REMOVAL OF CHROMIUM (VI) FROM ELECTROPLATING WASTEWATER USING TEA WASTE

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ABSTRACT: Electroplating wastewater contains some heavy metals that pollute natural water. It is very difficult to treat it by conventional treatment processes due to its high cost and difficult operating conditions. Adsorption in one of the alternatives. As cost is important parameter low cost adsorbent getting interest. In this paper adsorption of chromium from electroplating industry by tea waste has been studied at laboratory scale. Electroplating waste was obtained from an industry in Vadodara district. Some parameters like initial pH, dosage of adsorbent, agitation speed and contact time were studied and optimum operational condition for chromium removal of wastewater was derived.

Keywords Adsorption Process, Electroplating wastewaters, waste tea

1. INTRODUCTION

1.1 Background

Dangerous metals are discharged into the nature by different mechanical exercises like refining minerals, compost businesses, mining, tanneries, and pesticides and so on. The major harmful metal particles which are dangerous to people and condition are Cr, Fe, Se, V, Cu, Co, Ni, Cd, Hg, As, Pb, Zn and so forth. These overwhelming metals are of explicit worry because of their poisonous quality, bio-aggregation inclination and persistency in nature. Different administrative bodies have set the most extreme endorsed limits for the release of poisonous substantial metals in the earth. Still the waste water containing overwhelming metals are being discharged at higher fixation into nature which influences the strength of people just as condition.

Some Conventional strategies for expulsion of metal particles from water bodies incorporate substance precipitation, particle exchangers, concoction oxidation/decrease, switch assimilation, electro dialysis, and ultra-filtration and so on. Anyway these regular strategies have a few confinements, for example, less productivity, delicate working conditions, and creation of auxiliary muck and further the transfer is exorbitant. Another option is adsorption of substantial metals by enacted carbon for treating waste water yet the surprising expense of initiated carbon and its misfortune amid the recovery confines its application.

As of late consideration has been redirected towards the waste materials which are created in a few ventures and horticulture exercises. The real points of interest of adsorption over regular treatment techniques are: ease, high effectiveness, minimization of compound or organic muck, no extra supplement prerequisite, and recovery of adsorbents and plausibility of metal recuperation.

Agrarian materials especially those containing cellulose indicates potential metal adsorption limit. The fundamental segments of the farming waste materials incorporate hemicelluloses, lignin, extractives, lipids, proteins, basic sugars, water hydrocarbons, starch containing assortment of useful gatherings that encourages adsorption of overwhelming metals on to the surface. Horticultural waste materials are monetary and eco inviting because of their one of a kind compound piece, accessibility in huge amount, sustainable, low in expense and more effectiveness.

Different rural waste materials, for example, rice grain, rice husk, wheat grain, wheat husk, saw residue of different plants, bark of the trees, ground nutshells, coconut shells, dark gram husk, walnut shells, cotton seed frames, squander tea leaves, cassia fistula leaves, maize corn cob, sugarcane, apple, banana, orange strips, soybean bodies, grapes stalks, espresso beans, cotton stalks and so on has been attempted. These horticultural waste materials are utilized in the evacuation of metal particles either in their normal structure or after some physical or synthetic change.

The electroplating industry is one such type of industry whose effluent is directly released into the fresh water bodies thus polluting the environment. The electroplating industry is a large industry operated throughout the year and generates a large amount of effluent which contains high load of heavy metals. The industry uses electrolytic processes for plating heavy metals on the metal sheet. During this process large quantities of heavy metals remain in the effluent. These effluents are released out openly into the environment and get mixed into the rivers and ponds. The channel of the effluent may have many cracks through which the polluted water gets percolated deep inside to contaminate the ground water.

In this research, tea waste is used to remove heavy metal chromium (VI) from electroplating industry waste water.

2 ADSORPTION USING TEA WASTE

2.1 Material pre-treatment

Before the use of tea waste it needs to go through some pre-treatments like washing and sieving. Washing of tea waste remove the considerable amount of hydrolysable tannins, polysaccharides and proteins along with other coloured and soluble components to avoid contamination. Distilled water was used to wash tea waste. The tea waste was washed until a colourless solution remained. Washed Tea waste was then dried in an oven for 48 hours at 180°C. Then desired particle size of tea waste was obtained by sieving.

