



# The Corrosion of Mild Steel Checked by the Application of Octahydrodibenzo[a,d][8] Annulene-5,12-Dioxime and ZnS Filler in Corrosive Environments



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## Abstract

Mild steel is economical metal so it is used in different appliances of industries, railways, bridges, construction works and marine water. Marine water produces corrosive environment for this metal. Marine water is saline in character so it corrodes mild steel. Poly-butadiene is coated on the surface of mild steel for corrosion protection. But this coating does not provide sufficient protection of base metal. Marine water is major absorber of  $\text{CO}_2$ . It converts  $\text{CO}_2$  into  $\text{H}_2\text{CO}_3$ . Saline water and  $\text{H}_2\text{CO}_3$  interacts with poly-butadiene-coated mild steel and they exhibit chemical and electrochemical reactions. This chemical reaction produces swelling and dissolving corrosion and produce disbonding between carbon and carbon of poly-butadiene. These corrosive agents show osmosis or diffusion process and inter inside the base metal and develop corrosion cell. Metal generates corrosion reaction which produces several forms of corrosion like galvanic, pitting, crevice, stress, intergranular, blistering and embrittlement. For the protection from such types of corrosions synthesized organic compounds octahydrodibenzo[a,d][8]annulene-5,12-dioxime was used and this compounds was nano-coated on the surface of poly-butadiene-coated mild steel. During nano-coating used compound developed lot of porosities on the surface of poly-butadiene-coated mild steel and they were blocked by the use of filler materials ZnS. The corrosion rate of material was calculated by gravimetric and potentiostat. Composite barrier formation was analyzed by activation energy, heat of adsorption, free energy, enthalpy and entropy. The results of surface coverage area and coating efficiency were indicated that compound octahydrodibenzo[a,d][8]annulene-5,12-dioxime and ZnS have good coating and filling properties.

**Keywords:** Marine environment; Nano-coating; Filler; Corrosion; Polybutadiene; Mild steel

## Introduction

Corrosion is just like diabetes for materials. Corrosion control of materials is major problems. Develop nation's expense 5% their GNP for corrosion protection and repairing of materials. Corrosion occurs in metals, metalloids, ceramics, polymers, bio-metals, building materials and other type's metallic and nonmetallic equipments. There are several methods are applied for corrosion mitigation of materials [1] as per materials nature and environments. Generally most of metallic industries use metallic coatings but such types of coating do not provide good results in corrosive medium [2]. Now days some industries apply polymeric coating [3] for safe base metal in hostile atmosphere [4]. This coating does not provide protection [5] of base and its own. Disintegrations occur in base metal [6] and polymeric material [7]. Paint coating [8] is doing for building materials [9] and metal corrosion protection [10] in unfriendly atmosphere [11] but it is not protect itself and coating materials [12] for the attack of pollutants [13]. Organic [14] and inorganic inhibitors [15] are a very useful in acidic, basic and

moist oxygen environment [16]. These inhibitors do not give good results after certain period. There are several types of inhibitors [17] like anodic, cathodic and mixed [18] of types use as nature environments of pollutants [19]. They do not provide satisfactory results in against pollutants [20]. Nano-coating techniques [21] are a very suitable method for corrosion protection materials in any type's corrosive ambient. Nano-coating substances forms composite thin film barrier, top layer coating, thermal barrier coating, conversion coating and nano scale structural change coatings on surface of base materials as per materials applications in different fields [22] to safe it's against heat, light, temperature, particulates, acidic, basic, salt, marine atmosphere, flues gases and biological environment [23]. Such types of coating produce lot of porosities and corrosive agents are entered inside by osmosis or diffusion process and produce interior and exterior corrosion. In this research work octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime is used a nano-coating material and porosities are blocked by filler

ZnS. These materials create composite barrier and it works repeller against corrosive pollutants.

## Experimental

Poly-butadiene-coated mild steel kept in marine water and water sample was taken from Marina beach of Chennai (Tamil Nadu). Poly-butadine coated-mild steel (5X10X was in dipped into sea water and the corrosion rate was determined gravimetric methods at 283, 293,303,313 and 333 °K temperatures and time mentioned in given temperatures 2,5,8,11 and 14 days. The samples were nanocoated with octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and immersed into sea water and corrosion rate recorded at above mentioned temperatures and days. Nanocoated samples were coated by ZnS filler and dipped into sea water to calculate the corrosion rate given temperatures and days. The

corrosion potential, corrosion current, corrosion current densities were calculated by potentiostat technique. For these results pt electrode used as reference electrode, calomel as auxiliary electrode and polybutaine-coated mild steel sample electrode. Nanocoated compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime was synthesized by given methods as:

### Scheme1: synthesis of 4-chloro-1, 2-dihydronaphthalene

When 3, 4-dihydronaphthalen-1(2H)-one (25gm) is added into cold solution of benzene (50gm) containing  $\text{PCl}_5$  (30gm), the reaction mixture was stirred for one hour. The reaction mixture was quenched with  $\text{NaHCO}_3$  and did workup with diethyl ether. The solvent evaporated with rotator vapor. The product was purified by silica gel column chromatography and produced 89% 4-chloro-1, 2-dihydronaphthalene (Figure 1-3).

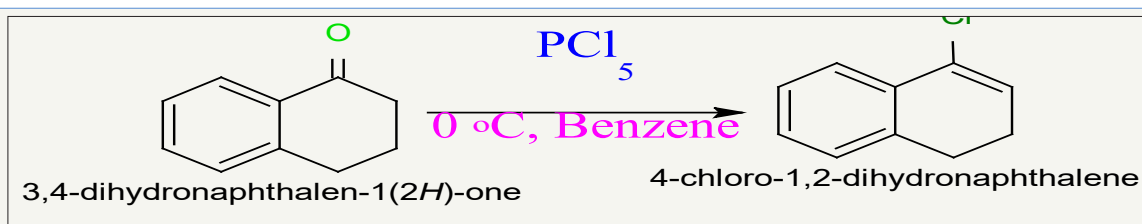


Figure 1: Physical properties of 4-chloro-1, 2-dihydronaphthalene.

