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## A practical approach to demonstrate the circular economy in remediation of textile dyes using nutraceutical industrial spent

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We used Nutraceutical Industrial Coriander Seed Spent (NICSS), a readily available, cheap, eco-friendly, and ready-to-use material, as an innovative adsorbent for the bioremediation of a bisazo Acid Red 119 (AR 119) dye, which is likely a mutagen from textile industrial effluents (TIE). A laboratory-scale experiment was tailored to demonstrate the framework of the circular economy (CE) in the remediation of textile dyes using Nutraceutical Industrial Spent to align with the principles of sustainability and valorization. An experimental  $q_e$  value of 97.00 mg g<sup>-1</sup> was obtained. For the practicality and effectiveness of the method, a two-level fractional factorial experimental design (FFED) was employed to determine variables that influence the adsorption capacity of NICSS. At optimal settings (pH of 1.4, adsorbent dosage of 6.000 g L<sup>-1</sup>, adsorbent particle size of 96  $\mu$ m, initial dye concentration of 599 mg L<sup>-1</sup>, adsorption duration of 173 min, orbital shaking speed of 165 rpm, and temperature of 35 °C), the maximum adsorption efficiency achieved through statistical optimization was 614 mg g<sup>-1</sup>. Six factors influencing the adsorption process were examined experimentally and were considered important for commercialization. Three orders of magnitude were applied to the identified variables in scaling experiments. Adsorption-equilibrium data were analyzed using nine isotherm models. The best fit was discovered to be the Vieth-Sladek adsorption isotherm model. The suitable mechanism for the overall rate of the adsorption process was a pseudo-second-order reaction: mass-transfer mechanistic studies were predicted to predominate over the diffusion process. NICSS was characterized using SEM and FTIR spectroscopy. Utilizing plastic trash, the dye-adsorbed NICSS that was recovered as "sludge" was utilized as a reinforcing material to create composites. Dye-adsorbed NICSS thermoplastic and thermoset composites were studied and compared with NICSS composites in terms of their physicomechanical and chemical properties.

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### 1. Introduction

The "circular economy" (CE) has become essential to the sustainability agenda as the world grapples with the need for more sustainable practices. The CE presents an economic approach focused on minimizing/eliminating waste while promoting the optimal use or complete reuse of resources. It

emphasizes the significance of embracing a holistic view of products and processes. Many countries are embracing practices aligned with the principles of the CE to enhance industrial productivity, competitiveness, and efficient use of resources.<sup>1</sup> Achieving circularity is a daunting challenge that requires a systemic approach.<sup>2</sup> However, the latest Circular Gap Report paints a worrying picture: adoption of the CE has stalled, and

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material usage has declined yearly.<sup>3</sup> One of the major reasons is that the CE is perceived as expensive. CE is a resource-intensive economy,<sup>4</sup> so it is desirable to demonstrate the principles of CE to all leading industries. Most countries have adopted rules on management of plastic waste, e-waste, construction, and demolition waste, as well as metals-recycling policies, to promote the CE. However, little attention has been paid to management of textile industrial effluents (TIE). This article explores how to drive change in this critical area of TIE management with simple laboratory experiments.

The United Nations has acknowledged water security to be one of the 17 goals for sustainable development.<sup>5</sup> Sustainable development aims to optimize economic growth by severing ties with wastewater discharge and water usage.<sup>6</sup> Water, therefore, represents a significant concern, hindrance, and avenue for the sustainable growth of enterprises that produce effluents that contain high concentrations of harmful and hazardous materials. According to the World Resources Institute, an estimated 5 trillion liters of water are used in the dyeing process, and an estimated 48–144 billion square meters of fabric from factory waste end up in landfills each year. Overall, the industry is responsible for 20% of the world's water pollution, which is enough water to quench the thirst of 110 million people for an entire year.<sup>7</sup> Furthermore, the textile industry dumps effluent straight into waterways due to lax laws in many developing nations, gravely harming the ecosystem and environment.<sup>8</sup> As a result, metrics for measuring the sustainability of textile production are crucial.

The sustainable index prioritizes the control of effluents by taking into account the effects of pollution on human health resulting from the manufacture of textiles.<sup>9</sup> The significance of waste treatment and water management in the textile industries has been highlighted by these shifts. The latter is seen as one of the major issues facing the modern economy, and the idea of valorization attempts to address it.<sup>10,11</sup> The process of transforming trash into components for additional use, or valorization, adds value above the initial cost of transformation. It draws attention to procedures and activities that lessen emissions and their negative effects on the environment. In contrast to linear economics, which prioritizes end-of-life considerations,<sup>12</sup> the CE envisions the idea of recycling trash from one process and using it as a resource for another.<sup>13</sup> To achieve the status of a sustainable economic system, both approaches make an effort to divorce economic growth from natural resources through the processes of reduction, reuse, recycling, and return.<sup>14</sup> Therefore, to change laboratory research in line with the most recent ideas, valorization and CE have been postulated.

Azo dyes are reactive dyes, representing about one-half of all the dyes in common use. They are synthetic compounds made from two aromatic amines joined by an azo bond. They are characterized by chemical groups that can form covalent bonds with textile substrates. They are synthesized by diazotization and coupling. Most azo dyes are toxic. Azo dyes are non-biodegradable and harmful to the environment. Some azo dyes release aromatic amines and generate carcinogenic metabolites when metabolized in the body.<sup>15</sup>

Azo dyes are widely utilized in the textile and related industries.<sup>16</sup> Due to their mutagenic and carcinogenic qualities, these substances are prohibited worldwide. Even though they are harmful, their proven benefits have made them indispensable to the textile industry. These include: (i) easy and affordable ways to synthesize them in aqueous media; (ii) a vast array of starting materials available; (iii) a wide spectrum of shades; (iv) high intensity and superior fastness of color; (v) versatility in applications on various substrates; (vi) an energy-saving dyeing process at 60 °C compared with the boiling temperature of its counterparts.<sup>17</sup> However, these colors are released as TIE because there are not enough appropriate disposal methods for the  $\sim 4.50 \times 10^5$  tonnes of dyes produced annually worldwide.<sup>18</sup> Thus, remediation of toxic dyes from TIE remains a daunting task.

AR119 is an azo dye. The published techniques for removal/remediation/decolorization of AR119 can be broadly categorized as biological,<sup>19</sup> chemical,<sup>20</sup> and physical.<sup>21,22</sup> Biological techniques cannot be used to break down AR119 dye, which is intended to withstand microbial breakdown and continue to be a stable, long-lasting colorant. The financial viability of the enterprises and sustainability of the environment are compromised by the high operational costs associated with treating and disposing of significant volumes of chemical sludge through the employment of chemical treatment methods.<sup>20</sup> Adsorption, a physical process, has been used to remediate AR119 with sewage sludge and ash.<sup>21,22</sup> Despite being efficient, the adsorption method has drawbacks, such as high investment, high operational costs, low efficiency, as well as the disposal problem of recyclability for real application in wastewater decontamination. These methods have serious limitations in the disposal of the dye-adsorbed sludge because they involve the Environmental Impact Factor (E-factor), and have a carbon footprint.<sup>23</sup> Because of these shortcomings, scholars have chosen to employ Nutraceutical Industrial Coriander Seed Spent (NICSS) as a superior and innovative adsorbent material for the bioremediation of AR119 from TIE. NICSS has seven main advantages: (i) it is readily available; (ii) it is environmentally friendly; (iii) it is abundant; (iv) it is low-cost; (v) it offers a workable solution for the valorization of Nutraceutical Industrial Spent (NIS); (vi) it presents opportunities for the creation of low-cost green composites using plastic waste and dye-adsorbed spent as filler material; (vii) NICSS has a competitive advantage over reported agriculture waste.

The market for nutraceuticals is expected to reach US\$ 336 billion by 2023, growing at a compound annual growth rate of 8%.<sup>24</sup> Nonetheless, a substantial 50–95% of waste is produced during the processing and manufacturing of the main components and/or active substances from nutraceuticals. Given the economics of nutraceuticals, it is possible that the total amount spent (for which data are not available) could reach millions of metric tonnes.

Coriander belongs to the Apiceae family and genus *Coriandrum* (*Coriandrum sativum* L.). With a yearly production of  $3.15 \times 10^5$  tonnes, India holds a larger share of the global export market for this spice than any other country, making it the top producer, consumer, and exporter.<sup>25</sup> NICSS is a by-product



**Table 1** Comparison of maximum adsorption capacities of various natural adsorbents

Adsorbent	Maximum adsorption capacity $q_e$ (mg g <sup>-1</sup> )	Reference
Maple sawdust	3.19	26
Hazelnut shell	16.23	27
Jute	17.18	28
Tree fern	39.80	29
Grape stalk waste	49.90	30
Black grain husk	49.97	31
Coriander seed spent (present study)	97.00	—

obtained by mechanical, thermal, and chemical methods that extract oleoresins as the main component. Thus, NICSS is not a useful feed or fertilizer. Currently, it is used as fuel with a low E-factor. A comparison of maximum adsorption capacities of various natural adsorbents is shown in Table 1.

The present study explores the porous nature of NICSS as an outstanding, ready-to-use biosorbent for the remediation of AR119, a toxic dye from TIE. Furthermore, little information is accessible on the elimination of colors from aqueous water and/or industrial effluents *via* adsorptive remediation and the appropriate disposal of the dye-adsorbed biosorbent ("sludge"). Our Research School has made innovative attempts to create composites using NIS, a filler material<sup>32–34</sup> and as adsorbent for the removal of harmful dyes.<sup>35–39</sup> Scant literature is available on the use of the CE concept in textile industries.<sup>37</sup>

The current work focused on the treatment of textile effluents employing the adsorption technique using NICSS. One of the main goals of this research was to build composites using plastic waste and dye-adsorbed NICSS, a waste material, as a filler and reinforcement material to fit into the concept of the CE.

## 2. Experimental

### 2.1. Preparation and characterization of the adsorbent

**2.1.1. Adsorbent preparation.** NICSS was purchased from a nearby factory that extracts oleoresin from coriander seeds. The waste was sun-dried, ball-milled, ground, and sieved following ASTM specifications.

**2.1.2. Adsorbent characterization.** The apparent composition of NICSS was scanned and visualized using an electronic microscope (LEO 435 VP; Japan). The functional groups included in the adsorbent were recognized using FTIR spectroscopy. Both AR119-loaded samples and NICSS before adsorption had their infrared spectra measured using an FTIR spectrometer (Inter-spec 2020; Spectro Lab, UK). The point of zero charge, or  $pH_Z$ , was firm enough to evaluate the NICSS surface charge. A stock solution of 0.1 M KCl was generated for the  $pH_Z$  determination, and 50 mL of this solution was divided among seven distinct 250 mL Erlenmeyer flasks. Initially, HCl and NaOH were recycled to modify the pH of the solutions between 2.0 and 12.0; 0.05 g of NICSS was added to each solution. A pH meter (Systronics-802; India) was cast-off to

determine the final pH of the solutions after these flasks were left for 24 h. The factors influencing the adsorption efficiency include pH, adsorbent dosage, adsorbent particle size, initial dye concentration, contact time, and temperature.