2.2 Methodology

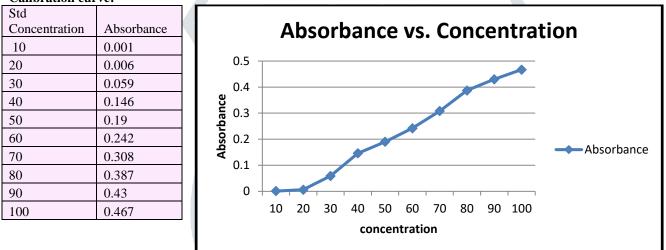
UV-VIS Spectrophotometer was used to measure chromium concentration using the method based on the reaction of Cr (VI) and diphenyl carbazide which forms a red-violet coloured complex. Standard calibration curve was obtained by taking different amount of Cr in $K_2Cr_2O_7$.

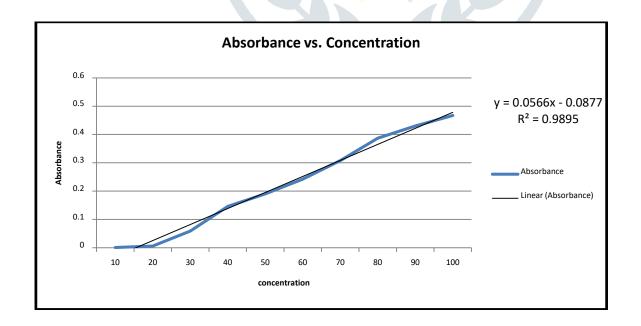
Experiments were conducted in the 250ml flask. 200ml of unknown sample was taken for each set of experiment. Experiments were done at different pH values ranging from 1-3, different adsorbent dose ranging from 3 to 5 g/200ml, different agitation speeds 200, 360 and 480 rpm. Samples were taken at time interval of 15 min (15, 30, 45, 60, 75, 90, 105, and 120) to find optimum contact time. Magnetic stirrer was used to maintain speed.

Initial pH of wastewater was 3.0. To vary pH, 1.0 m HCl was used.

Initial chromium concentration was 117.474 µg/ml.

Calibration curve:





3. Results and discussion

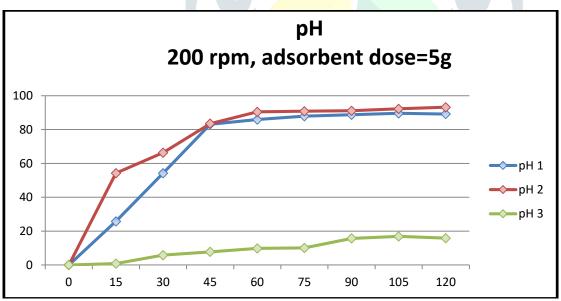
3.1 Effect of pH

The pH of the solution is an effective controlling parameter. Thus the ion concentration was examined at different pH, covering a range of 1.0 to 3.0.

A graph was plotted of % reduction vs. Time keeping other parameters constant. Speed was constant 200rpm, adsorbent dose was taken 5g, and effects of pH 1.0, 2.0 and 3.0 were examined.

U'	1 /	
	pH = 1	
Time	Concentration(µg/ml) % reduction	
(min)		
0	117.474 0	
15	87.222	25.75
30	53.756	54.24
45	19.8275	83.12
60	16.55	85.911
75	14.232	87.88
90	13.225	88.742
105	12.22	89.59
120	12.69	89.19
	pH = 3	
Time	Concentration(µg/ml)	% reduction
(min)		
0	117.474	0
15	116.542	0.79
30	110.632	5.82
45	108.36	7.758
60	105.98	9.784
75	105.66	10.054
90	99.002	15.724
105	97.664	16.86
120	98.887	15.822

	pH = 2					
Time	Concentration(µg/ml)	% reduction				
(min)						
0	117.474	0				
15	53.756	54.24				
30	39.551	66.33				
45	19.367	83.51				
60	11.225	90.44				
75	10.75	90.84				
90	10.435	91.117				
105	9.09	92.262				
120	8.002	93.188				



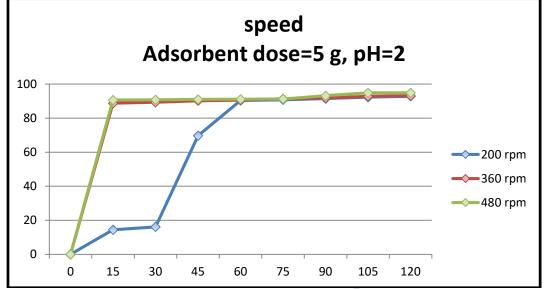
The maximum removal of chromium was found at pH = 2.