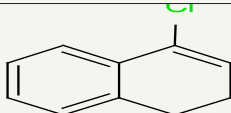
	Molecular Formula	= C <sub>10</sub> H <sub>9</sub> Cl
	Formula Weight	= 164.63146
	Composition	= C(72.96%) H(5.51%) Cl(21.53%)
	Molar Refractivity	= 47.73 ± 0.4 cm <sup>3</sup>
	Molar Volume	= 141.2 ± 5.0 cm <sup>3</sup>
	Parachor	= 355.3 ± 6.0 cm <sup>3</sup>
	Index of Refraction	= 1.590 ± 0.03
	Surface Tension	= 40.0 ± 5.0 dyne/cm
	Density	= 1.16 ± 0.1 g/cm <sup>3</sup>
	Dielectric Constant	= Not available
	Polarizability	= 18.92 ± 0.5 10 <sup>-24</sup> cm <sup>3</sup>
	Monoisotopic Mass	= 164.039278 Da
	Nominal Mass	= 164 Da
	Average Mass	= 164.6315 Da
	M+	= 164.038729 Da
M-	= 164.039827 Da	
[M+H] <sup>+</sup>	= 165.046554 Da	
[M+H] <sup>-</sup>	= 165.047652 Da	
[M-H] <sup>+</sup>	= 163.030904 Da	
[M-H] <sup>-</sup>	= 163.032002 Da	

Figure 2: <sup>1</sup>H NMR of 4-chloro-1, 2-dihydronaphthalene.

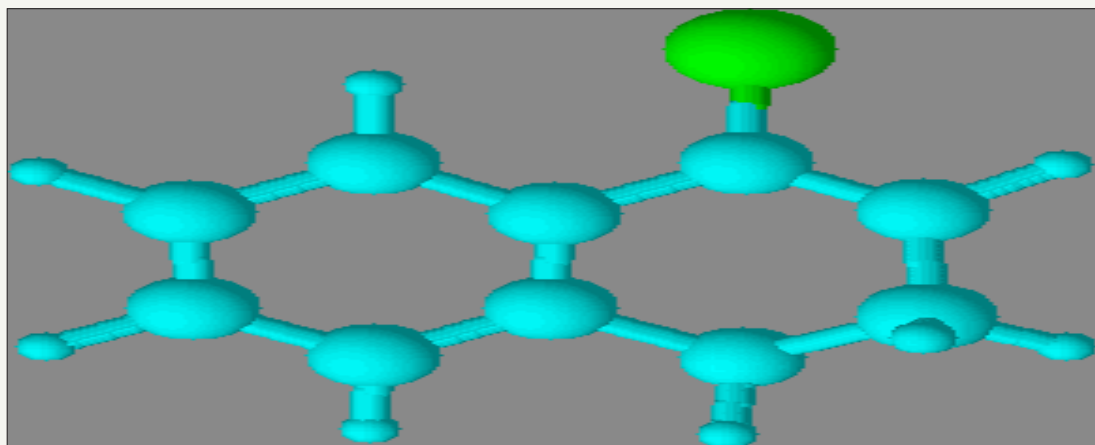
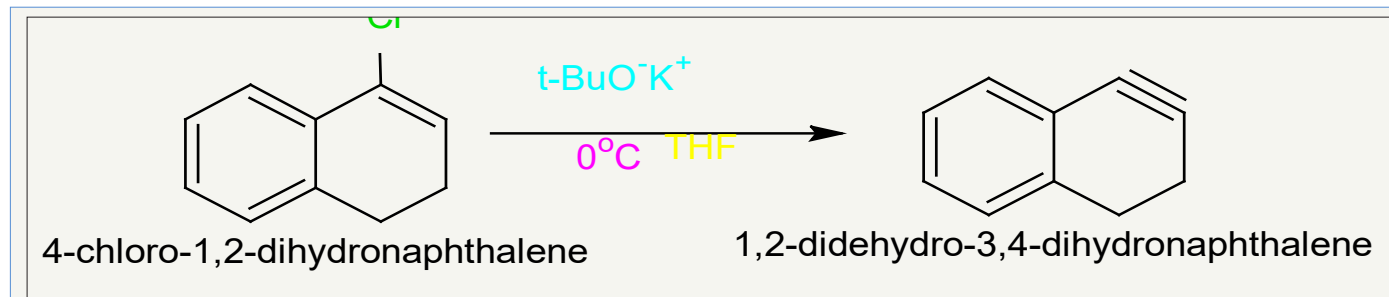


Figure 3: XRD of 4-chloro-1,2-dihydronaphthalene.

**Scheme2: synthesis of 1,2-didehydro-3,4-dihydronaphthalene**

4-Chloro-1, 2-dihydronaphthalene (10gm) kept in two neck round bottle flask and potassium t-butoxide (25gm) dissolved in THF solution. This solution poured into 4-Chloro-1,

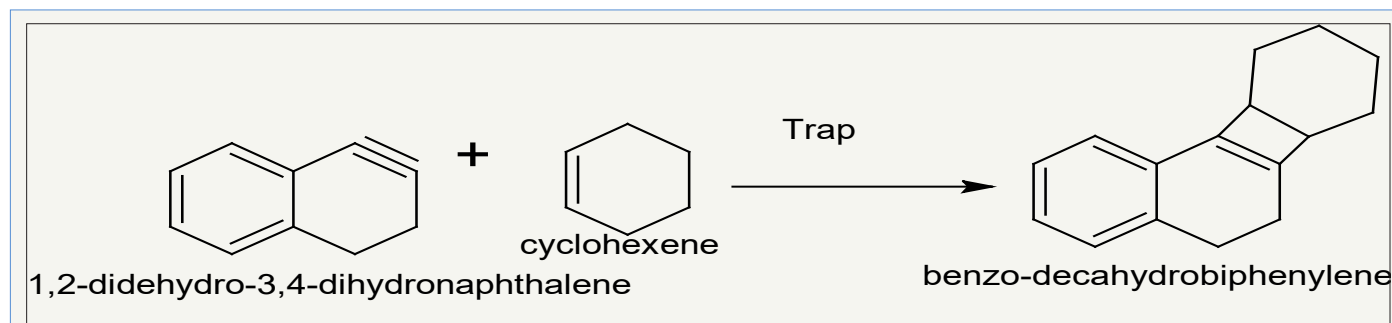
2-dihydronaphthalene and reaction temperature 0 °C. The reaction was mixture stirring four hours after completion reaction added cyclohexene as trapping agent and again stirring reaction more two hours. After work up got adduct 90% of 1, 2-didehydro-3, 4-dihydronaphthalene (Figure 4 & 5).