### 2.2. Studies on composites

**2.2.1. Preparation of AR119 dye-adsorbed NICSS.** First, 100 g of commercial AR119 dye was applied to a 100 L barrel. Ten litres of TIE were used to dissolve the colour. A 20 mm plastic rod was used to manually mix the fluid after 5 kg of commercial NICSS had been transferred. With periodic stirring, the solution was left for ~24 h. Using a cloth, the dye-adsorbed NICSS was separated, and the precipitate was extensively cleaned with distilled water until the filtrate was nearly colourless. Air drying was used to absorb the blue dye from the NICSS. The resulting lump-containing powder was crushed, sieved through a 177  $\mu$ m mesh, and allowed to dry for 24 h at 60 °C in an oven. The powder was placed in an airtight-lidded container and allowed to cool. "Dye-modified NICSS powder" (dm-NICSS) was the term used to describe AR119-dye adsorbed NICSS.

**2.2.2. Preparation of composites.** Several procedures were used to prepare thermoplastic bio-composites of polypropylene (polymer matrix), NICSS (filler material), and dm-NICSS (filler material). Reliance (India) provided polypropylene (PP) (H110MA) (MFI = 11.0 g/10 min). For 12 h, NICSS and dm-NICSS were oven-dried at 100 °C. As a coupling agent, maleic anhydride-grafted-PP, or MAg-PP, was employed. The PP/dm-NICSS and PP/NICSS thermoplastic composites were created in three steps. First, the PP resin was dry-blended in a tumble mixer with weights of 10, 20, 30, 40, and 50% (w/w) NICSS and dm-NICSS. Then, master batches were melted together. Second, the polymer matrix, coupling reagent, and filler material were combined in a co-rotating, fully intermeshing twin-screw extruder. Third, after being gathered and allowed to cool, the extrudates were ground into pellets. Dumbbell examples of the granular blends were created using injection moulding, and their physicomechanical and chemical characteristics were examined.

Using 2% (v/v) methyl ethyl ketone as a catalyst, thermoset composites of unsaturated polyester resin (USP) and dm-NICSS and NICSS in ratios of 2, 5, 10, 15, and 20% (w/w) were created. A straightforward process was chosen. After the liquid had been stirred to create a homogenised slurry, it was carefully placed in a glass frame that met the necessary measurements. We let the slurry dry in air. A pressure plate was applied to the resulting thermoset for ~3 h. A sample with the necessary dimensions was used to examine the chemical and physicomechanical characteristics of the material. Relevant ASTM procedures were used to prepare thermoplastic and thermoset composites to examine their properties: ASTM D 570-98, ASTM D 638-95, ASTM D 792-00, and ASTM D 2240.

## 3. Results and discussion

### 3.1. Surface characterization of the adsorbent

Surface characterization of NICSS was done through SEM, and exhibited a slightly porous structure (Fig. 1). Upon adsorption



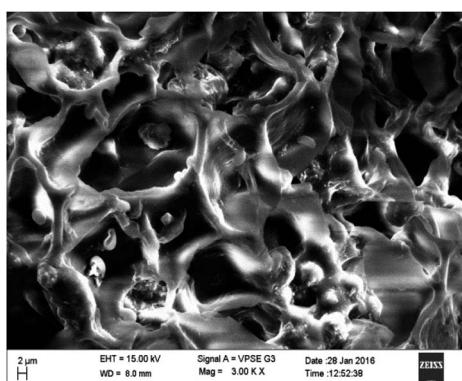


Fig. 1 SEM of NICSS.

(Fig. 2) of AR119 dye, some pores were filled with the adsorbate, forming a thin film over the particle. The IR spectra of NIFGS (Fig. 3) suggested that the absorption at  $3339\text{ cm}^{-1}$  was due to the stretching of hydroxyl groups of cellulose, hemicellulose, and lignin present in NICSS. The bands at  $2924\text{ cm}^{-1}$  and  $2851\text{ cm}^{-1}$  were due to the C-H stretching of saturated carbon.

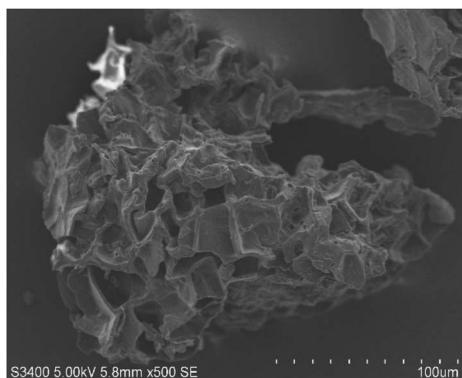


Fig. 2 SEM of AR119-NICSS.

The bands at  $1621$  and  $1775\text{ cm}^{-1}$  were due to C=O stretching. The bands at  $1390\text{ cm}^{-1}$ ,  $1607\text{ cm}^{-1}$ , and  $2923$  were assigned to C-O-C, C-O, and CH vibrations, respectively. The fingerprint region of the biological material NICSS indicated complex bands, and accurate assignment of the peaks was difficult in the absence of a standard sample. However, the presence of cellulose and lignin indicated that the spent material had hydrophilic and hydrophobic properties.

### 3.2. Effect of pH and the initial dye concentration

The pH of the solution affects the adsorption capability of NICSS. First, the pH affects the properties of the adsorbent surface. Second, the pH affects the chemistry of the dye solution. In order to validate the effectiveness of the adsorbent under investigation and scale it up to commercial levels, the parameter pH is crucial. A positively charged bisazo dye, AR119, will be present at lower pH, aiding in the adsorption of NICSS, a cellulose substance rich in -OH groups. The positively charged dye eventually loses its positive nature as the pH rises, which makes the dye adsorb less readily. The curve shape in Fig. 4 was in line with the anticipated chemistry.

The initial concentration of AR119 dye had a significant impact on the adsorption capability of NICSS. The findings are shown in Fig. 5. The curve form indicated that the starting concentration of the dye in the range under study had little bearing on the ability of the adsorbent (NICSS) to remove the adsorbate (AR119) according to percent removal. When the design is changed to improve the commercial viability of the technique, this observation becomes even more crucial.

### 3.3. Effect of adsorbent dosage

Due to its ability to determine the economic viability of the process, the dose parameter will also have a significant impact on commercialization of the process. The adsorbent dosage range of  $0.025\text{--}0.300\text{ g}$  ( $0.500\text{--}6.000\text{ g L}^{-1}$ ) was examined. The removal capacity of the dye affected only the lower

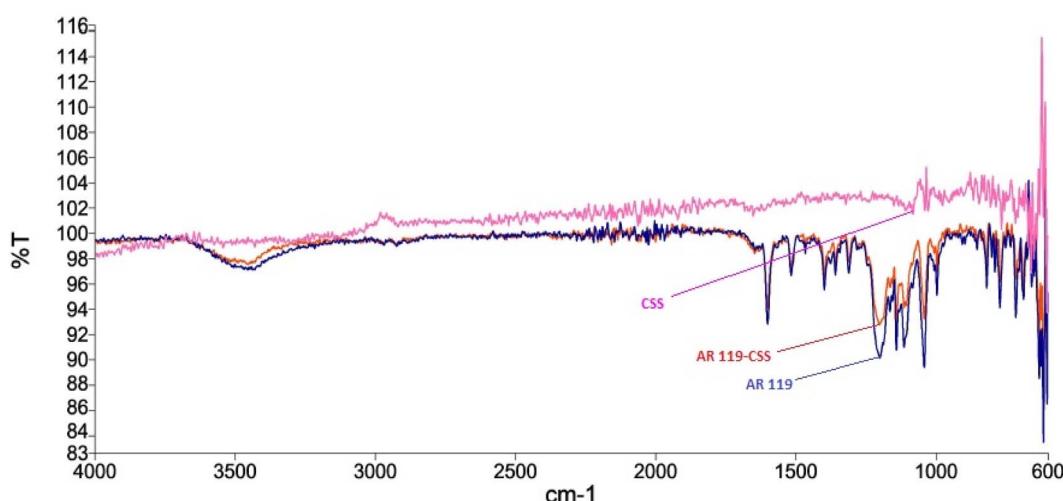


Fig. 3 FTIR spectroscopy ranges of AR119 dye, NICSS and AR119 dye adsorbed on NICSS.



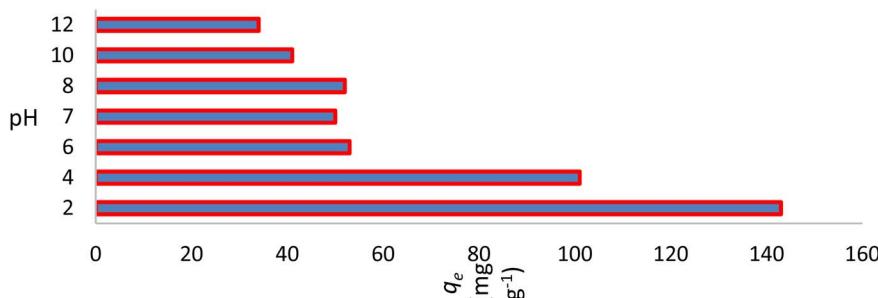


Fig. 4 Effect of pH.

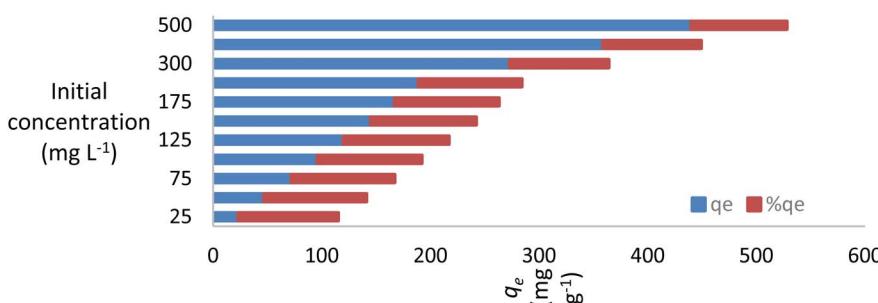


Fig. 5 Effect of the initial dye concentration.

concentrations (0.500 to 1.000 g L<sup>-1</sup>) and remained nearly constant up to 6.000 g L<sup>-1</sup>. Increasing the number of trials with minimal amounts of the adsorbent significantly increased the removal efficiency of the dye by NICSS, an observation that is likely of scientific and economic importance (Fig. 6).

