

Excellent Anti-bacterial Activity of Poly(o-toluidine)-DBSA/ ZnO Nanocomposite

Dalal K. Thbayh*¹ Rawnaq A. Talib¹ E. H. Al – Tememe^{1,2}

1.University of Basra, Polymer Research Canter, Basra, Iraq

2.iversity of Basra, Pharmacy College, Basra, Iraq

Abstract

This work presents a study of the biological application (antibacterial activity) of ZnO/poly(o-toluidine) (POT) doped with organic acids dodecylbenzene sulfonate acid (DBSA) nanocomposites synthesized by in-situ polymerization of (o-toluidine) monomer in presence of 5% ZnO. The FTIR spectroscopy confirms the existence of an interaction between POT-DBSA matrix and ZnO particles. Scanning electron microscopy reveals the nanostructure nature of the obtained composite. The antibacterial activity of POT-DBSA/ZnO nanocomposite and POT-DBSA studied by agar well diffusion method, was found to increase with increasing concentration meanwhile POT/DBSA/ZnO exhibits better antibacterial activity compared to POT/DBSA and POT separately.

Keywords: Poly(O-toluidine); Organic acid; ZnO; nanocomposite; antibacterial activity.

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1. Introduction

Synthesizing and doping poly (o-toluidine) with organic acid such as DBSA (dodecylbenzene sulfonate acid) is useful for various applications. Poly (o-toluidine) (POT) as dopant is one of the most conducting polymer promising to substitute polyaniline (D. Macinnes 1988, Watanabe, Mori et al. 1989, Wei, Focke et al. 1989) due to its good chemical stability with high conductivity, easy polymerization, since polyaniline shows difficult solubility in majority of solvents. Therefore, POT doped with DBSA containing long alkyl chains could be soluble in common solvents like chloroform, xylene, m-cresol and toluene (T. Taka 1994, Athawale, Kulkarni et al. 2002, Ebrahim, Kashyout et al. 2009). To solve the problem of poor chemical, thermal and mechanical properties of conducting polymers for particular applications, a considerable amount of research work has been devoted to prepare hybrid inorganic/organic composites, which are formed by inserting inorganic particles to the polymer chain structure in order to obtain unique properties (Khan, Khalid et al. 2010, Ansari and Mohammad 2011, Khan, Baig et al. 2011, Khan and Baig 2012, Khan and Baig 2013, Khan and Baig 2013).

Zinc oxide (ZnO) considered as one of the versatile inorganic compound belonging to II–VI group compound, is an n-type semiconductor with a direct wide band gap (3.37 eV) and a large exciton binding energy at room temperature (about 60 meV) (Sui, Shao et al. 2007).

ZnO-based nanostructured materials represent an important class of semi conductive oxides that can be used in many applications because of its low cost with a diversity of forms, such as nanorods (Park, Kim et al. 2002, Hassan, Hassan et al. 2011), nanofibers, nanoparticles, nanobelts, nanoflowers (Wang, Qian et al. 2004, Izaki, Watanabe et al. 2008) and nanowires (Zhang, Ram et al. 2012). G. Sudhana and co-workers (Sudhana, Thenmozhi et al. 2012) studied the antibacterial activity of POT prepared by chemical oxidation method in the presence / absence of ultrasonic irradiation, it's results shows the antibacterial activity and conductivity of POT prepared in the presence of ultrasonic irradiation is better than its absence. In recent decades, many works have been synthesized of polymer-metallic nanocomposite and studied their antibacterial activities (Khan, Radhakrishnan et al. 2018).

This work involves preparation of composites from inorganic 5% weight ratios ZnO nanoparticles and organic conducting polymer of poly (o-toluidine) doped with dodecylbenzene sulfonate (POT-DBSA) by in-situ chemical oxidative polymerization method. The inhibition zone experiment was applied to evaluate the antibacterial activity of POT-DBSA/ZnO nanocomposite and POT-DBSA by using three types of bacteria *Escherichia Coli* and *Pseudomonas aeruginosa* (Gram negative) and *Staphylococcus aureus* (Gram positive), respectively.