**Figure 4:** Physical properties of 1, 2-didehydronaphthalene.

	Molecular Formula	= C <sub>10</sub> H <sub>8</sub>
	Formula Weight	= 128.17052
	Composition	= C(93.71%) H(6.29%)
	Molar Refractivity	= 41.23 ± 0.4 cm <sup>3</sup>
	Molar Volume	= 119.7 ± 5.0 cm <sup>3</sup>
	Parachor	= 309.8 ± 6.0 cm <sup>3</sup>
	Index of Refraction	= 1.604 ± 0.03
	Surface Tension	= 44.8 ± 5.0 dyne/cm
	Density	= 1.07 ± 0.1 g/cm <sup>3</sup>
	Dielectric Constant	= Not available
	Polarizability	= 16.34 ± 0.5 10 <sup>-24</sup> cm <sup>3</sup>
	Monoisotopic Mass	= 128.0626 Da
	Nominal Mass	= 128 Da
Average Mass	= 128.1705 Da	
M+	= 128.062052 Da	
M-	= 128.063149 Da	
[M+H] <sup>+</sup>	= 129.069877 Da	
[M+H] <sup>-</sup>	= 129.070974 Da	
[M-H] <sup>+</sup>	= 127.054227 Da	
[M-H] <sup>-</sup>	= 127.055324 Da	

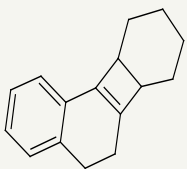
**Figure 5:** H<sup>1</sup>NMR of 1, 2-didehydronaphthalene.

**Scheme3: Synthesis of benzo-decahydrobiphenylene**

**Figure 6:** Physical properties of benzo-decahydrobiphenylene.

When 1, 2-didehydro-3, 4-dihydronaphthalene was used with cyclohexene, it was trapped by 1,2-didehydro-3,4-

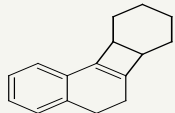
dihydronaphthalene to yield benzo-decahydrobiphenylene (Figure 6-8).



Molecular Formula	= C <sub>16</sub> H <sub>18</sub>
Formula Weight	= 210.31412
Composition	= C(91.37%) H(8.63%)
Molar Refractivity	= 66.33 ± 0.4 cm <sup>3</sup>
Molar Volume	= 191.6 ± 5.0 cm <sup>3</sup>
Parachor	= 490.1 ± 6.0 cm <sup>3</sup>
Index of Refraction	= 1.608 ± 0.03
Surface Tension	= 42.7 ± 5.0 dyne/cm
Density	= 1.09 ± 0.1 g/cm <sup>3</sup>
Dielectric Constant	= 3.09 ± 0.2
Polarizability	= 26.29 ± 0.5 10 <sup>-24</sup> cm <sup>3</sup>
Monoisotopic Mass	= 210.140851 Da
Nominal Mass	= 210 Da
Average Mass	= 210.3141 Da
M+	= 210.140302 Da
M-	= 210.141399 Da
[M+H] <sup>+</sup>	= 211.148127 Da
[M+H] <sup>-</sup>	= 211.149224 Da
[M-H] <sup>+</sup>	= 209.132477 Da
[M-H] <sup>-</sup>	= 209.133574 Da

Figure 7: H<sup>1</sup>NMR of benzo-decahydrobiphenylene.

ChemNMR <sup>1</sup>H Estimation



Estimation quality is indicated by color: **good**, **medium**, **rough**

Protocol of the H-1 NMR Prediction (Lib=SU Solvent=DMSO 300 MHz):

Node	Shift	Base + Inc.	Comment (ppm rel. to TMS)
CH 2.15	1.44		cyclohexane
	0.68		1 alpha -C=C from methine
	0.03		1 beta -C=C from methine
CH 2.15	1.44		cyclohexane
	0.68		1 alpha -C=C from methine
	0.03		1 beta -C=C from methine
CH2 2.34,2.24	0.00		1.37 methylene
	0.63		1 alpha -C=C
	0.29		1 beta -1 C *C *C *C *C *1
CH2 1.41,1.31	0.00		1.44 cyclohexane
	0.00		1 beta -C=C from methylene
	-0.08		generalconnections
CH2 1.41,1.31	0.00		1.44 cyclohexane
	0.00		1 beta -C=C from methylene
	-0.08		generalconnections
CH 7.34	7.26		1-benzene
	0.04		1-C=C
	0.00		1-C-C
	0.04		generalconnections
CH 7.44	7.26		1-benzene
	-0.05		1-C=C
	-0.08		1-C-C
	0.31		generalconnections
CH 7.27	7.26		1-benzene
	-0.05		1-C=C
	-0.10		1-C-C
	0.16		generalconnections
CH 7.65	7.26		1-benzene
	-0.12		1-C=C
	0.00		1-C-C
	0.51		generalconnections
CH2 2.59	1.37		methylene
	1.22		1 alpha -1 C *C *C *C *C *C *1
	0.00		1 beta -C=C
CH2 1.53,1.43	0.00		1.44 cyclohexane
	0.04		generalconnections
CH2 1.53,1.43	0.00		1.44 cyclohexane
	0.04		generalconnections

1H NMR Coupling Constant Prediction

shift	atom	index	coupling partner, constant and vector
2.15		12	
		11	7.0 H-C-C-H
		16	7.0 H-C-CH-H
2.15		11	
		12	7.0 H-C-C-H
		13	7.0 H-C-CH-H
2.29		8 diastereotopic	-12.4 H-C-H
		10	7.1 H-CH-C-H-H
1.36		16 diastereotopic	-12.4 H-C-H
		12	7.0 H-C-C-H
		15	7.1 H-CH-C-H-H
1.36		13 diastereotopic	-12.4 H-C-H
		11	7.0 H-CH-C-H
		14	7.1 H-CH-C-H-H
7.34		3	
		5	7.5 H-C *C -H
		6	1.5 H-C *CH *C -H
7.44		4	
		6	7.5 H-C *C -H
		5	1.5 H-C *CH *C -H
7.27		5	
		3	7.5 H-C *C -H
		6	7.5 H-C *C -H
		4	1.5 H-C *CH *C -H
7.65		6	
		4	7.5 H-C *C -H
		5	7.5 H-C *C -H
		3	1.5 H-C *CH *C -H
2.59		10	
		9	7.1 H-CH-C-H-H
1.48		15 diastereotopic	-12.4 H-C-H
		16	7.1 H-CH-C-H-H
		14	7.1 H-CH-C-H-H
1.48		14 diastereotopic	-12.4 H-C-H
		13	7.1 H-CH-C-H-H
		15	7.1 H-CH-C-H-H

Figure 8:

## Scheme4: Synthesis of octahydrodibenzo [a,d] [8] annulene-5, 12-dione

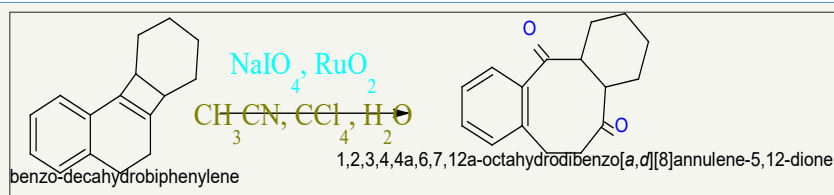


Figure 9: Physical properties of octahydrodibenzo[a,d][8]annulene-5,12-dione.