### 3.4. Effect of adsorbent particle size

At neutral pH, the maximal adsorption capability of the AR119 dye was discovered at a preliminary concentration of 100 mg L<sup>-1</sup> (Fig. 7). This observation aligns with the theory that, as particle size increases, and surface area decreases. The extent

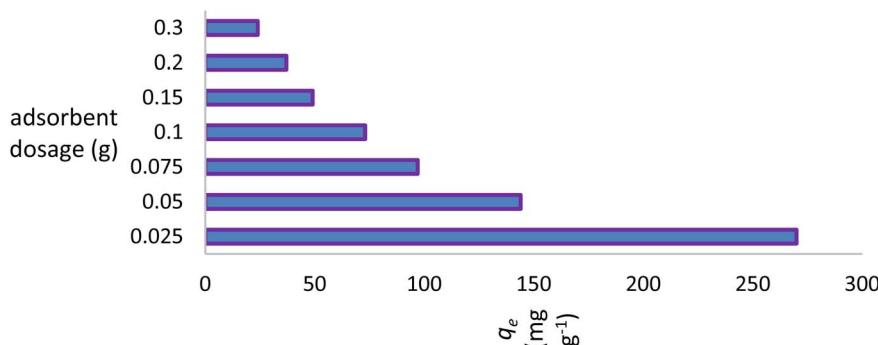


Fig. 6 Influence of the amount of adsorbent used.

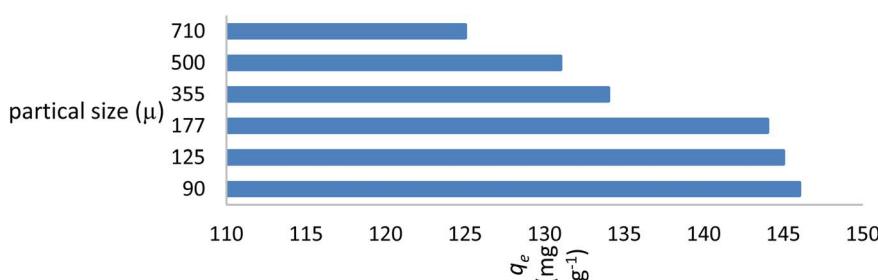


Fig. 7 Impact of the size of adsorbent particles.



range of  $\geq 125 \mu\text{m} \leq 177 \mu\text{m}$  (80-mesh ASTM) was chosen for additional research because particles of this size range are frequently used in the fabrication of composites, and screening particles with smaller sizes requires more time and money.

### 3.5. Effect of contact time

Adsorption is a commonly employed separation technique in the removal of hazardous and toxic materials from industrial effluents. Using kinetic models, one can analyse adsorption kinetics by tracking the uptake of the dye by the adsorbent over time under constant pressure and starting dye concentration. Fig. 8 shows how contact time affected the adsorption of AR119 dye onto NICSS. Based on those data, we concluded that the adsorption process proceeded nearly spontaneously and that the dye was removed with an efficiency of  $\sim 80\%$  of the rapid adsorption seen in the first 60 min. However, a modest increase in adsorption was seen as the contact period increased to 90 min, after which it gradually achieved equilibrium in  $\sim 180$  min.

### 3.6. Effect of temperature

The dye-adsorption process of NICSS as a function of solution temperature was tested, keeping in mind our goal of scaling up to commercial applications. Fig. 9 shows how temperature affected the adsorption of AR119 dye onto NICSS. It may be concluded from the data that the adsorption process was

essentially unaffected by the temperature range under study ( $30^\circ\text{C}$ – $50^\circ\text{C}$ ).<sup>40</sup>

### 3.7. Adsorption isotherms

The parameters of adsorption isotherms reflect the interactions of adsorbate molecules with adsorbent surfaces. This is achieved through Langmuir and Freundlich isotherm models by predicting the adsorption capacity. The Langmuir isotherm model implies that adsorption is a monolayer process that occurs on the surface of the adsorbent, and it involves a finite number of similar adsorption sites processing uniform energies.<sup>41</sup> In the Langmuir isotherm model, the separation factor  $R_L$  has a crucial role.<sup>42</sup> The calculated  $R_L$  values, between 0.052 and 0.214, remained constant, indicating favorable and nearly linear AR119 dye adsorption on NICSS.

The Freundlich isotherm model suggests that the adsorption process takes place on a heterogeneous surface.<sup>43</sup> However, from Langmuir and Freundlich models, no definite inference was obtained for the homogeneity or heterogeneity of the process of adsorption. Therefore, the Jovanovic isotherm<sup>44</sup> model was attempted (Table 2 and Fig. 10).

Redlich–Peterson, Brouers–Sotolongo, Vieth–Sladek, Toth, Sips, and Radke–Prausnitz were the six other three-parameter isotherm models we tried for academic interest. The Redlich–Peterson isotherm model<sup>45</sup> is an improved version of the Langmuir–Freundlich isotherm model, and a correction factor

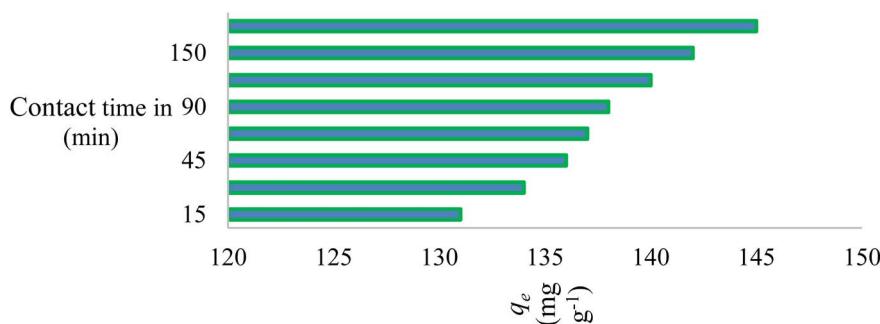


Fig. 8 Influence of exposure time on dye adsorption.

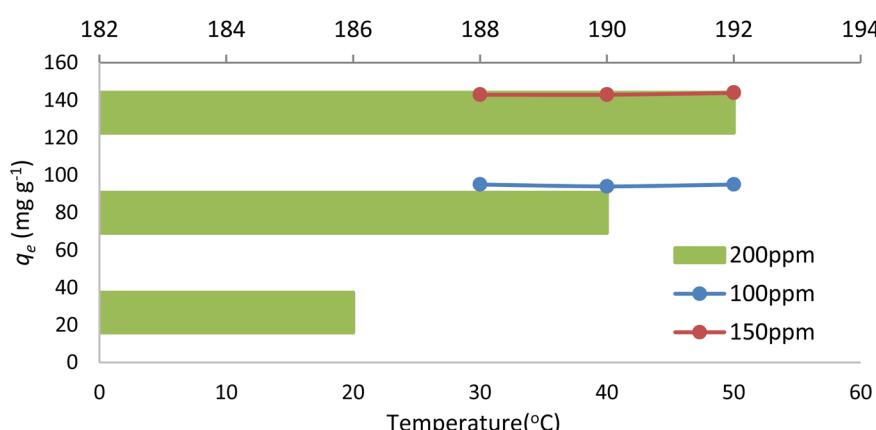


Fig. 9 Impact of temperature on dye adsorption.



Table 2 Intended limitations for isotherms involving two parameters

Langmuir	Freundlich		Jovanovic	
$Q_m$	607.65	$K_F$	46.62	$Q_m$
$K_S$	0.037	$n_F$	1.831	$K_J$
				0.042

(“g”) is included. The Brouers–Sotolongo isotherm<sup>46</sup> is similar to the Vieth–Sladek isotherm. The KBS and  $\alpha$  in the equation represent the adsorption power and active-site distribution, respectively, of the adsorbent–adsorbate system (Fig. 11).

The Vieth–Sladek isotherm<sup>47</sup> is useful for such solutes that are adsorbed because some specific isotherms simplify linear components (Henry's law) and nonlinear components (Langmuir equation) according to the solute dissolved in the amorphous region of the adsorbent. The Toth isotherm<sup>48</sup> is useful for heterogeneous adsorption systems. The Sips isotherm,<sup>49</sup> depending on the adsorbate concentration, can be either the Langmuir isotherm equation or Freundlich isotherm equation

(Fig. 12). The Radke–Prausnitz isotherm<sup>50</sup> experimentally observed the  $Q_m$  value, and the expected  $Q_m$  value was not near the trial  $q_e$  value. These models discuss more complicated equations, but they help make the mechanism of adsorption clearer. The value of  $R^2$  alone could not be considered for our experiment because it can be applied only to linear models.  $\chi^2$  values, however, are applicable only if model data and experimental data are similar. The  $Q_m$ ,  $\chi^2$ , and  $R^2$  values of all nine models are presented in Tables 3 and 4. The values of all these models, as well as the actual experimental data ( $q_e$ ), could provide useful guidance for further research to develop new models for solving adsorption phenomena occurring in the AR119–NICSS system.

### 3.8. Adsorption kinetics

The pseudo-second-order model proposed by Ho and McKay had the best fit at all the dye concentrations and temperatures investigated, as evidenced by the coefficient of determination ( $R^2$ ) and chi-square ( $\chi^2$ ) values. The AR119 dye molecules moved

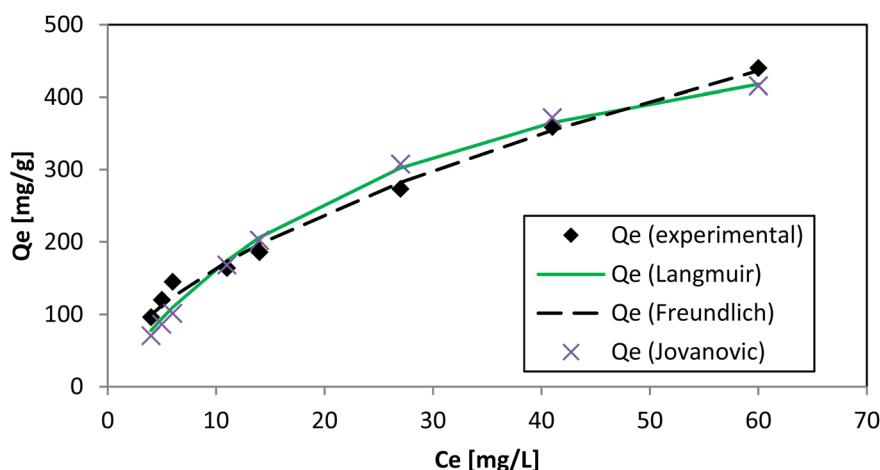


Fig. 10 Langmuir, Freundlich, and Jovanovic adsorption isotherms.

