 35 | -2.13 | -2.78 | -3.26 |
| 45 | -2.5 | -3.5 | -4.1 |

TABLE 5 | Calculated physical and quantum chemical quantities of ERY at HF/6-311G** level.

| QCP | Physical parameters |
|------|---------------------|
| HOMO (\(-1\)) (eV) | -9.385 | DM (debye) | 4.421 |
| LUMO (\(-A\)) (eV) | 0.372 | \(\alpha\) (au) | 416.124 |
| \(\Delta E_{\text{gap}}\) (L-H) (eV) | 9.757 | \(\beta\) (au) | 169.795 |
| \(\mu\) (eV) | -4.506 | \(\Delta E\) (cal/mol) | -2,467.184 |
| \(\eta\) (eV) | -4.879 | \(\Delta H\) (cal/mol) | -2,467.129 |
| \(\omega\) (eV) | 2.081 | \(\Delta G\) (cal/mol) | -2,467.269 |
| \(\omega'\) (au) | 0.016 | \(\Delta E_{\text{vib.thermal}}\) (cal/mol) | 728.461 |
| \(\omega''\) (au) | 0.182 | \(\Delta E_{\text{vib.thermal}}\) (cal/mol) | 728.461 |
| \(\Delta N\) (eV) | 0.032 | \(C_v\) (cal/molK) | 201.476 |
| \(\Delta N_{\text{back-donat}}\) (eV) | -1.220 | \(S\) (cal/molK) | 294.519 |
| \(\Delta N_{\text{max}}\) (eV) | 0.924 | \(S\) (cal/molK) | 209.940 |

Table 4, in order to understand the nature of ERY adsorption on the three adsorbents using Eqs. 7–9. The three adsorbents showed negative values of \( \Delta H \) and the values were \(-6,200, -8,500, \) and \(-9,600 \) kcal/mol for zeolite, CA, and ZCA, respectively, and this shows that the
adsorption is exothermic. The positive values of \( \Delta S \) for ERY on the three adsorbents showed some orderliness on the surfaces
adsorbtion process occurred mainly in the monolayer system.
were estimated at 207.438 cal/molK and 294.519 cal/molK, respectively, whereas the vibrational part of these quantities were determined at 201.476 cal/molK and 209.940 cal/molK, respectively.

In addition, the QCPs are used successfully to assess the chemical reactivity tendency of ERY. In past, corrosion inhibition efficiency was reported to increase with an increase of the electron-donating ability in case \( \Delta N \) < 3.6, and vice versa for \( \Delta N \) > 3.6 (Lukovits et al., 2001). According to the \( \Delta N \) (0.032 < 3.6) and electron-donating power values, the adsorption of ERY toward the studied adsorbents is easily noticed to be actualized via the charge transfer from the ERY compound to studied adsorbents. Moreover, the possible nucleophilic (HOMO) and electrophilic (LUMO) attack sites of ERY compound are shown in Figure 13. The HOMO density was mostly amplified over the dimethyl amin (-N(CH3)2) functional and partially be scattered on the oxacyclohexane ring. On the other side, the LUMO broadens on the surrounding of -(C=O)-functional group of ERY. In addition, the MEP graphs displayed the richness of the electron by red color (V < 0).