Molecular Formula	= C <sub>14</sub> H <sub>12</sub> O <sub>2</sub>
Formula Weight	= 242.31292
Composition	= C(79.31%) H(7.49%) O(13.21%)
Molar Refractivity	= 69.02 ± 0.3 cm <sup>3</sup>
Molar Volume	= 216.7 ± 3.0 cm <sup>3</sup>
Parachor	= 549.8 ± 6.0 cm <sup>3</sup>
Index of Refraction	= 1.549 ± 0.02
Surface Tension	= 41.4 ± 3.0 dyne/cm
Density	= 1.118 ± 0.06 g/cm <sup>3</sup>
Dielectric Constant	= Not available
Polarizability	= 27.36 ± 0.5 10 <sup>-24</sup> cm <sup>3</sup>
Monoisotopic Mass	= 242.13068 Da
Nominal Mass	= 242 Da
Average Mass	= 242.3129 Da
M+	= 242.130131 Da
M-	= 242.131228 Da
[M+] <sup>+</sup>	= 243.137956 Da
[M+] <sup>-</sup>	= 243.139053 Da
[M-H] <sup>+</sup>	= 241.122306 Da
[M-H] <sup>-</sup>	= 241.123403 Da

Figure 10: H<sup>1</sup>NMR of octahydrodibenzo [a, d] [8] annulene-5, 12-dione.

**ChemNMR <sup>1</sup>H Estimation**

Estimation quality is indicated by color: good, medium, rough

Protocol of the H-1 NMR Prediction (Lib=SU Solvent=DMSO 300 MHz):

Node	Shift	Base + Inc.	Comment (ppm rel. to TMS)
CH 7.83	7.26		1-benzene
	0.63		1-C (=O)JC
	0.00		1-CC
	-0.06		generalconnections
CH 7.48	7.26		1-benzene
	0.08		1-C (=O)JC
	-0.08		1-CC
	0.22		generalconnections
CH 7.30	7.26		1-benzene
	0.08		1-C (=O)JC
	-0.10		1-CC
	0.06		generalconnections
CH 7.59	7.26		1-benzene
	0.18		1-C (=O)JC
	0.00		1-CC
	0.15		generalconnections
CH 3.16	1.44		cyclohexane
	1.50		1 alpha-C (=O)-1 1 1 1 1 1 1 1 1 1 from methine
	0.22		1 beta-C=O from methine
	1.44		cyclohexane
CH 2.83	1.44		cyclohexane
	0.86		1 alpha-C=O from methine
	0.53		1 beta-C (=O)-1 1 1 1 1 1 1 1 1 1 from methine
CH2 1.64,1.385000	1.44		cyclohexane
	0.15		1 beta-C (=O)-1 1 1 1 1 1 1 1 1 1 from methylene
	-0.08		generalconnections
CH2 1.73,1.475000	1.44		cyclohexane
	0.24		1 beta-C (=O)-C from methylene
	-0.08		generalconnections
CH2 1.53,1.430000	1.44		cyclohexane
	0.04		generalconnections
CH2 1.53,1.430000	1.44		cyclohexane
	0.04		generalconnections
CH2 2.83	1.37		methylene
	1.22		1 alpha-1 1 1 1 1 1 1 1 1 1
	0.24		1 beta-C (=O)-C
CH2 2.83,2.730000	1.37		methylene
	1.12		1 alpha-C (=O)JC
	0.29		1 beta-1 1 1 1 1 1 1 1 1 1

**<sup>1</sup>H NMR Coupling Constant Prediction**

shift	atom	index	coupling partner	constant and vector
7.83	3	5	7.5	H-C-C-H
		6	1.5	H-C-C-H-H
7.48	4	6	7.5	H-C-C-H
		5	1.5	H-C-C-H-H
7.30	5	3	7.5	H-C-C-H
		6	7.5	H-C-C-H
		4	1.5	H-C-C-H-H
7.59	6	4	7.5	H-C-C-H
		5	7.5	H-C-C-H
		3	1.5	H-C-C-H-H
3.16	11	10	7.0	H-C-C-H
		16	7.0	H-C-C-H-H
2.83	10	11	7.0	H-C-C-H
		13	7.0	H-C-C-H-H
1.51	16	diastereotopic	-12.4	H-C-H
		11	7.0	H-C-C-H
		15	7.1	H-C-C-H-H
1.60	13	diastereotopic	-12.4	H-C-H
		10	7.0	H-C-C-H
		14	7.1	H-C-C-H-H
1.48	15	diastereotopic	-12.4	H-C-H
		16	7.1	H-C-C-H-H
		14	7.1	H-C-C-H-H
1.48	14	diastereotopic	-12.4	H-C-H
		13	7.1	H-C-C-H-H
		15	7.1	H-C-C-H-H
2.83	7	8	7.1	H-C-C-H-H
2.78	8	diastereotopic	-12.4	H-C-H
		7	7.1	H-C-C-H-H

Figure 11:

Adduct (20gm) oxidized into benzo-decahydrobiphenylene after workup 87% yield of octahydrodibenzo [a,d] [8] annulene-5, 12-dione was obtained (Figure 9-11). The reaction was quenched with H<sub>2</sub>O and

#### Scheme5: Synthesis of octahydrodibenzo[a,d][8]annulene-5,12-dioxime

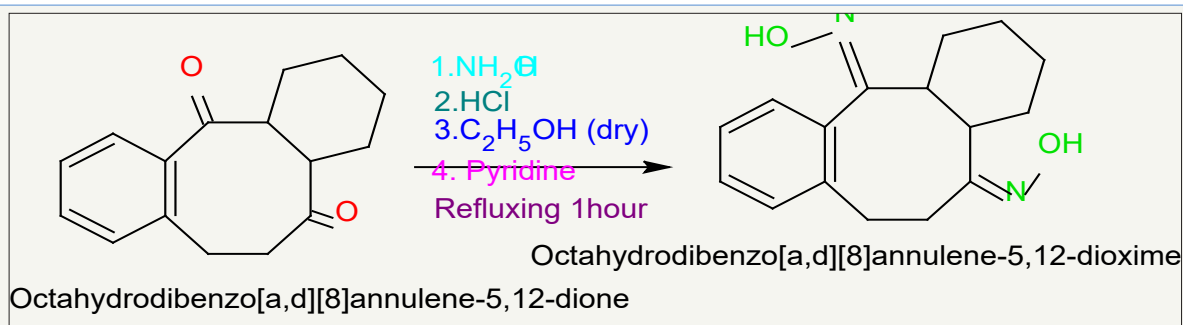


Figure 12: Physical properties of octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime.