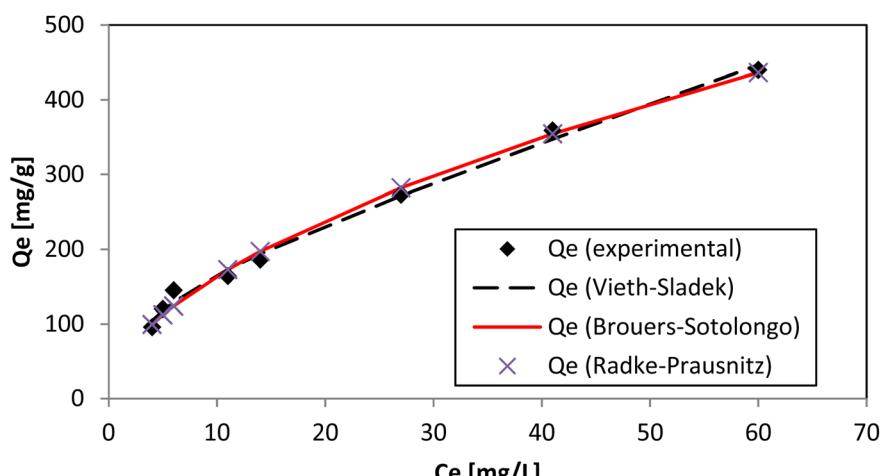


Fig. 11 Adsorption isotherms of Vieth–Sladek, Brouers–Sotolongo, and Radke–Prausnitz.

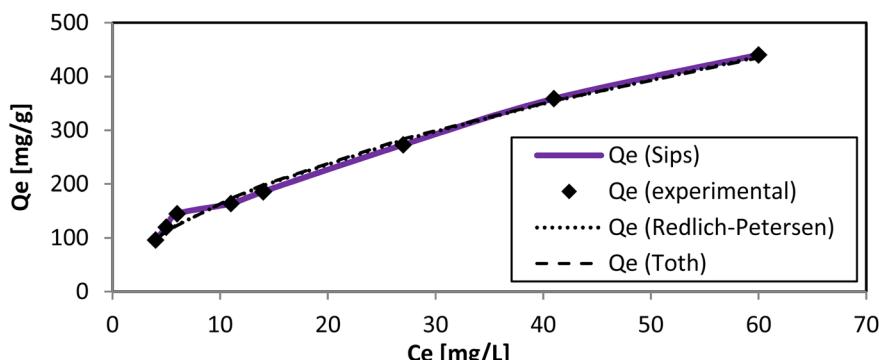


Fig. 12 Sips, Redlich–Petersen, and Toth adsorption isotherms.

Table 3 Designed constraints for isotherms with three parameters

Redlich–Peterson	Toth	Sips	Vieth–Sladek	Brouers–Sotolongo	Radke–Prausnitz
$A_{RP}$	17190.5	$Q_m$	7 679 696	$Q_m$	38298.3
$B_{RP}$	368.076	$n_{TO}$	0.076	$K_s$	$4.61 \times 10^{-6}$
$g$	0.454	$b_{TO}$	1.508	$m_s$	0.546

Table 4 Numerical limitations for fitting isotherm models

Isotherms	Langmuir	Freundlich	Jovanovic	Radke–Prausnitz	Redlich–Peterson	Toth	Sips	Vieth–Sladek	Brouers–Sotolongo
SSE	4163.6	825.8	5927.3	826.2	826.8	957.0	826.2	650.0	825.9
$\chi^2$	24.937	5.159	37.125	5.161	5.164	5.823	5.158	3.818	5.159
$R^2$	0.97	0.99	0.96	0.99	0.99	0.99	0.99	0.99	0.99

from the solution to the solid surface and then diffused into the pores of NICSS. Thus, it may be inferred that multi-step adsorption, which is not rate-limiting, had arisen. For kinetic investigations, three concentrations of AR119 dye were used: 100, 150, and 200 ppm. The difference in the adsorption rate at different temperatures was provided by kinetics studies conducted at 303 K, 313 K, and 323 K. The pseudo-first-order model,<sup>51</sup> pseudo-second-order model,<sup>52</sup> intra-particle diffusion (Weber–Morris) model,<sup>53</sup> Dumwald–Wagner model,<sup>54</sup> and film diffusion model<sup>55</sup> (Table 5) were used as non-linear methods to

examine the adsorption kinetic data (Microsoft Excel® 2010). Table 6 displays the estimated parameters.

When testing initial AR119 dye concentrations of 100, 150, and 200 ppm at different temperatures, the pseudo-second-order model aligned more closely with the experimental results compared with the pseudo-first-order model (Fig. 13–15). After reaching maximum adsorption, the initially very fast adsorption rate gradually slowed down to become stagnant. The temperature increased along with the adsorption capacity ( $q_e$ ). These findings suggested no rate limitation caused by adsorption

Table 5 Theoretical and experimental determinations of model parameters

Initial concentration [ppm]	Temp. [K]	$q_{e,\text{expt}}$ [mg g <sup>-1</sup> ]	Pseudo-first order			Pseudo-second order		
			$Q_{\text{mpred}}$ [mg g <sup>-1</sup> ]	$k_1$	$R^2$	$\chi^2$	$Q_{\text{mpred}}$ [mg g <sup>-1</sup> ]	$k_2$
100	303	95	89.62	$2.53 \times 10^{-1}$	0.66	0.22	92.91	$8.27 \times 10^{-3}$
	313	94	92.11	$2.85 \times 10^{-1}$	0.82	0.05	94.31	$1.24 \times 10^{-2}$
	323	95	93.00	$2.53 \times 10^{-1}$	0.84	0.09	96.00	$8.81 \times 10^{-3}$
150	303	143	106.13	$7.81 \times 10^{-2}$	0.78	4.46	126.44	$7.51 \times 10^{-4}$
	313	143	118.07	$1.18 \times 10^2$	0.85	4.77	65.88	$1.71 \times 10^{-3}$
	323	144	123.42	$6.21 \times 10^{-2}$	0.85	4.97	152.08	$4.50 \times 10^{-4}$
200	303	186	174.55	$3.09 \times 10^{-1}$	0.65	0.15	178.24	$7.71 \times 10^{-3}$
	313	190	176.19	$2.91 \times 10^{-1}$	0.66	0.21	180.60	$6.36 \times 10^{-3}$
	323	192	183.77	$2.49 \times 10^{-1}$	0.50	0.92	191.51	$3.57 \times 10^{-3}$



Table 6 Planned restrictions for diffusion representations

Initial concentration [ppm]	Temp. [K]	Film diffusion model		Weber–Morris model		Dumwald–Wagner	
		$R^1$ [min $^{-1}$ ]	$R^2$	$k_{ist}$ [mg g $^{-1}$ s $^{-0.5}$ ]	$R^2$	$K$ [min $^{-1}$ ]	$R^2$
100	303	0.0277	0.99	1.98	0.99	0.027	0.99
	313	0.0395	0.93	1.29	0.89	0.039	0.93
	323	0.0452	0.92	1.72	0.87	0.044	0.92
	303	0.0192	0.98	10.06	0.98	0.015	0.96
	313	0.0275	0.96	12.27	0.99	0.023	0.94
	323	0.0320	0.95	12.96	0.99	0.027	0.93
150	303	0.0584	0.89	2.23	0.96	0.015	0.95
	313	0.0158	0.99	2.71	0.99	0.015	0.99
	323	0.0584	0.89	4.77	0.99	0.059	0.89
	303	0.0192	0.98	10.06	0.98	0.015	0.96
	313	0.0275	0.96	12.27	0.99	0.023	0.94
	323	0.0320	0.95	12.96	0.99	0.027	0.93
200	303	0.0584	0.89	2.23	0.96	0.015	0.95
	313	0.0158	0.99	2.71	0.99	0.015	0.99
	323	0.0584	0.89	4.77	0.99	0.059	0.89

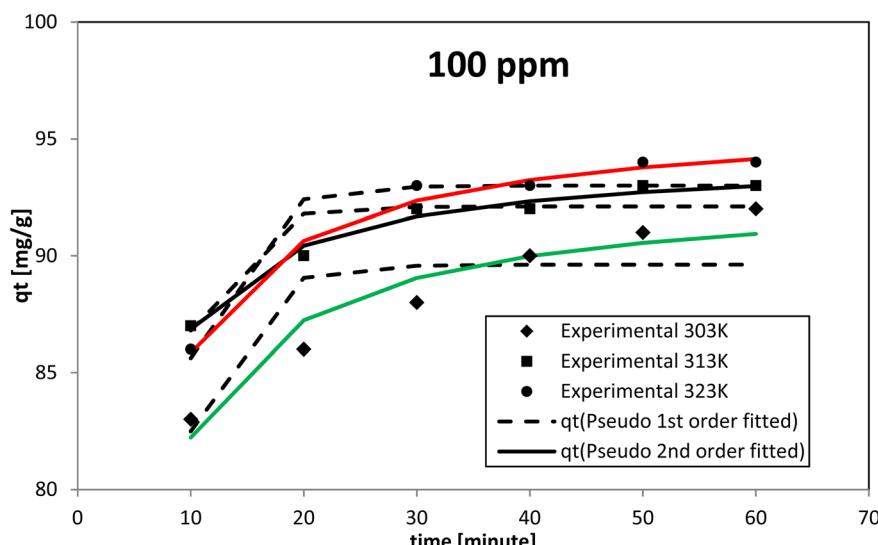


Fig. 13 Adjustments of kinetic models to an initial concentration of 100 ppm.

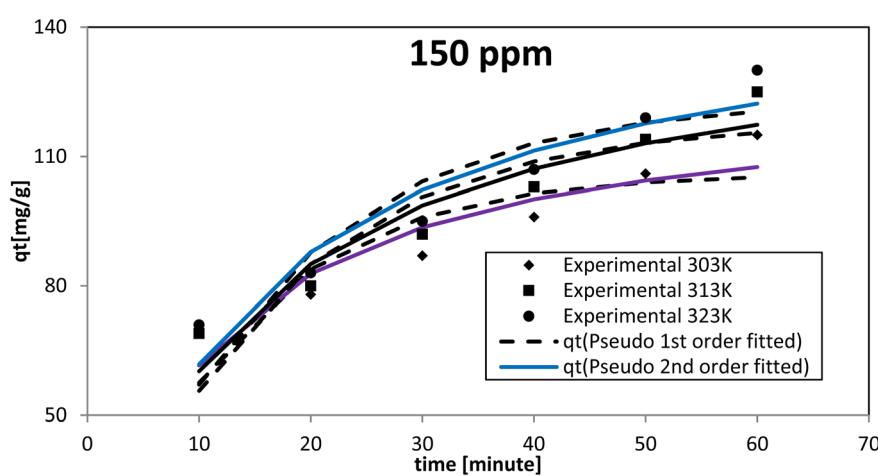


Fig. 14 Adaptations of kinetic models for a starting concentration of 150 ppm.

processes. Additionally, this information indicated that in a multi-stage adsorption process, solute particles transitioned from the bulk solution to the solid surface, and then proceeded to diffuse into the pores of NICSS.