2. Experimental

2.1 Synthesis of poly-o-toluidine doped with dodecyl benzene sulfonic acid (POT-DBSA)

The polymerization of the monomer o-toluidine (Fisher scientific company, USA), was initiated by the drop wise addition of the oxidizing agent, (5.8 gm, 1M) ammonium persulphate (APS) (NH₄)₂S₂O₈ (Sigma Aldrich, USA). 5.4 gm organic acid Dodecylbenzene sulfonate (DBSA) (Sigma Aldrich, USA) was used for doping POT. The POT to ammonium persulphate mixing in similar volume ratios is achieved by continuous stirring using a magnetic stirrer for 24 h. A green colored gel was obtained (Ahmad, Riaz et al. 2012, Khan and Baig 2013, Thbayh DK 2018) which was filtered, and repeatedly washed with distilled water to remove the excess of acid and impurities then finally dried at 70°C for 6 h in an oven. The final product was a green powder (POT-DBSA).

2.2 Synthesis of POT-DBSA/ ZnO nanocomposite

POT-DBSA/ZnO nanocomposite was prepared by in situ oxidative polymerization of o-toluidine in the presence of ZnO (Sigma Aldrich, USA) nanoparticles (NPs). 5wt% of ZnO NPs were poured into solution of o-toluidine-DBSA and polymerization was affected by the addition of solution containing oxidant ammonium persulphate prepared in 1 M HCl. The addition of oxidant solution led to the polymerization of adsorbed o-toluidine on ZnO NPs resulting in the appearance of light greenish black precipitate colored solution which was kept under continuous stirring for 24 h. The reaction mixture was then filtered, washed with double distilled water to remove excess acid. The nanocomposite thus prepared was dried at 70°C for 6 h in an oven, converted into powders and stored in desiccator for further investigations.

2.3 Antibacterial activity

The Muller Hinton agar was used for the growth of *Escherichia coli* and *Pseudomonas aeruginosa* (Gram negative) and *Staphylococcus aureus* (Gram positive) bacteria to study the antibacterial activity of POT-DBSA/ ZnO nanocomposite and POT-DBSA, by using the diffusion in the agar method. The bacterial strain was cultured on the nutrient agar broth to produce a new colony of age 24 h, then to obtain a bacterial growth one new colony from the growth was inoculated on a 5 ml of nutrient broth and was incubated in 37°C for 6 h. A 0.1 ml from the growth was separated on the Muller-Hinton agar. The inoculated agar surface was poured in 4 mm diameter for every pore. The pores of cultured media were filled by 100, 200, 300, 400 and 500 µg/ml of POT-DBSA/ZnO nanocomposite and POT-DBSA, the controlled solution was 10 ml of Dimethyl- sulphoxide. Then the dishes were kept in the incubator at 37°C for a period of 18-24 h. The inhibition zone diameters in mm were measured around the pores.

2.4 Characterizations

The morphology of nano composites films was investigated by field emission scanning electron microscopy (FESEM) using FEINova Nano SEM 450. Fourier transform infrared (FT-IR) spectra were recorded on FT-IR 4200 Jasco.

3. Results and Discussion

3.1 FTIR spectroscopy analysis

The FT-IR spectrum of the POT-DBSA/ (5 wt% ZnO) nanocomposite is shown in Fig 1. The characteristic bands of POT-DBSA are as follows: the peak at 1604 cm⁻¹ is ascribed to the stretching vibration of quinoid rings; the peak at 1406 cm⁻¹ depicts the presence of benzenoid ring units (Kulkarni, Viswanath et al. 2004, Ebrahim, Gad et al. 2010); the peak at 1170 cm⁻¹ is assigned to a vibration band of the dopant ion, which is formed during protonation of POT with DBSA (Rao, Sathyanarayana et al. 2002, Kulkarni, Viswanath et al. 2004); the peak at 3121 cm⁻¹ is attributed to the free N-H stretching vibration (Kulkarni, Viswanath et al. 2004), the peak at 1004 cm⁻¹ corresponds to the C-H in plan bending vibration (Kulkarni, Viswanath et al. 2004); while the peaks at 682 and 582 cm⁻¹ are associated to C-H out of plane bending vibration (Kulkarni, Viswanath et al. 2004) and C-N-C bonding mode of aromatic ring, respectively. The characteristic peak at 436 cm⁻¹ belongs to the Zn-O stretching mode (Hamedani and Farzaneh 2006, Kakazey, Vlasova et al. 2007, Geetha and Thilagavathi 2010, Zhao, Cai et al. 2011, Nosrati, Olad et al. 2012). The frequency data obtained, and their assignments are presented in Table 1.