| Table 6 | NBO results for ERY at HF/6-311G** level. |
|---------|------------------------------------------|
| Donor (i) | ED/e  | Acceptor (j) | EDj/e | \( \Delta\text{E}_{\text{gap}} \)/kcal/mol | E(i)-E(j)/a.u | F(i,j)/a.u |
| \( \sigma \text{ C31}-\text{C43} \) | 1.97272 | \( \text{I}^{+} \text{ C12}-\text{C36} \) | 0.04962 | 4.77 | 1.06 | 0.064 |
| \( \sigma \text{ C31}-\text{H72} \) | 1.96794 | \( \sigma \text{ C15}-\text{H17} \) | 0.04194 | 9.47 | 1.08 | 0.091 |
| LP (2) O1 | 1.94227 | \( \sigma \text{ C22}-\text{C26} \) | 0.04328 | 9.84 | 1.11 | 0.093 |
| LP (2) O2 | 1.94036 | \( \sigma \text{ C18}-\text{H54} \) | 0.02444 | 8.56 | 1.18 | 0.090 |
| LP (2) O3 | 1.96470 | \( \sigma \text{ C17}-\text{C19} \) | 0.03290 | 10.64 | 1.12 | 0.097 |
| LP (2) O4 | 1.93923 | \( \sigma \text{ C22}-\text{C26} \) | 0.04338 | 9.30 | 1.11 | 0.091 |
| LP (2) O5 | 1.93444 | \( \sigma \text{ C37}-\text{H77} \) | 0.02965 | 11.13 | 1.15 | 0.102 |
| LP (2) O6 | 1.84800 | \( \sigma \text{ C28}-\text{C30} \) | 0.04813 | 10.84 | 1.10 | 0.098 |
| LP (2) O7 | 1.94152 | \( \sigma \text{ C51}-\text{H117} \) | 0.01658 | 9.50 | 1.12 | 0.093 |
| LP (2) O8 | 1.96521 | \( \sigma \text{ C26}-\text{C28} \) | 0.03160 | 9.21 | 1.11 | 0.090 |
| LP (2) O9 | 1.96742 | \( \sigma \text{ C24}-\text{C30} \) | 0.04813 | 9.41 | 1.13 | 0.092 |
| LP (2) O10 | 1.96687 | \( \sigma \text{ C32}-\text{C35} \) | 0.04718 | 8.51 | 1.10 | 0.087 |
| LP (2) O11 | 1.96259 | \( \sigma \text{ C31}-\text{C35} \) | 0.03166 | 9.18 | 1.10 | 0.090 |
| LP (2) O12 | 1.91457 | \( \sigma \text{ C21}-\text{C36} \) | 0.05236 | 24.58 | 1.10 | 0.148 |
| LP (2) O13 | 1.88472 | \( \sigma \text{ C24}-\text{C30} \) | 0.04813 | 10.84 | 1.10 | 0.098 |
| LP (1) N14 | 1.90376 | \( \sigma \text{ C49}-\text{H111} \) | 0.02255 | 10.74 | 1.07 | 0.097 |
and poorness by blue color (V > 0) fields of the ERY compound. As expected, the charge transfer zeolites are minerals that contain mainly aluminum and silicon compounds—C=O groups were covered by red color to electrophilic attacks, and the H Atom of the -O-H group was marked by blue color for the nucleophilic attacks. Also, the saturated C-chain of ERY presented neutral attitude for both nucleophiles and electrophiles because it is covered by green color.

The chemical reactivity of many kinds of molecular systems (Mustafa and Serdaroglu, 2017; Jacob et al., 2020; Serdaroglu, 2020; Serdaroglu et al., 2020; Al-Otaibi et al., 2021) has also been clarified by using the results of the second-order perturbative energy analysis. Table 6 summarized the lowering of the stabilization energy, possible interaction types, and the occupancies of both donor and acceptor orbitals. As expected, the mainly saturated structure of the ERY compound, the dominant interactions contributed to $E(2)$ (62.33 kcal/mol) were sourced from the charge transfer to anti-bonding orbital $\sigma^*$ O13-C39 (ED$_i = 0.15098e$) from nonbonding orbital LP (2) O6 (ED$_i = 1.84800e$). Also, the hyperconjugations due to the charge movement from each filled orbital $\sigma$ C31-C43 and $\sigma$ C31-H72 to unfilled orbital $\sigma^*$ C12-O36 were calculated with the $E(2)$ of 4.77 kcal/mol and 2.39 kcal/mol, respectively, even if they did not contribute much to the $E(2)$. From Table 6, the remaining interactions were due to the anumeric interactions, and the highest energy interactions among them were predicted as the interaction LP (2) O13 (ED$_i = 1.88472e$) $\rightarrow \sigma^*$ O6-C39 (ED$_i = 0.07615e$) in 42.55 kcal/mol. Similarly, the charge movement from the lone pair of the oxygen atom known as the strong electron-donating of the -C=O group to neighbor orbitals also had great responsibility of energy lowering. Namely, the LP (2) O12 $\rightarrow \sigma^*$ C21-C36 ($E(2) = 24.58$ kcal/mol), LP (2) O12 $\rightarrow \sigma^*$ C31-C36 ($E(2) = 24.89$ kcal/mol), and LP (2) O13 $\rightarrow \sigma^*$ C20-C39 ($E(2) = 24.31$ kcal/mol) interactions also had significant roles in the lowering of the stabilization energy. As known well, the -NH2 group also has strong capability of electron-donating. From Table 6, the charge movement from the N atom of the -NH2 group to each of $\sigma^*$ C28-H67, $\sigma^*$ C48-H108, and $\sigma^*$ C49-H111 interactions was calculated with the energy of 10.72, 10.83, and 10.74 kcal/mol, respectively. Here, it can be considered that these interactions have significant responsibility of the possible intermolecular interactions due to the charge movement that existed in a molecular system, affecting the polarity distribution on the surface.