The Dumwald–Wagner model was employed to determine the true rate constant of absorption ( $K$ ) (Fig. 16), which incorporates adjustments for experimental diffusion (Table 6). Our experimental data clearly showed numerous levels of linearity at



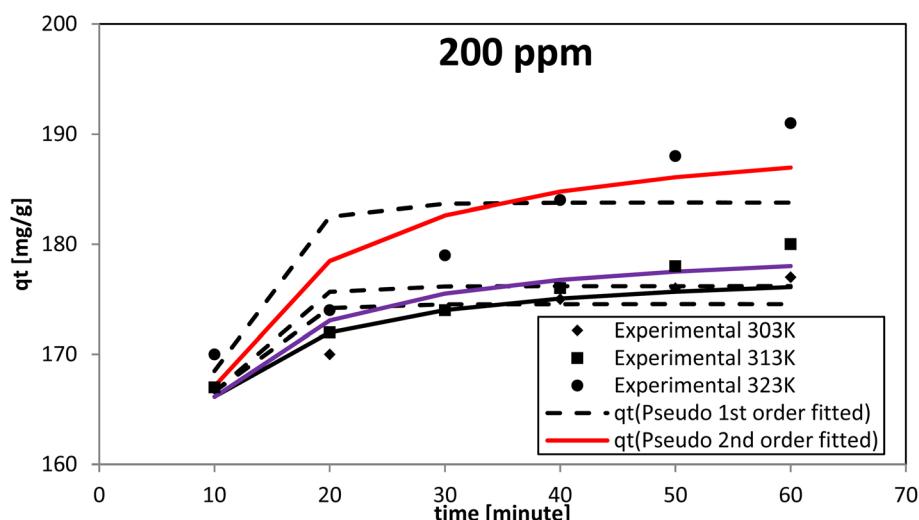


Fig. 15 Fits of kinetic models for an initial concentration of 200 ppm.

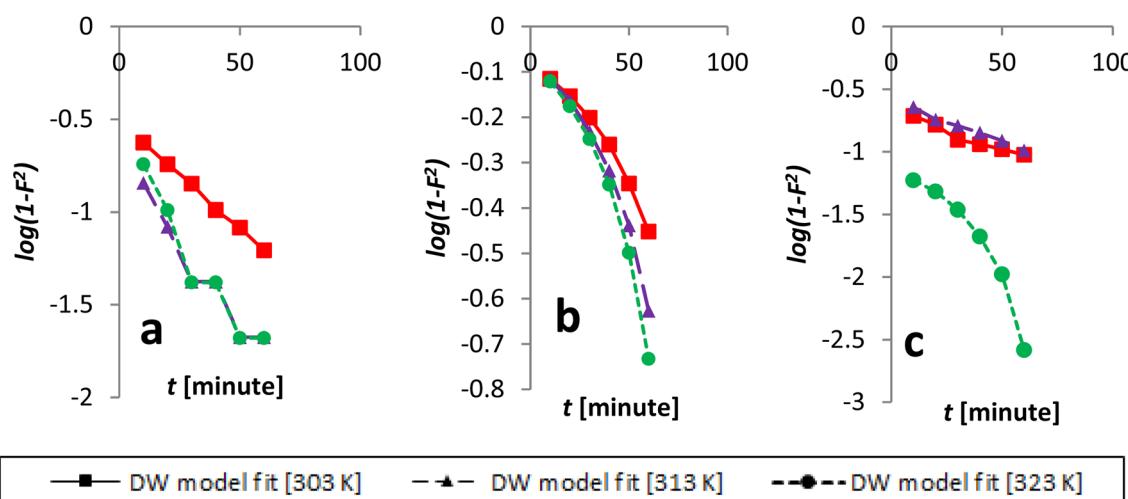


Fig. 16 Kinetics data, beginning with an initial concentration of AR119, was modeled according to the Dumwald–Wagner model: (a) 100 ppm, (b) 150 ppm and (c) 200 ppm.

all solute concentrations, as per the Weber–Morris model (Fig. 17). The adsorption rate was high at first (100 ppm) and lower temperatures. After that, it shifted and took a different linear trajectory before eventually stabilising with time. At higher temperatures, the rate became more linear. This was especially true after fitting the film diffusion model with data at higher temperatures (Fig. 18 and Table 6). These data suggested that the rate obstruction caused by adsorption was negligible due to diffusional constraints at higher temperatures. Hence, diffusion is a mechanism that limits rate. The solute is first absorbed rapidly onto the particle surface, generating a coating that subsequently slows down further diffusion and alters the absorption rate.

### 3.9. Adsorption thermodynamics

The primary variables of the communication method are energy and entropy. The spontaneity of adsorption is signified by the

change in Gibbs free energy ( $\Delta G^\circ$ ). Adsorption was efficient when the change in free energy ( $\Delta G^\circ$ ) related to the adsorption process was negative (Fig. 19 and 20). Table 7 provides estimations of the thermodynamic parameters.

### 3.10. Statistical optimization using a reduced factorial experimental framework

Tests were conducted using various combinations of the six independent factors to analyze their individual and combined impacts. The variance analysis of the quadratic regression model, as detailed in Table 8, highlighted the significance of the singular and interactive impacts of these factors. There was reasonable agreement between the modified  $R^2$  value of 89.6% and projected  $R^2$  value of 70.7%. The high coefficient of variance (16.2%) and  $R^2$  value of 91.0% of the model ensure that it could be utilized to navigate the design space. The expected reactions and experimental results showed a strong connection, as seen



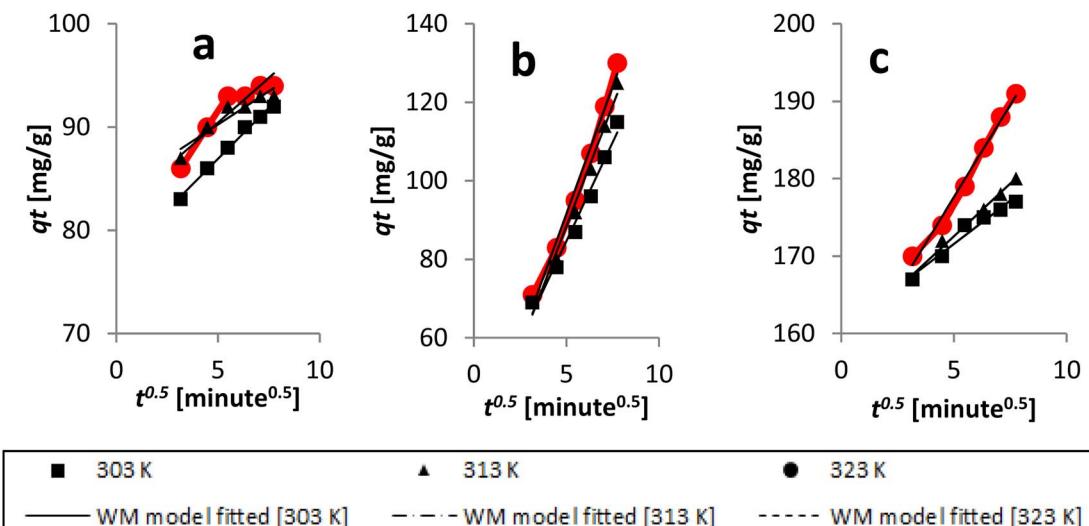


Fig. 17 Kinetics data, with an initial concentration of AR119, were analyzed using the Weber–Morris model: (a) 100 ppm, (b) 150 ppm and (c) 200 ppm.

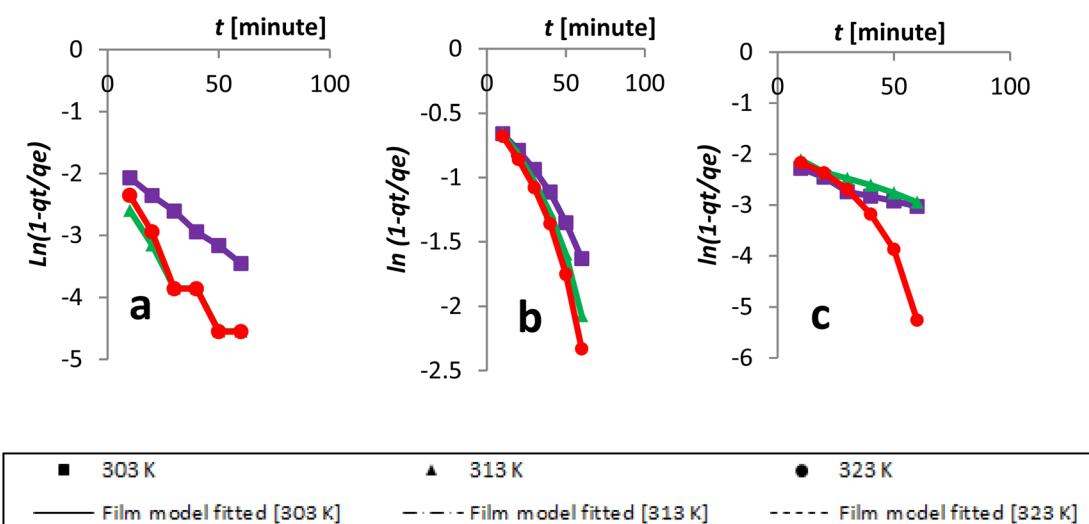


Fig. 18 Kinetics data, with an initial concentration of AR119, were modeled according to the film diffusion model: (a) 100 ppm, (b) 150 ppm and (c) 200 ppm.

in the comparison graph of actual *versus* predicted values (Fig. 21). The following equation displays the regression equation that was derived from the investigation:

$$\begin{aligned} \text{Adsorption} = & -15.1 + 28.0 \times A + 4.6 \times B + 203.1 \\ & \times C - 12.6 \times D - 75.7 \times E - 53.4 \times F - 0.3 \\ & \times AB + 8.8 \times AC + 8.6 \times BC - 22.0 \times A^2 + 3.7 \\ & \times B^2 - 4.2 \times C^2 + 0.6 \times D^2 + 88.6 \times E^2 + 39.6 \times F^2 \end{aligned}$$

By applying the second-order polynomial equation, optimal variable values could be achieved by maximizing the interaction terms derived from multiple regression analysis, as suggested by the factorial fixed effects design (FFED). At the optimal settings, which included a pH of 1.4, an adsorbent dosage of

6.000 g L $^{-1}$ , an adsorbent particle size of 96  $\mu\text{m}$ , an initial dye concentration of 599 mg L $^{-1}$ , an adsorption duration of 173 min, and an orbital shaking speed of 165 rpm at a temperature of 35 °C, the maximum adsorption efficiency achieved through statistical optimization was 614 mg g $^{-1}$ . The final phase of the investigation entailed generating a 3D response surface and contour plots to analyze the effects of two independent variables within the statistical optimization process.