3.2 Effect of speed

The speed considerably affects the removal of chromium concentration. Speed 200rpm, 360rpm and 480rpm were examined.

				360 rpm	
200 rpm		Time (min)	Concentration(µg/ml)	% reduction	
Time (min)	Concentration(µg/ml)	% reduction	0	117.474	0
0	117.474	0	15	13.221	88.745
15	100.517	14.434	30	12.554	89.31
30	98.521	16.133	45	11.5	90.21
45	35.5695	69.72	60	11.03	90.61
60	11.32	90.363	75	10.09	91.41
75	10.849	90.764	90	9.62	91.81
90	10	91.487	105	8.4	92.84
105	9	92.338	120	8	93.189
120	84	92.84			

480 rpm			
Time(min)	Concentration(µg/ml)	% reduction	
0	117.474	0	
15	11.003	90.633	
30	10.849	90.764	
45	10.566	91.005	
60	10.435	91.117	
75	10.283	91.24	
90	8	93.189	
105	6	94.892	
120	6	94.892	

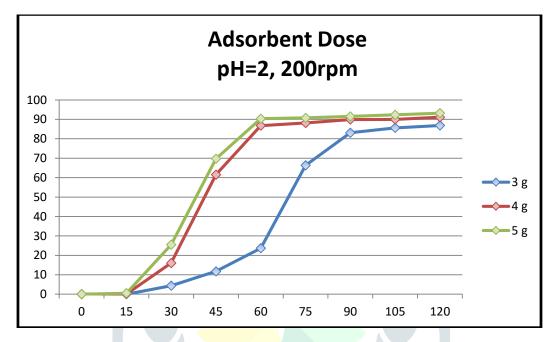


From results we can conclude that 200 rpm speed is enough for chromium removal as there is no more difference between % reduction at 200, 360 and 480 rpm

3.3. Effect of Adsorbent dos

3 g		4 g			
Time	Concentration(µg/ml)	% reduction	Time	Concentration(µg/ml)	% reduction
(min)			(min)		
0	117.474	0	0	117.474	0
15	117.474	0	15	117.361	0.096
30	112.323	4.385	30	98.62	16.05
45	103.650	11.768	45	45.221	61.506
60	89.68	23.66	60	15.52	86.789
75	39.551	66.332	75	13.96	88.117
90	19.828	83.122	90	11.856	89.908
105	16.89	85.622	105	11.784	89.969
120	15.447	86.851	120	10.47	91.087

	5 g	
Time(min)	Concentration(µg/ml)	% reduction
0	117.474	0
15	116.897	0.491
30	87.561	25.464
45	35.5695	69.721
60	11.32	90.364
75	10.849	90.765
90	10	91.487
105	9	92.339
120	8	93.19



Maximum removal was observed at 5g adsorbent dose. 93% removal was obtained at 25 gL⁻¹ adsorbent dose.

3.4 Effect of contact time

From above all experiments we can conclude that 60min is an optimum contact time. Maximum removal was observed upto 60min. After that removal was almost constant. 90% removal has been observed at 60min.

4. Conclusion

Following conclusion can be drawn from the present study:

- 1. The present experiment shows that the tea waste is an effective adsorbent for the removal Cr (VI).
- 2. The adsorption process is the function of pH, agitation speed, Adsorbent dose, and contact time.
- 3. Adsorption is maximum at pH=2.
- 4. As adsorbent dose increases, removal of chromium ion increases.
- 5. With the increase in contact time, adsorption increases, and maximum adsorption takes place in 60 min.
- 6. Agitation speed of 200rpm is enough for this process because all agitation speeds show almost same results.

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