	Molecular Formula	= C <sup>16</sup> H <sup>20</sup> N <sup>2</sup> O <sup>2</sup>
	Formula Weight	= 272.3422
	Composition	= C(70.56%) H(7.40%) N(10.29%) O(11.75%)
	Molar Refractivity	= 76.44 ± 0.5 cm <sup>3</sup>
	Molar Volume	= 208.6 ± 7.0 cm <sup>3</sup>
	Parachor	= 558.5 ± 8.0 cm <sup>3</sup>
	Index of Refraction	= 1.653 ± 0.05
	Surface Tension	= 51.3 ± 7.0 dyne/cm
	Density	= 1.30 ± 0.1 g/cm <sup>3</sup>
	Dielectric Constant	= Not available
	Polarizability	= 30.30 ± 0.5 10 <sup>-24</sup> cm <sup>3</sup>
	Monoisotopic Mass	= 272.152478 Da
	Nominal Mass	= 272 Da
	Average Mass	= 272.3422 Da
M+	= 272.151929 Da	
M-	= 272.153026 Da	
[M+H] <sup>+</sup>	= 273.159754 Da	
[M+H] <sup>-</sup>	= 273.160852 Da	
[M-H] <sup>+</sup>	= 271.144104 Da	
[M-H] <sup>-</sup>	= 271.145201 Da	

Figure 13: XRD of octahydrodibenzo [a,d] [8] annulene-5, 12-dihydrazone.

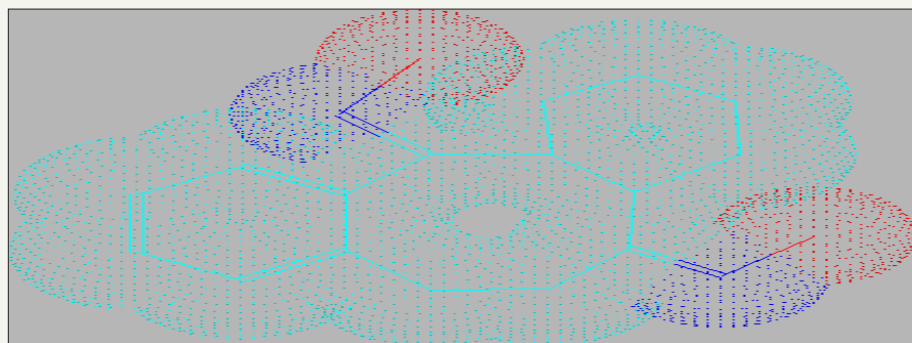


Figure 14:

Octahydrodibenzo [a,d] [8] annulene-5,12-dione (30g), hydroxylamine hydrochloride (50g) and 135ml of dry ethanol were taken and adding 70ml of pyridine and refluxing reaction mixture for two hours. Solvent was removed by use of rotator vapor and water was added in reaction mixture and after cooling the reaction by ice, solution was stirred until oxime crystallized. Solid was filtered and washed with a little water and then dried. The product was re-crystallized with ethanol and 71% yield of octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime was obtained (Figure 12-14).

### Results and Discussion

Marine water generates hostile environment for polybutadiene-coated mild steel. Corrosion is control in such hostile environment by the use of nanocoating and filler materials. Polybutadiene-coated mild steel corrosion rates were studied in marine water environment at 283, 293, 303, 313 and 323 °K temperatures

after interval of 2, 5, 8, 11 and 14days with the help of gravimetric method equation  $K(\text{mmpy})=13.56X (W/D At)$  (where W=weight loss of test coupon expressed in kg, A=area of test coupon in square meter, D=Density of the material in  $\text{kg.m}^{-3}$ ) and their values recorded in Table 1. Similarly, the samples of poly-butadiene-coated mild steel nanocoated with octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler were immersed into marine water and corrosion rate was calculated on above mentioned temperatures and days and their values were mentioned in Table 1. Figure 15 plotted between corrosion rate K(mmpy) versus times (t) in days which produced straight line and it indicated that corrosion rate of polybutadiene-coated mild steel increased without coating and their values were reduced after nanocoating and filler compounds. The results of Table 1 were shown that the corrosion rate of material is reduced by the action of nanocoating and filler compounds.

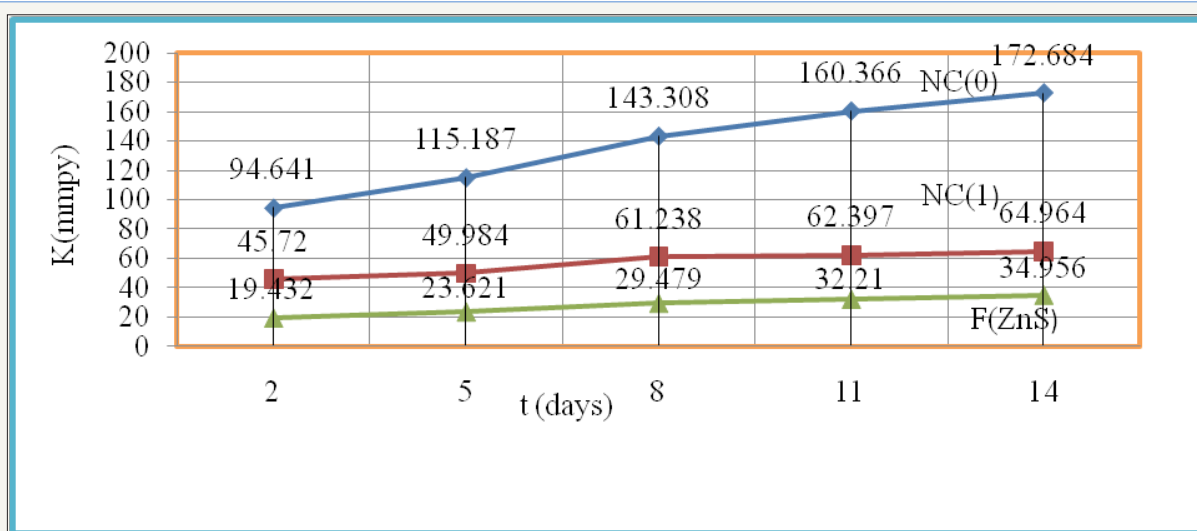


Figure 15: K (mmpy) Vs t (days) for nanocoating of NC (1) & ZnS filler on polybutadiene-coated mild steel.

Table 1: Corrosion rate of polybutadine-coated mild steel nanocoated with octahydrodibenzo [a,d] [8] annulene-5,12-dioxime[NC(1)] and ZnS filler in marine water.