Fig. 22–25 show 3D graphs revealing the relationship between time and temperature (AB), concentration (AC), adsorbent particle size (AD), adsorbent dosage (AE), and pH (AF), where A is time, B is temperature, C is concentration, D is adsorbent particle size, E is adsorbent dosage, and F is pH. Time was shown to have a beneficial effect on adsorption ability.

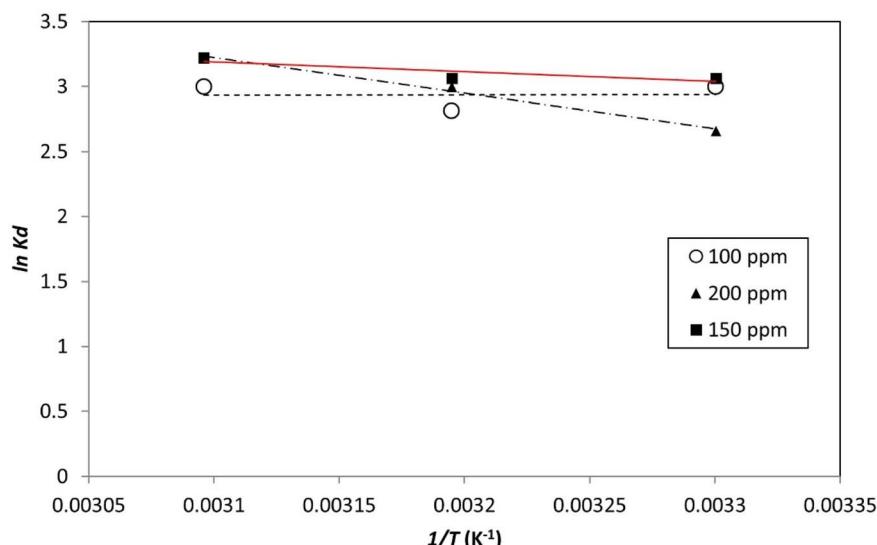


Fig. 19 Plotting the thermodynamic equilibrium constant *versus* the reciprocal of temperature ( $1/T$ ), the enthalpy and Gibbs free energy of the process was determined.

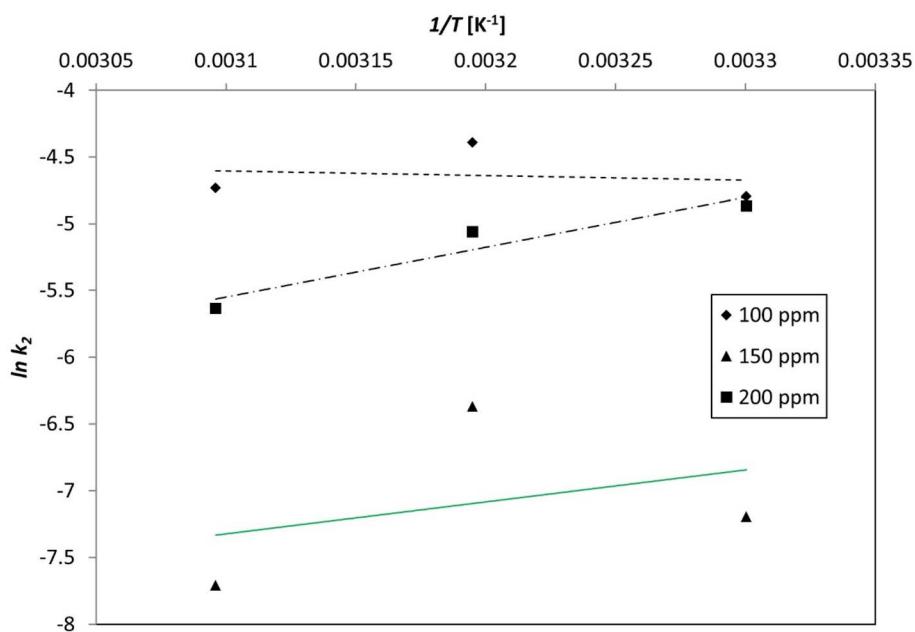


Fig. 20 Plotting the pseudo-second-order kinetic constant as a function of the inverse of temperature ( $1/T$ ) unveiled the activation energy of the process.

Adsorption capacity was positively impacted by time according to 3D graphs that plotted time against all other variables. We could speed-up the adsorption process by increasing the duration in addition to the temperature, particle size, and dye concentration. The “ideal” duration for the adsorption process was 173 min, and increasing the period further was found to enhance adsorption. Adsorption capacity was positively impacted by temperature increases. Adsorption capacity increased in tandem with temperature over time.

It has been noted that a rise in the starting concentration of a dye boosts the adsorption capacity. A particle size of 96  $\mu$ m

was the maximum for adsorption; at larger sizes, the reactivity decreased. Higher initial dye concentrations led to increased levels of adsorption. The maximum pH at which the enhanced adsorption capacity can occur was around 1 and, as time increased, so did the sorption capacity. Positive numbers denote an incremental effect; for instance, a substantial increase in sorption capacity was caused by a temperature increase ( $B$ ). Similarly, a negative number denoted a negative effect; in the case of adsorption dose ( $D$ ), this meant that a rise in adsorption dosage resulted in a fall in the adsorption capacity of the adsorbent. Therefore, the surface and contour



Table 7 Thermodynamic limitations of the AR119-NICSS method

Initial concentration [ppm]	Temperature [K]	$\Delta G^\circ$ [kJ mol <sup>-1</sup> ]	$\Delta S^\circ$ [J mol <sup>-1</sup> K <sup>-1</sup> ]	$\Delta H^\circ$ [kJ mol <sup>-1</sup> ]	ln A	$E_a$ [kJ mol <sup>-1</sup> ]
100	303	-7.55	198.67	-1.31	-3.52	24.08
	313	-7.32				
	323	-8.05				
150	303	-7.72	380.26	51.57	-14.74	-165.41
	313	-7.97				
	323	-8.64				
200	303	-6.69	810.74	189.63	-17.16	-258.96
	313	-7.79				
	323	-8.64				

Table 8 ANOVA<sup>a</sup>

Source	Sum of squares	Degree of freedom	Mean square	F value	P-value
Model	380042.7	15	25336.2	63.9	<0.001**
A	4515.5	1	4515.5	8.4	0.0011**
B	115.2	1	115.2	0.3	0.5913
C	78941.2	1	78941.2	199.0	<0.001**
D	650.8	1	650.8	1.6	0.2034
E	24492.9	1	24492.9	61.7	<0.001**
F	13544.5	1	13544.5	34.1	<0.001**
AB	1.9	1	1.9	0.0	0.9443
AC	221.0	1	221.0	0.6	0.4573
BC	100.9	1	100.9	0.3	0.6152
$A^2$	4373.8	1	4373.8	8.0	0.0013**
$B^2$	175.0	1	175.0	0.4	0.5082
$C^2$	31.4	1	31.4	0.1	0.7792
$D^2$	0.4	1	0.4	0.0	0.9750
$E^2$	8991.8	1	8991.8	22.7	<0.001**
$F^2$	2812.7	1	2812.7	7.1	0.0091**
Residual	37689.6	95	396.7		
Total	417732.3	110			

<sup>a</sup> Significant figures. + suggestive significance ( $0.05 < p < 0.10$ ). \* moderately significant ( $0.01 < p \leq 0.05$ ). \*\* strongly significant ( $p \leq 0.01$ ).

plots (Fig. 22–25) graphically depict the combined outcome of two limitations on biosorption.

It has been discovered that the quadratic model created for process optimization is useful for understanding the relationship between independent variables and how it affects the adsorption method, as well as for forecasting the maximum adsorption capacity. Adsorption was amplified from  $420 \text{ mg g}^{-1}$  to  $614 \text{ mg g}^{-1}$ , a rise of  $>46.2\%$ , as a result of statistical optimization.

### 3.11. Characterization of composites

Thermoplastic bio-composites comprising polypropylene and unsaturated polyester thermoset composites were investigated for their physicomechanical characteristics. Furthermore, an assessment was conducted on the impact of ageing on the tensile strength and chemical resistance of unsaturated polyester thermoset composites. Tables 9–12 provide more specific results.

According to a study on the physicomechanical characteristics of thermoplastic polypropylene composites, adding more NICSS or dm-NICSS filler increased the tensile modulus,

reduced elongation at break, and increased tensile strength. However, when compared with plain PP, flexural characteristics significantly improved. Table 9 illustrates how the inclusion of hydrophilic NICSS affected the water-adsorption properties by causing a weight increase.

Our findings indicate that thermoset composites made of dmNICSS/NICSS and USP had superior dimensional stability compared with USP. Additionally, it verified enhanced resistance to all chemical reagents examined, with the exception of sodium hydroxide (10% w/v). On the other hand, the tensile strength declined as the filler content increased, most likely as a result of lower interfacial adhesion and decreased interaction between the filler material and polymer matrix.

### 3.12. Application of the proposed method to textile industrial effluent

A simple procedure was developed to compare the remediation process for AR119 dye in water as well as in textile industrial effluent, and these are explained elsewhere.<sup>39</sup> The detailed procedure involved nine steps. Step 1 involved 500 mL of solution 1 being transferred to a 1 L conical flask. Then, 5 g of



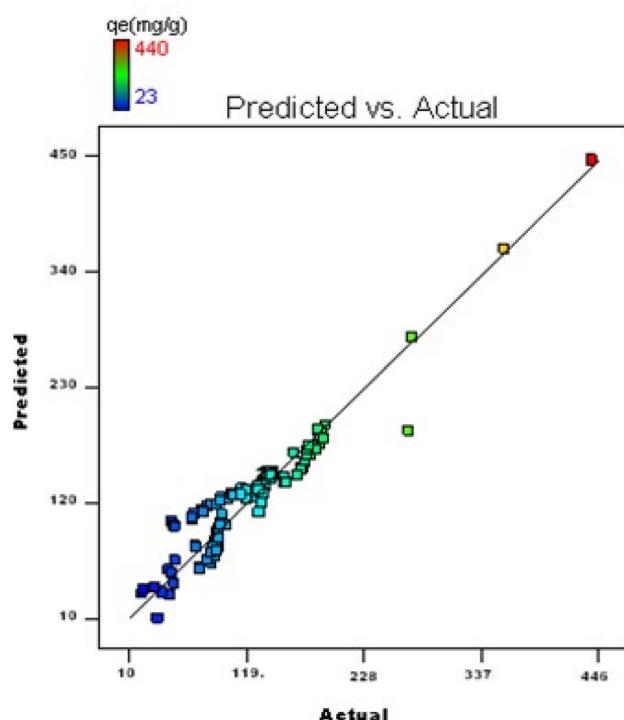


Fig. 21 Comparison of real and expected values.

