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Regular Article

# Trimellitated sugarcane bagasse: A versatile adsorbent for removal of cationic dyes from aqueous solution. Part I: Batch adsorption in a monocomponent system



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#### ABSTRACT

Trimellitated-sugarcane bagasse (STA) was used as an environmentally friendly adsorbent for removal of the basic dyes auramine-O (AO) and safranin-T (ST) from aqueous solutions at pH 4.5 and 7.0. Dye adsorption was evaluated as a function of STA dosage, agitation speed, solution pH, contact time, and initial dye concentration. Pseudo-first- and pseudo-second-order, Elovich, intraparticle diffusion, and Boyd models were used to model adsorption kinetics. Langmuir, Dubinin-Radushkevich, Redlich-Peterson, Sips, Hill-de Boer, and Fowler-Guggenheim models were used to model adsorption isotherms, while a Scatchard plot was used to evaluate the existence of different adsorption sites. Maximum adsorption capacities for removal of AO and ST were 1.005 and 0.638 mmol  $g^{-1}$  at pH 4.5, and 1.734 and 1.230 mmol  $g^{-1}$  at pH 7.0, respectively. Adsorption enthalpy changes obtained by isothermal titration

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#### Nomenclature

AO	auramine-O	υ	number of degrees of freedom
$a_{\rm R}$	Redlich-Peterson constant (L mmol <sup>-1</sup> )	Р	number of variables of the model
α	initial adsorption rate (mmol $g^{-1}$ min <sup>-1</sup> )	Р	equilibrium pressure
h	Langmuir hinding constant (L mmol <sup><math>-1</math></sup> )	Po	saturation pressure
D	colid characteristic energy towards a reference		proudo first order
D	solid characteristic energy towards a reference	FFU	pseudo-mst-order
	adsorbate (mol <sup>2</sup> kJ <sup>2</sup> )	$pH_{PZC}$	point of zero charge
BET	Brunauer, Emmett, and Teller	PSO	pseudo-second-order
BIH	Barret, Javner, and Halenda	pwg	percent weight gain (%)
ß,	desorption constant ( $\sigma$ mmol <sup>-1</sup> )	Pv	pyridine
Paes	Redlich Deterson exponent (dimensionless)	r y	adcomption conscitut (mmol $a^{-1}$ )
р С	Redicti-retersoil exponent (dimensionless)	q	
C	concentration at time t or equilibrium (mmol L )	$q_{\rm s}$	maximum adsorption capacity (mmol g )
С	interaction energy among adsorbed molecules	$q_{\rm i,int}$	heats in the reaction cell at the $i^{tn}$ injection in the
	$(k mol^{-1})$		presence of the STA adsorbent (kJ)
CN	coordination number	<b>()</b> :	heats in the reaction cell at the $i^{th}$ injection in the
Л	effective diffusion coefficient $(m^2 min^{-1})$	91,011	absonce of the STA adsorbant (kl)
	NN dimethal asta wide	0	absence of the STA ausoident $(K_j)$
DMA	N,N-dimethylacetamide	Q <sub>max</sub>	maximum adsorption capacity (mmol g <sup>-1</sup> )
DTGS	deuterated triglycine sulfate detector	Q <sub>max,e</sub>	maximum adsorption capacity at equilibrium
D-R	Dubinin-Radushkevich		$(\text{mmol g}^{-1})$
$\Delta_{\rm ads} H$	adsorption enthalpy change $(kI mol^{-1})$	Ore-ads	maximum re-adsorption capacity (mmol $g^{-1}$ )
A Hdye-dye	enthalow changes associated with the dve-dve inter-	R	dve removal percentage (%)
∠ads <sup>1</sup>	actions on the adsorbant surface ( $kI$ mol <sup>-1</sup> )		Podlich Deterson
	actions on the ausorbent surface (kj mor)	N-F p <sup>2</sup>	Reditch-reterson
$\Delta_{\rm ads} H^{\rm aye \ sm}$	enthalpy changes associated with the formation of	R <sup>2</sup>	determination coefficient