N C & F	Temp(°K)	283	293	303	313	323	C(mM)
	Time(days)	2	5	8	11	14	
NC(0)	K	94.641	115.187	143.308	160.366	172.684	0
	Log K	1.976	2.061	2.156	2.205	2.237	
NC(1)	K	45.72	49.984	61.238	62.397	64.964	50
	Log K	1.66	1.698	1.787	1.795	1.812	
	log(K/T)	1.109	1.166	1.268	1.291	1.322	
	θ	0.5169	0.566	0.5721	0.6109	0.6237	
	(1-θ)	0.4831	0.444	0.4279	0.3891	0.3263	
	(θ/1-θ)	1.069	1.274	1.336	1.57	1.657	
	log(θ/1-θ)	0.028	0.105	0.125	0.195	0.219	
	%CE	51.69	56.6	57.21	61.09	62.37	



F(ZnS)	K	19.432	23.621	29.479	32.21	34.956	10
	Log K	1.288	1.373	1.469	1.507	1.543	
	log(K/T)	0.74	0.84	0.95	1.004	1.053	
	$\theta$	0.7946	0.7949	0.7942	0.7987	0.7967	
	(1- $\theta$ )	0.2054	0.2051	0.2058	0.2013	0.2039	
	( $\theta/1-\theta$ )	3.868	3.875	3.859	3.967	3.918	
	log( $\theta/1-\theta$ )	0.587	0.588	0.586	0.598	0.593	
	%CE	79.46	79.49	79.42	79.87	79.67	

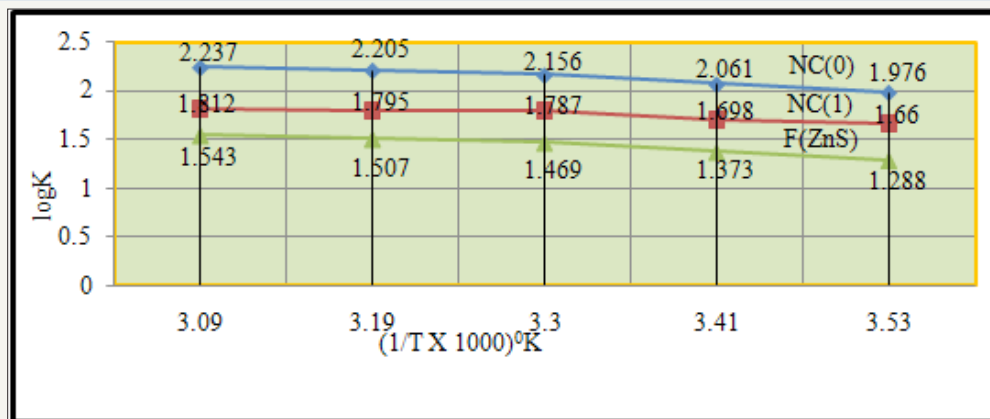


Figure 16: log K Vs 1/T for nanocoating NC (1) & ZnS filler on poly-butadiene-coated mild steel.

The octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime is an electronic rich compound and large molecular weight so it is suitable for nanocoating materials. Nanocoating compound is coordinated its electron to ZnS filler thus they can form strong barrier on the surface of poly-butadiene-coated mild steel. This barrier stops osmosis or diffusion process of saline water. Studied the effect of temperature on the poly-butadiene-coated mild steel at mentioned above temperatures and their results were written in Table 1, it observed that corrosion rate of material increased

without nanocoating but its values were reduced with nanocoating and filler compounds such types trends clearly noticed in Figure 16 which plotted between logK versus 1/T found to be a straight line. Nanocoating and filler compounds were formed a stable barrier with on the surface of poly-butadiene-coated mild steel. This barrier has thermal stability and suppressed the attack of Cl<sup>-</sup> ions. Nanocoating and filler materials were developed composite barrier i.e. stable in saline water as temperature increased.

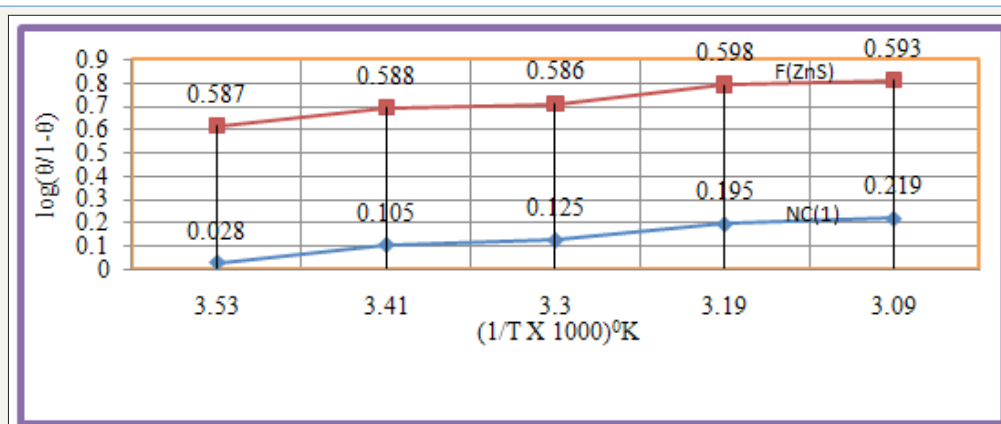


Figure 17: log ( $\theta/1-\theta$ ) Vs 1/T for nanocoating of NC (1) & ZnS filler on polybutadiene-coated mild steel.

The plot of log ( $\theta/1-\theta$ ) versus 1/T depicted a linear graph as shown in Figure 17. The plot of Figure 17 and Table 1 results confirmed that nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler increased log( $\theta/1-\theta$ ) as temperature enhanced. The values of log( $\theta/1-\theta$ ) increased with octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime but its values

more increased with ZnS filler in marine water system. The surface coverage area ( $\theta$ ) of nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler was calculated by equation  $\theta = (1 - K/K_0)$  (where K is the corrosion rate before coating and  $K_0$  is the corrosion rate after coating) and their values were mentioned in Table 1. Figure 18 plotted between surface coverage



(θ) versus temperature(T) which indicated that filler compound covered more surface area with respect of nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime. The low dose of nanocoating and filler compounds were occupied more surface

coverage area as temperatures were increased. These results were given information that as temperatures were risen nanocoating and filler compounds were accommodated more surface areas.