NICSS was added to the conical flask and the solution was agitated at  $\sim 700$  rpm using magnetic stirrer. At the end of 15 min, agitation was stopped and solution was passed through number-42 Whatman filter paper using the Büchner funnel apparatus. If the filtrate was not clear, the filtration was repeated and absorbance recorded. For step 2, the dye-adsorbed NICSS on the filter paper of step 1 was carefully transferred from a Büchner funnel to a watch glass and kept in an oven at  $60$   $^{\circ}\text{C}$  for 24 h for drying. The dried powder was scraped using a spatula and transferred to a watch glass. For step 3, the second portion of 5 g NICSS was added to the conical flask containing

the filtrate solution from step 1, and agitation was continued on a magnetic stirrer at  $\sim 700$  rpm for 15 min. At the end of 15 min, the agitation was stopped and the solution was passed through number-42 Whatman filter paper using the Büchner funnel apparatus. If the filtrate was not clear, the filtration was repeated and absorbance recorded. In step 4, the dye-adsorbed NICSS on the filter paper of step 3 was carefully transferred from a Büchner funnel to a watch glass and kept in an oven at  $60$   $^{\circ}\text{C}$  for 24 h for drying. The dried powder was scraped using a spatula and transferred to a watch glass. In step 5, the third portion of 5 g NICSS was added to the conical flask containing filtrate solution from step 3 and agitation was continued on a magnetic stirrer at  $\sim 700$  rpm for 15 min. At the end of 15 min, agitation was stopped and the solution was passed through number-42 Whatman filter paper using the Büchner funnel apparatus. If the filtrate was not clear, the filtration was repeated and absorbance recorded. At step 6, the dye-adsorbed NICSS on the filter paper of step 5 was carefully transferred from a Büchner funnel to a watch glass and kept in an oven at  $60$   $^{\circ}\text{C}$  for 24 h for drying. The dried powder was scraped using a spatula and transferred to a watch glass. For step 7, the fourth portion of 5 g NICSS was added to the conical flask containing the filtrate solution from step 5 and agitation was continued on a magnetic stirrer at  $\sim 700$  rpm for 15 min. At the end of 15 min, agitation was stopped and the solution passed through number-42 Whatman filter paper using the Büchner funnel apparatus. If the filtrate was not clear, the filtration was repeated and absorbance recorded. At step 8, the dye-adsorbed NICSS on the filter paper of step 7 was carefully transferred from a Büchner funnel to a watch glass and kept in an oven at  $60$   $^{\circ}\text{C}$  for 24 h for drying. The dried powder was scraped using a spatula and transferred to a watch glass. At step 9, steps 1 to 8 were repeated with solution 2.

This procedure was also adopted using NICSS as an adsorbent. Recovery of the dye and allied substances to the extent of 75%, 85%, 94%, and 96% after 15, 30, 45, and 60 min, respectively, was obtained. This observation was in line with the

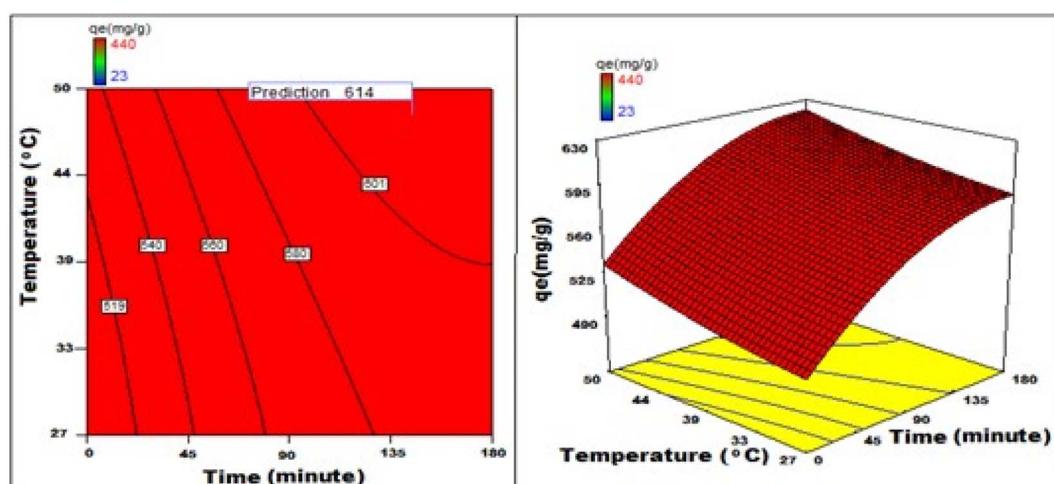


Fig. 22 Fluctuations in the capacity for adsorption over time as influenced by pH.



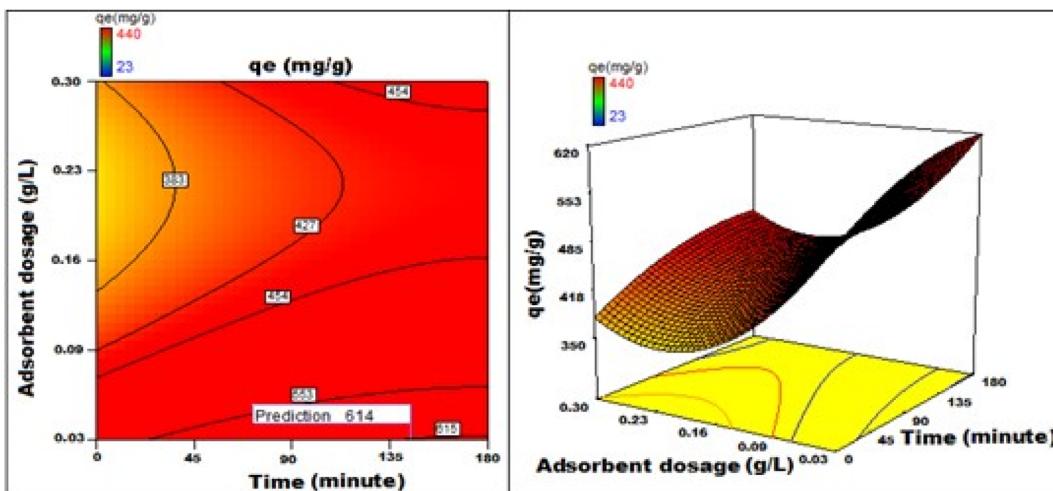


Fig. 23 Change in adsorption capacity over time as a function of the amount of adsorbent used.

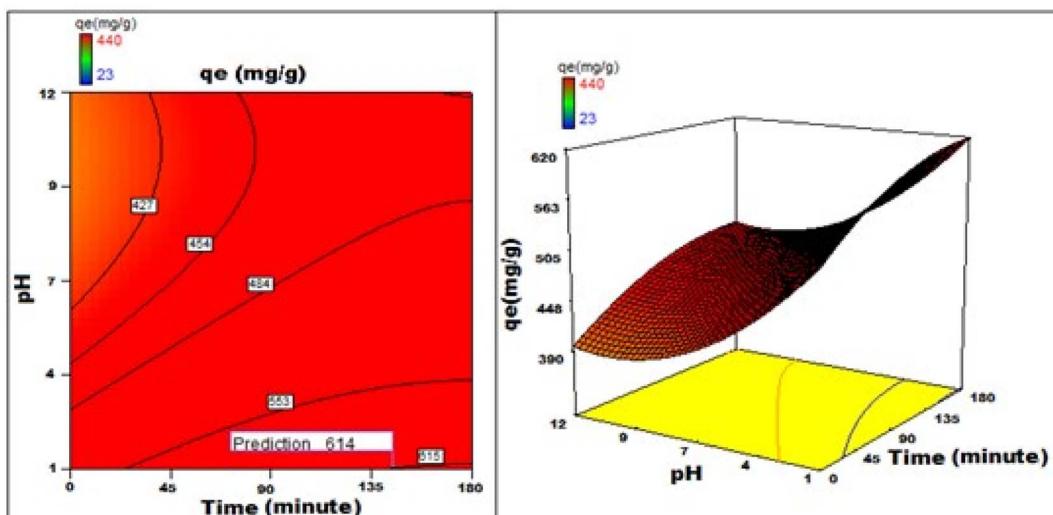


Fig. 24 Variation in the adsorption capability with time in connection to pH.

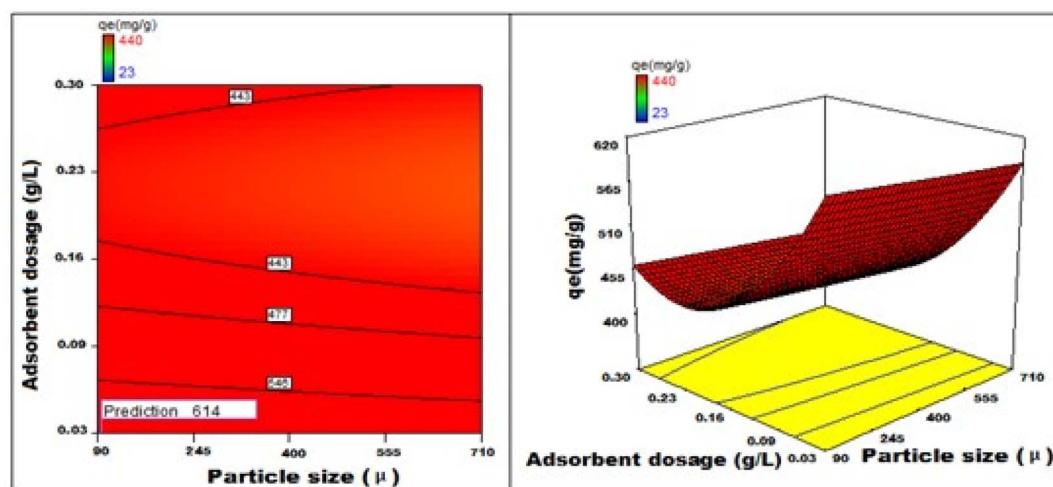


Fig. 25 Fluctuations in adsorption capability correlated with the size of the particles and quantity of adsorbent deployed.