	dye-adsorbent interactions (kJ mol <sup><math>-1</math></sup> )	RSS	residual sum of squares
$\Delta_{ads}H^{desol}$	enthalpy changes associated with the desolvation of	SB	sugarcane bagasse
	adsorption sites on the STA surface and dve mole-	ST	safranin-T
	cules in the bulk solution (kI mol <sup><math>-1</math></sup> )	STA	trimellitated sugarcane bagasse
Е	characteristic energy of adcorption $(kJ mol^{-1})$		STA loaded with auramine O
	characteristic energy of ausorption (Kj mor	SIA-AU	STA loaded with acfurin T
EDX	energy dispersive X-ray spectroscopy	51A-51	SIA-loaded with safranin-1
Edes	desorption efficiency (%)	STA-DAO	STA after desorption of auramine-O
$E_{\rm re-ads}$	re-adsorption efficiency (%)	STA-DST	STA after desorption of safranin-T
8	adsorption potential (kI mol $^{-1}$ )	te	equilibrium time (min)
f	fractional loading	to	the time constant (min)
J F C	Fowler Cuggenheim	0	fractional coverage
	Fourier transforms informed an estimation	U	indutional coverage
FIIK	Fourier transform infrared spectroscopy	V	volume of the dye solution (L)
γe	dye activity coefficient at equilibrium	Wi	weighting coefficient
h	initial adsorption rate of the pseudo-second-order	W <sub>STA</sub>	weight of STA (g)
	kinetic model (mmol $g^{-1}$ min <sup>-1</sup> )	W <sub>STA dve</sub>	weight of the STA loaded with a dye (g)
H-B	Hill-de Boer	W'STA	weight of STA in $W_{STA, dvo}(g)$
IARC	international Agency for Research on Cancer	1W/ ,	weight of dve not desorbed from the STA adsorbent
ITC	isothermal titration calorimetry	w dye	after the description (mg)
IIC ID		,,	
IR	infrared	W'dye	weight of dye adsorbed on the SIA in the re-
$k_1$	pseudo-first-order rate constant (min <sup>-1</sup> )		adsorption experiment (mg)
$k_2$	pseudo-second-order rate constant (g mmol <sup><math>-1</math></sup> min <sup><math>-1</math></sup> )	$\chi^2$	chi-square
K <sub>2</sub>	thermodynamic equilibrium constant (dimension-	$\chi^2$ red	reduced chi-square
- a	less)	<i>V</i> .	experimental data point
V	$\begin{array}{c} 1 \\ 1 \\ 1 \\ 1 \\ 1 \\ 1 \\ 1 \\ 1 \\ 1 \\ 1 $	$\hat{y}_1$	estimated data point calculated by the model
KHB		<i>y</i> i	estimated data point calculated by the model
<i>K</i> <sub>i</sub>	intraparticle diffusion rate constant (mmol g	Z	charge of the adsorbate
	$\min^{-1/2}$ )		
K <sub>FG</sub>	Fowler-Guggenheim constant (L mmol <sup>-1</sup> )	Subscripts	
KR	Redlich-Peterson constant (L $g^{-1}$ )	ads	adsorption
<i>II</i>	ionic strength (mol $I^{-1}$ )	ro ada	re adcomption
n	number of carboxylic acid groups (mmol $a^{-1}$ )	re-aus	
"COOH	Since model parameter accepted with the hotor	e	equilibrium
п	sips model parameter associated with the netero-	exp	experimental
	geneity of the adsorption system	t	time
ni	amount of dye (in mol)	Т	theoretical value obtained from the model
Ν	number of experimental data points		
	-		

Safranin-T Desorption Isothermal titration calorimetry calorimetry (ITC) ranged from  $-21.07 \pm 0.25$  to  $-7.19 \pm 0.05$  kJ mol<sup>-1</sup>, indicating that both dyes interacted with STA by physisorption. Dye desorption efficiencies ranged from 41 to 51%, and re-adsorption efficiencies ranged from 66 to 87%, showing that STA can be reused in new adsorption cycles. ITC data combined with isotherm studies allowed clarification of adsorption interactions.

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