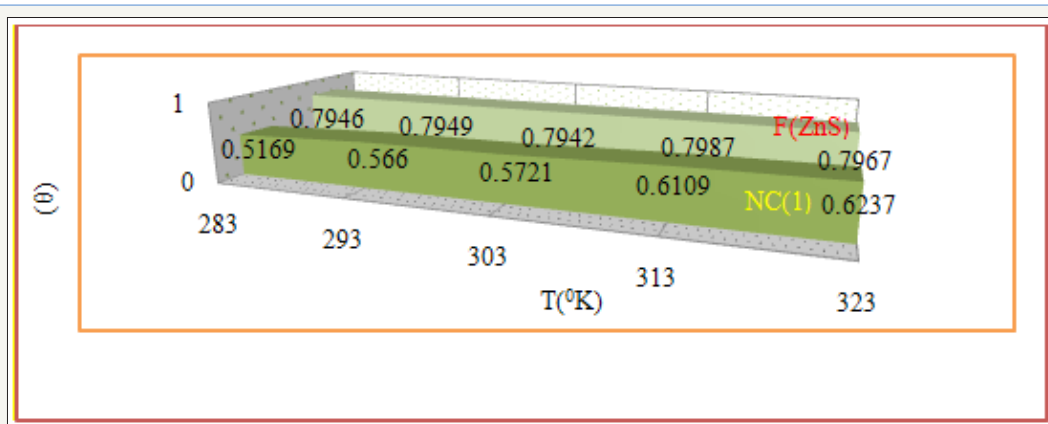


Figure 18: θ Vs T for nanocoating of NC (1) & ZnS filler on the poly-butadiene-coated mild steel.

The percentage coating efficiency for octahydrobenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler were calculated by equation, %CE=(1-K /K<sub>o</sub>) X 100 (where CE=coating efficiency, K=Corrosion rate after coating, K<sub>o</sub>=corrosion rate before coating) and their values were given in Table 1. Figure 19 plotted between percent CE (percentage coating efficiency) versus T (temperature) indicated that filler compound enhanced percentage coating efficiency with respect of nanocoating compound. Nanocating and filler compounds were electron rich compound so they have more binding capacities. Activation energy of polybutadiene-coated mild steel, nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and

ZnS filler were determined by Arrhenius equation, d/dt (log K)=E<sub>a</sub>/RT<sup>2</sup> (where T is temperature in Kelvin, R is universal gas constant and E<sub>a</sub> is the activation energy of the reaction) and Figure 16 which plotted between log K versus 1/T. The calculated values of activation energies were mentioned in Table 2. Poly-butadiene-coated mild steel produced high activation energies at different temperatures in marine water whereas nanocoating and filler compounds exhibited lower activation energy. The results of activation energies were shown that nanocoating and filler compounds adhered on the surface of poly-butadiene-coated mild steel by chemical bonding.

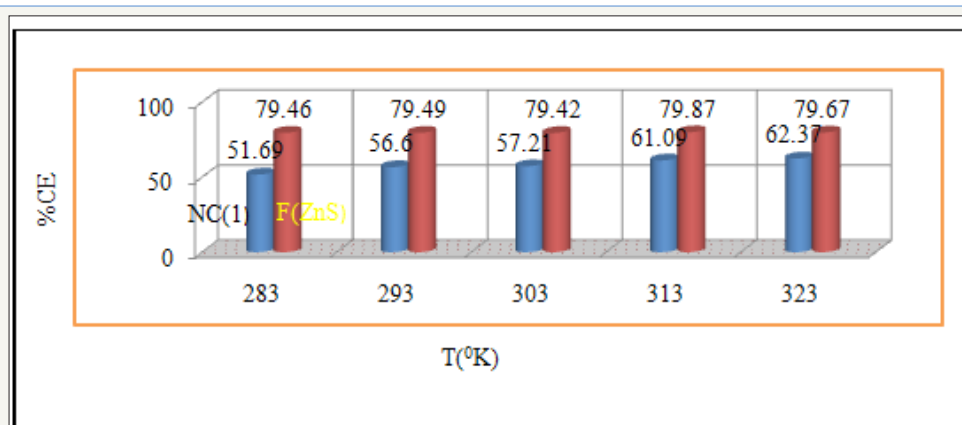


Figure 19: θ Vs T for nanocoating of NC (1) & ZnS filler on poly-butadiene-coated mild steel.

Table 2: Thermal parameters of nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler coated on poly-butadiene-coated mild steel in marine water

Thermal Parameters	283 °K	293 °K	303 °K	313 °K	323 °K
E <sub>a</sub> (0)	133.43	134.44	136.1	139.19	132.23
E <sub>a</sub> , NC(1)	113.15	110.76	112.81	109.53	107.11
q, NC(1)	-1.89	-6.84	-7.89	-11.89	-12.94
ΔG,NC(1)	-214.91	-210.13	-208.92	-202.45	-197.94

$\Delta H, NC(1)$	-75.11	-76.06	-80.07	-78.8	-78.11
$\Delta S, NC(1)$	-82.15	-83.92	-87.57	-88.16	-89.07
$\theta, NC(1)$	0.5169	0.566	0.5721	0.6109	0.6237
$E_a, NC(ZnS)$	86.97	89.56	92.73	91.96	91.2
$q, NC(ZnS)$	-39.63	-38.35	-36.99	-36.49	-35.05
$\Delta G, NC(ZnS)$	-189.82	-188.89	-188.87	-184.93	-181.23
$\Delta H$	-50.02	-54.83	-60.03	-61.28	-62.27
$NC(ZnS)$	-67.94	-71.47	-75.42	-61.28	-78.8
$\Delta S, NC(F)$	-67.94	-71.47	-75.42	-77.08	-78.8
$\theta, NC(ZnS)$	0.7946	0.7949	0.7942	0.7987	0.7967

Table 2 Thermal parameters of nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler coated on poly-butadiene-coated mild steel in marine water. Heat of adsorption of nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler were calculated by Langmuir equation,  $\log(\theta/1-\theta)=\log(A.C)-(q_{ads}/2.303RT)$  (where T is temperature in Kelvin and  $q_{ads}$  heat of adsorption) and Figure 17 plotted against  $\log(\theta/1-\theta)$  versus  $1/T$  which produced straight lines. Heat adsorption values found to be negative with nanocoating and filler compounds so the formation of composite barrier is

a chemical process. These compounds were adhered with base material by chemical bonding. Free energies of both compounds give information that coating is an exothermic process. It is clear by data calculated for octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler with the help of equation,  $\Delta G=-2.303RT \log(33.3K)$  (where R is universal gas constant, T be temperature and K corrosion rate) and their values were written in Table 2. Both compounds were produced a negative heat of adsorption which indicated that they formed chemical bonding during nanocoating process.