**Desorption Study**

The stability and reusability of the three adsorbents are especially critical for widespread applications. The adsorption–desorption recycling test, as shown in Figure 14, was used to investigate the adsorbents’ stability further. The adsorbents were washed twice with ethanol after each run and then reused for the next stage of adsorption (Jodeh et al., 2018; Gholamiyan et al., 2020). The findings show that there is no significant loss of adsorption site after three runs, showing that...
the three adsorbents are more reliable. After the first three regeneration cycles, the adsorption efficiency loss of the three adsorbents to ERY was only about 5.04%. The result was attributed to the reduction of the binding sites in imprinted polymer matrix during regeneration cycles (Barrett et al., 1951). Therefore, the three adsorbents can be reused at least three times without significantly decreasing their adsorption capacities.

**CONCLUSION**

With an increase in the number of studies and research on the fate of pharmaceuticals, personal care products, and their environmental effects on human beings, many researches have been published. As the population and economies grow, numerous antibiotics are increasingly being used in bio-manufacturing, livestock farming, and pharmaceutical industries. The QCPs revealed that the adsorption of ERY toward the studied adsorbents actualize via the charge transfer from the ERY compound to studied adsorbents, because of the ΔN (0.032 < 3.6) and electro-donating power values. The MEP plots pointed out that the -C=O groups were covered by red color to electrophilic attacks and the H Atom of the -O-H group was marked by blue color for the nucleophilic attacks. The NBO analysis of ERY indicated that the anomeric and hyper-conjugative interactions have chief responsibility for the possible intermolecular interactions because of the charge movement affecting the polarity distribution on the surface.

The three adsorbents zeolite, cellulose acetate, and ZCA were used to study the removal of ERY from aqueous liquid prepared in the lab. Several characterizations were done on both the adsorbents and ERY including SEM, XRD, FTIR, and TGA. A brief summary of the results was shown in Abstract, and more details about the results were presented in the Results section.

In summary, the three adsorbents showed very high removal efficiency and reached more than 98% using the fiber. Those adsorbents showed very high reusability, and this will save a lot of money and protect environment.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material; further inquiries can be directed to the corresponding authors.

**AUTHOR CONTRIBUTIONS**

SJ: wrote the manuscript, IE: experimental work, OH: editing the manuscript, YM: did most of the plots. GH and SS: financial aid, OD: explored some data. Soheila Gholamiyan: formal analysis and concept creation. Majid Hamzehloo: formal analysis, conceptualization, control, validation, and visualization. Abdolhadi Farokhnia’s responsibilities include systematic analysis, writing—review and editing, conceptualization, supervision, confirmation, and visualization.

**ACKNOWLEDGMENTS**

The authors would like to thank both the Palestinian Water Authority and the Middle East Desalination Research Center (MEDRC) for their help and financial support through the I2 Innovation program. The authors would like to express their gratitude to the Shahid Chamran University of Ahvaz and the University of Tehran for their financial support of this research. The authors would also like to thank Turkey’s Scientific and Technological Research Council (TUBITAK). The High Performance and Grid Computing Center (TR-Grid e-Infrastructure) has been used for all quantum chemical calculations.

**REFERENCES**

Abujaber, F., Zougagh, M., Jodeh, S., Rios, Á., Guzmán Bernardo, F. J., and Rodríguez Martín-Doimeadios, R. C. (2018). Magnetic Cellulose Nanoparticles Coated with Ionic Liquid as a New Material for the Simple and Fast Monitoring of Emerging Pollutants in Waters by Magnetic Solid Phase Extraction. *Microchemical J.* 137, 490–495. doi:10.1016/j.microc.2017.12.007

Al-Otaibi, J. S., Mary, Y. S., Mary, Y. S., Kaya, S., and Serdaroglu, G. (2021). DFT Computational Study of Trihalogenated Aniline Derivative’s Adsorption onto Graphene/fullerene/fullerene-like Nanocages, X12Y12 (X = Al, B, and Y = N, P). *J. Biomol. Struct. Dyn.* 27 (4), 1–14. doi:10.1080/07391102.2021.1914172

Al-Shaalan, N. H., Ali, L., AlOthman, Z. A., Al-Wahaibi, L. H., and Alabdulmonem, H. (2019). High Performance Removal and Simulation Studies of Diuron Pesticide in Water on MWCNTs. *J. Mol. Liquids* 289, 111039. doi:10.1016/j.molliq.2019.111039

Ali, I., Alharbi, O. M. L., Alothman, Z. A., and Alwarthan, A. (2019). Modeling of Fenuron Pesticide Adsorption on CNTs for Mechanistic Insight and Removal in Water. *Environ. Res.* 170, 389–397. doi:10.1016/j.envres.2018.12.066

Ali, I., Alharbi, O. M. L., Alothman, Z. A., and Alwarthan, A. (2018). Facile and Eco-Friendly Synthesis of Functionalized Iron Nanoparticles for Cyanazine Removal in Water. *Colloids Surf. B: Biointerfaces* 171, 606–613. doi:10.1016/j.colsurfb.2018.07.071