Table 9 Physicomechanical properties of thermoplastic polypropylene composites<sup>a</sup>

Properties	Percent composition of polymer matrix and filler material											
	PP : NICSS						PP : dm-NICSS					
	100 : 00	90 : 10	80 : 20	70 : 30	60 : 40	50 : 50	100 : 00	90 : 10	80 : 20	70 : 30	60 : 40	50 : 50
Tensile strength (MPa)	30.2	29.2	28.1	26.3	23.9	18.2	30.3	29.3	28.2	25.9	22.8	19.1
Tensile modulus (MPa)	1035	1354	1543	1758	1722	1613	1035	1358	1532	1747	1739	1634
Tensile elongation at break (%)	151	12.7	9.6	5.7	3.3	2.6	151	12.9	9.9	5.4	3.4	2.3
Flexural strength (MPa)	32.8	49.5	51.7	53.6	55.2	NRR	33.8	50.7	52.7	55.1	55.6	NRR
Flexural modulus (MPa)	821	492	1582	1747	2089	NRR	821	1126	1569	1782	2142	NRR
Density (g cm <sup>-3</sup> )	0.894	0.923	0.983	1.009	1.053	NRR	0.899	0.923	0.981	1.009	1.057	NRR
Surface hardness (shores D)	65	69	71	74	79	NRR	65	69	70	73	81	NRR
Water absorption in 48 h (%)	0.009	0.09	0.23	0.75	2.62	NRR	0.009	0.09	0.25	0.76	2.82	NRR

<sup>a</sup> NRR: Non-reproducible results.

Table 10 Effect of aging on the tensile strength of unsaturated polyester thermoset composites

Properties	Percentage composition									
	USP : NICSS					USP : dm-NICSS				
	100 : 00	95 : 05	90 : 10	85 : 15	80 : 20	100 : 00	95 : 05	90 : 10	85 : 15	80 : 20
<b>Density (g mL<sup>-1</sup>)</b>										
Experimental	1.214	1.216	1.223	1.217	1.234	1.214	1.214	1.223	1.226	1.227
Theoretical	—	1.217	1.225	1.235	1.236	—	1.217	1.225	1.235	1.236
Surface hardness (shores) ± 2	88.5	89.7	91.3	0.93	1.02	88.5	90.1	90.9	91.5	92.1
Void content (%)	—	0.42	0.84	0.89	1.00	—	0.40	0.81	0.91	1.06
Specific tensile strength (kN m kg <sup>-1</sup> )	36.5	25.1	24.6	20.7	18.6	36.5	25.3	24.7	20.7	18.4

Table 11 Physicomechanical properties of unsaturated polyester thermoset composites

Properties	Percent composition										
	USP : NICSS						USP : dm-NICSS				
	100 : 00	90 : 10	80 : 20	70 : 30	60 : 40	50 : 50	100 : 00	90 : 10	80 : 20	70 : 30	60 : 40
Density (g cm <sup>-3</sup> ) [experimental]	1.213	1.216	1.223	1.217	1.234	1.213	1.214	1.224	1.226	1.227	1.216
Density (g cm <sup>-3</sup> ) [theoretical]	—	1.216	1.225	1.235	1.236	—	1.216	1.225	1.235	1.236	1.217
Surface hardness (shores) [±2]	88.5	89.7	91.3	0.93	1.02	88.5	90.1	89.9	91.5	92.1	89.7
Void content (%)	—	0.42	0.84	0.90	1.00	—	0.40	0.81	0.91	1.06	0.42
Specific tensile strength (kN m kg <sup>-1</sup> )	36.5	25.1	24.6	20.7	18.6	36.5	25.3	24.7	20.7	18.4	25.1

kinetic results, where the solute is adsorbed onto the surface, quickly forming a film. Later on, the adsorption is retarded due to the formation of the film, thereby causing a change in absorption rates. The scaling up of the experiment was done using 10 g, 20 g, and 50 g of NICSS and using 1, 2 and 5 L of AR119 dye dissolved in TIE. In all cases, the results did not exceed ±2% error. The scale-up experimental data yielded promising and reliable results about the process, which could be extended to a higher scale to operate in an industrial environment. The powder and filtrate solutions after the adsorption of constituents of the solution on NICSS are shown in Fig. 26 and 27.

### 3.13. Regeneration of the adsorbent and cost analysis

Dye-adsorbed NICSS could be regenerated for reuse along with the recovery of the adsorbed NICSS. However, the proposition may not prove economical because the process and solvent costs will be much higher when compared with the recovered adsorbent cost (<1 US\$ for 50 kg of NICSS). In addition, enhancement of E-factor<sup>23</sup> is not desirable because it will add to the unprecedented load of environmental toxicants. However in our research team, work is in progress to use dye-adsorbed nutraceutical industrial spent as filler/reinforcing material, and encouraging results have emerged, some of which have been published elsewhere.<sup>35</sup>



Table 12 Studies on the chemical resistance of unsaturated polyester thermoset composites

Chemical reagents	Percentage change in weight after seven days								
	Neat USP		USP : NICSS				USP : dm-NICSS		
	100 : 00	95 : 05	90 : 10	85 : 15	80 : 20	95 : 05	90 : 10	85 : 15	80 : 20
Water	1.07	2.15	3.13	3.35	4.94	2.16	3.15	3.41	4.95
10% (v/v) acetic acid	0.26	0.37	0.35	0.38	0.47	0.31	0.34	0.37	0.47
10% (v/v) hydrochloric acid	0.31	0.36	0.45	0.47	0.51	0.35	0.43	0.46	0.47
10% (v/v) nitric acid	0.35	0.40	0.59	0.65	0.67	0.39	0.57	0.63	0.65
10% (v/v) ammonium hypochlorite	0.65	0.72	0.77	0.80	0.84	0.70	0.76	0.79	0.83
10% (v/v) sodium hydroxide	2.87	3.33	4.35	5.94	7.79	3.17	4.27	5.91	7.68

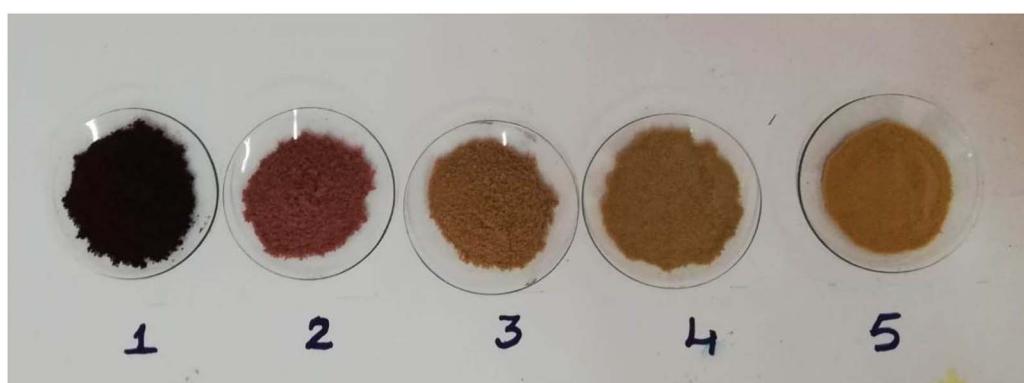


Fig. 26 : Powders 1 to 4: fresh samples of NICSS added to AR119-TIE solution after every 15 min, filtered and the residue dried in an oven. Sample 5: NICSS.

For academic curiosity, preliminary investigations were carried out to desorb the adsorbed toxicants on NICSS. The solvents we aimed to use were methyl alcohol, ethyl alcohol, acetone, and dimethylformamide. However, dimethylformamide was not used because it is toxic. Use of acetone created turbidity after three cycles. Methyl alcohol and ethyl alcohol displayed

almost similar results. Methyl alcohol was toxic compared with ethyl alcohol. However, the cost of ethyl alcohol is exorbitantly high compared with the cost of the material. Five cycles were carried out with an efficiency of ~90%. However, the idea of recycling is not a desirable proposition because the cost of recycling is approximately 500-times than the cost of the adsorbent.

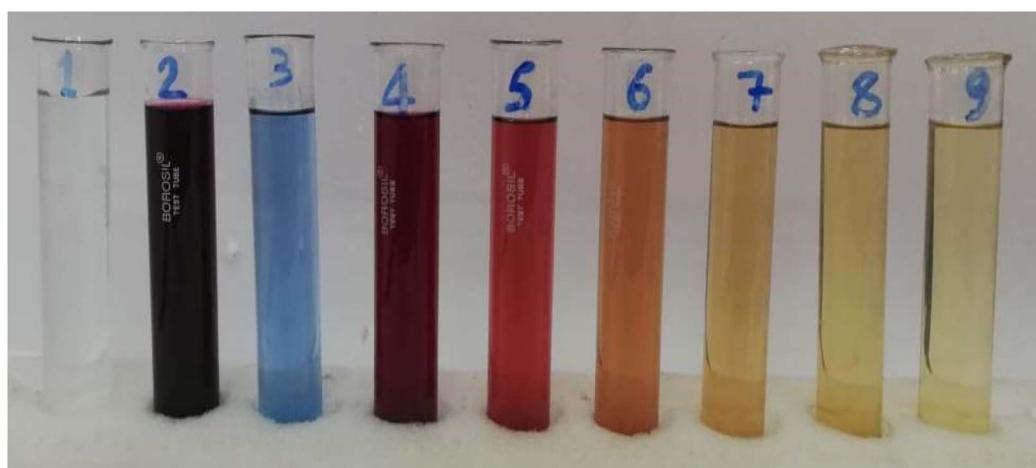


Fig. 27 Color of the solutions before and after adsorption. (1) Distilled water; (2) AR119 in distilled water; (3) TIE; (4) AR119 in TIE; (5) filtrate after adsorption of dye on NICSS after 15 min; (6) 30 min; (7) 45 min; (8) 60 min; (9) filtrate of NICSS in distilled water.



## 4. Conclusions

NICSS has been utilized as a biosorbent for the treatment of harmful dyes, adding value to the waste produced in the nutraceutical sector. NICSS showed efficacy as an adsorbent for AR119 in an aqueous solution. The adsorption mechanism of AR119 was nearly spontaneous and endothermic. The results aligned closely with the pseudo-second-order kinetic model. Intra-particular diffusion was significantly impacted by the film, and the method was primarily physical. The FTIR and SEM spectra verified that AR119 was adsorbed onto NICSS. Using by-products from nutraceutical industries as adsorbents to tackle pollution offers a more sustainable, cost-effective, and efficient solution. As an assumption, tonnes of renewable trash and waste materials are being produced by the unprecedently fast ascent of the nutraceutical industries to prominence in the global economy. The ecosystem is being adversely affected by this. Under such circumstances, the agro-industrial sector, which includes the nutraceutical industry, is unable to sustainably contaminate the environment. Efficient management of nutraceutical expenditures and/or waste is urgently needed to maintain the development of the nutraceutical industry. The key elements of recycling include reutilization, renewal, and retrieval. These are the remedial strategies used in environmental management employing the concepts of the CE. In brief, the textile industry is facing new challenges with emerging trends such as sustainability, transparency, and new businesses with consumption models derived from the waste of one industry to be used as a raw material for another industry and/or a process. This aligns with the CE model based on product design and development, waste collection and sorting, and effectively reusing waste as a starting material for the next process.

## Data availability

Data are available upon request to the corresponding author.

## Conflicts of interest

No conflicts of interest exist among the authors.

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