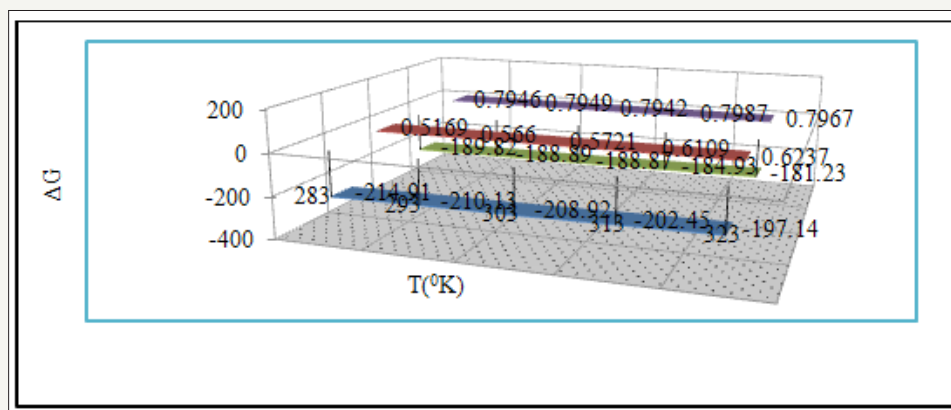


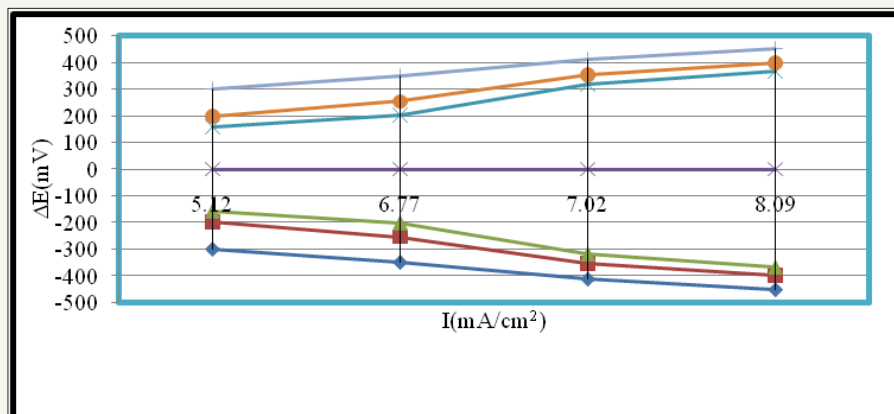
Figure 20:  $\Delta G$  Vs T for ( $\theta$ ) for nanocoating of NC (5) & ZnS filler on poly-butadiene-coating mild steel.

Enthalpy and entropy are very important thermal parameters which give information about nanocoating and filler compounds bonding nature with base materials. The values of enthalpy and entropy for octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler were calculated by transition state equation,  $K=RT/Nh \log(\Delta S^\ddagger/R) \times \log(-\Delta H^\ddagger/RT)$  (where N is Avogadro's constant, h is Planck's constant,  $\Delta S^\ddagger$  is the change of entropy activation and  $\Delta H^\ddagger$  is the change of enthalpy activation) and Figure 20 and their values were recorded in Table 2. The results of enthalpy and entropy were

shown that both compounds were adsorbed with the base materials by chemical bonding. Such coating is an exothermic process. The negative entropy indicated that nanocoating and filler compounds accommodated on the surface of poly-butadiene-coated mild steel in an ordered matrix. Figure 20 plotted between free energy ( $\Delta G$ ) and temperatures for surface coverage area ( $\theta$ ) occupied by octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler that confined free energy decreased when temperatures increased but at this moment surface coverage area was enhanced.

Table 3: Potentiostat polarization of octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS nanocoated on poly-butadiene-coated mild steel in marine water.

NC	$\Delta E$ (mV)	$\Delta I$	$\beta_a$	$\beta_c$	$I_{corr}$ (mA/cm <sup>2</sup> )	K (mmpy)	$\theta$	%CE	C(mM)
NC(0)	-451	112	172	151	8.69	264.69	0	0	0
NC(1)	-412	99	112	165	7.02	213.82	0.71	71	50
F(ZnS)	-350	85	101	175	6.77	206.21	0.88	88	10



**Figure 21:**  $\Delta E$  (mV) Vs  $I$  (mA/cm<sup>2</sup>) for nanocoating of NC (1) & filler ZnS on poly-butadiene-coated mild steel.

The corrosion potential, corrosion current and corrosion current density of poly-butadiene-coated mild steel, nanocoated octahydrodibenzo [a,d] [8] annulene and ZnS filler were calculated by equation,  $\Delta E/\Delta I = \beta_a \beta_c / 2.303 I_{\text{corr}} (\beta_a + \beta_c)$  (where  $\Delta E/\Delta I$  is the slope which linear polarization resistance ( $R_p$ ),  $\beta_a$  and  $\beta_c$  are anodic and cathodic Tafel slope respectively and  $I_{\text{corr}}$  is the corrosion current density in mA/cm<sup>2</sup>) and Tafel plot between electrode potential ( $\Delta E$ ) versus current density ( $I$ ) and their values were mentioned. It was observed that electrode potential and corrosion current density were high with polybutadiene whereas anodic current density increased and cathodic density current reduced. But octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and ZnS filler reduced electrode potential and corrosion current density. Tafel plot of Figure 21 and the results of Table 3 indicated that nanocoating and filler compounds minimized anodic current and maximized cathodic current. These results confirmed that nanocoating and filler compounds developed a strong barrier on the poly-butadiene-coated mild steel which neutralized the attack of saline water and carbonic acid.

The corrosion current was obtained by above equation and Tafel graph of Figure 21 for poly-butadiene-coated mild steel, nanocoated octahydrodibenzo [a,d] [8] annulene and ZnS filler and these values were put in equation,  $C. R (\text{mmpy}) = 0.1288 I (\text{mA/cm}^2) \times \text{Eq. Wt} (\text{g}) / \rho (\text{g/cm}^3)$  (where  $I$  is the corrosion current density  $\rho$  is specimen density and Eq. Wt is specimen equivalent weight) and their values were recorded in Table 3. It was noticed that corrosion rate increased with poly-butadiene-coated mild steel and these values were reduced with nanocoating and filler compounds. The results of corrosion rate measured by weight loss experiment which confirmed the results of potentiostat.

## Conclusion

It is very difficult to control corrosion of marine water. Poly-butadiene-coated mild steel uses in marine water for different works but this material is face corrosion problem. In this research, it is tried to check corrosion of poly-butadiene mild steel by the application of nanocoating compound octahydrodibenzo [a,d] [8] annulene-5, 12-dioxime and filler ZnS. The corrosion activities of poly-butadiene was studied at 283 °K, 293 °K, 303 °K, 313 °K

and 323 °K temperatures and the concentration of nanocoating and filler compounds were taken in 50mM and 10mM. The results of surface coverage areas and coating efficiencies of nanocoating and filler compounds were indicated these compounds had more coverage capability. Nanocoating and filler compounds results of activation energy, heat of adsorption, free energy, enthalpy and entropy were shown that these compounds were attached with base material by chemical bonding. They can form composite thin film barrier which is passive in corrosive environment. Filler material blocks the porosities of nanocoating compounds and stop osmosis or diffusion process of corrosive agents.

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