Ali, I., Alharbi, O. M. L., Alothman, Z. A., and Badjah, A. Y. (2018). Kinetics, Thermodynamics, and Modeling of Amido Black Dye Photodegradation in Water Using Co/TiO2 Nanoparticles. *Photochem. Photobiol. Sci.* 17 (4), 1–14. doi:10.1038/s41574-018-0080-z

Armaroli, T., Simon, L. J., Digne, M., Montanari, T., Bevilacqua, M., Valtchev, V., et al. (2006). Effects of crystal Size and Si/Al Ratio on the Surface Properties of H-ZSM-5 Zeolites. *Appl. Catal. A: Gen.* 306, 78–84. doi:10.1016/j.apcata.2006.03.030

Arshad, M. S., Khan, U., Sadiq, A., Khalid, W., Hussain, M., Yasseen, A., et al. (2020). Coronavirus Disease (COVID-19) and Immunity Booster green Foods: A Mini Review. *Food Sci. Nutr.* 8 (8), 3971–3976. doi:10.1002/fsn3.1719

Azzaoui, K., Mejdoubi, E., Lamhamdi, A., Jodeh, S., Hamed, O., Berrabah, M., et al. (2017). Preparation and Characterization of Biodegradable Nanocomposites Derived from Carboxymethyl Cellulose and Hydroxyapatite. *Carbohydr. Polym.* 167, 59–69. doi:10.1016/j.carbpol.2017.02.092

Bagaev, A. V., Bukanova, T. V., and Chubarenko, I. P. (2021). Spring Cold Water Intrusions as the Beginning of the Cold Intermediate Layer Formation in the Baltic Sea. *Estuarine, Coastal Shelf Sci.* 250, 107141. doi:10.1016/j.ecss.2020.107141

Barchiesi, M., et al. Presence and Fate of Microplastics in the Water Sources: Focus on the Role of Wastewater and Drinking Water Treatment Plants. *Journal of Water Process Engineering*. 2020: p. 101787.
Ma, H., Li, X., Zhu, C., Chen, F., Yang, T., and Chen, X. (2020). Liberation and Recovery of Cr from Real Tannery Sludge by Ultrasound-Assisted Supercritical Water Oxidation Treatment. J. Clean. Prod. 267, 122064. doi:10.1016/j.jclepro.2020.122064.

Martucci, A., Pasti, L., Marchetti, N., Cavazzini, A., Dondi, F., and Alberti, A. (2012). Adsorption of Pharmaceuticals from Aqueous Solutions on Synthetic Zeolites. Microporous Mesoporous Mater. 148 (1), 174–183. doi:10.1016/j.micromeso.2011.07.009

McLean, A. D., and Chandler, G. S. (1980). Contracted Gaussian Basis Sets for Molecular Calculations. I. Second Row Atoms. Z. 11=18The Journal of Chemical Physics. J. Chem. Phys. 72 (10), 5639–5648. doi:10.1063/1.4439890

McQuarrie, D. A. (1973). Statistical Thermodynamics.

Mery-Araya, C., Lear, G., Perez-Garcia, O., Astudillo-Garcia, C., and Singhal, N. (2019). Using Carbon Substrate as a Selection Pressure to Enhance the Potential of Aerobic Granular Sludge Microbial Communities for Removing Contaminants of Emerging Concern. Bioresour. Technol. 290, 121705. doi:10.1016/j.biotech.2019.121705

Mostafapour, F. K., Dashitzade, M., and Balarak, D. (2018). Adsorption Thermodynamics, Kinetics and Mechanism for the Adsorption of Erythromycin onto Multi-Walled Carbon Nanotubes. J. Pharm. Res. Int. 24(6), 1–11.

Mustafa, E., and Serdaroglu, G. (2011). A DFT Study of Determination of the Reactive Sites of the Acetylcholine and its Agonists: In the Gas Phase and Dielectric Medium. Int. J. Quan. Chem. 111(10): p. 2464–2475.

Removal of Erythromycin From Wastewater by Ultrasound-Assisted Supercritical CO2 Extraction. Res. Chem. Intermediates. 2020. 46(1): p. 961–982.

Serdaroglu, G., Kaya, S., and Touri, R. (2020). Eco-friendly Sodium Gluconate and Trisodium Citrate Inhibitors for Low Carbon Steel in Simulated Cooling Water System: Theoretical Study and Molecular Dynamic Simulations. J. Mol. Liquids. 319, 114108.

Jodeh et al. Removal of Erythromycin From Wastewater

Copyright © 2021 Jodeh, Erman, Hamed, Massad, Hanbali, Samhan, Dagdag, Kaya and Serdaroglu. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.