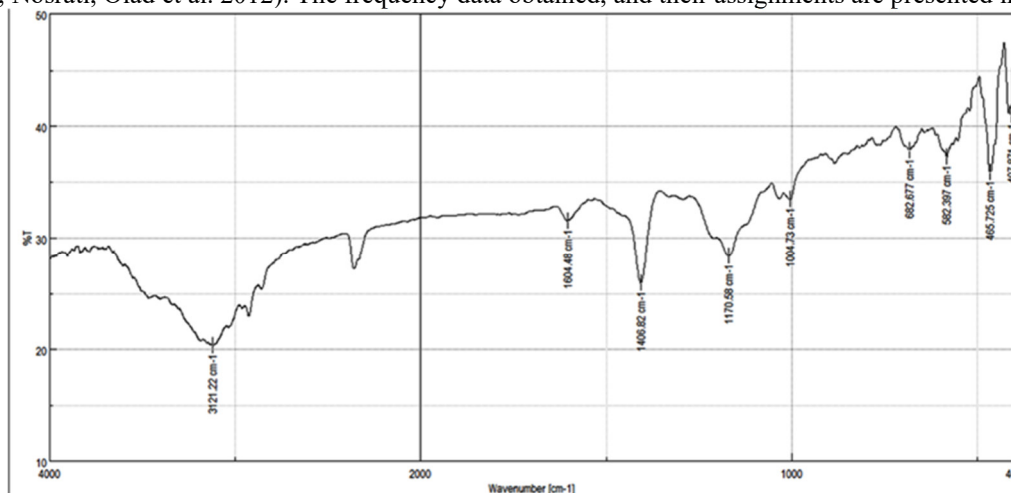


Figure 1. FTIR spectrum of POT-DBSA/5% ZnO nanocomposite.

Table 1. Characteristic frequencies of in-situ chemically synthesized modes of the POT-DBSA/5% ZnO nanocomposite.

Assignment	Wave number (cm ⁻¹) POT-DBSA/5%ZnO
Zn-O	407.8 465.7
C-N-C bonding mode of aromatic ring	582.3
C-H out of plane bonding in benzenoid ring	682.6
C -H in plane bonding	1004.7
Vibration band of the dopant ion	1170.5
C = C stretching mode of the benzenoid rings	1406.8
C=C stretching mode of the quinoid rings	1604.4
H-N stretching vibration	3121.2

3.2 FESEM and EDX analysis

Figure 2 shows the basic chemical structure of POT, DBSA, and ZnO. The EDX spectrum of POT-DBSA/5% ZnO shown in the Table 2, reveals the presence of Zn, O, S, and C elements. It can be deduced that the presence of Zn element in composite results from the interaction of POT-DBSA with ZnO nanoparticles. The SEM image of nanocomposite film is shown in Fig 3. It can be observed that the crystals of ZnO NPs are fully covered by POT and the composite shows a globular-type nanostructure consisting of small globules and pores. This result indicates that ZnO NPs are encapsulated into the POT matrix hence can significantly improve the compactness of the polymer. The average diameter of POT-DBSA/ZnO grains is found to be 37 nm.

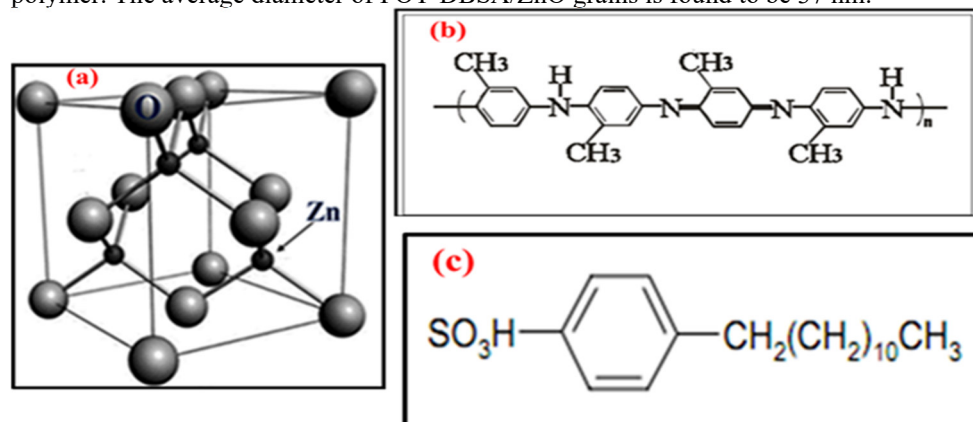


Figure 2. The basic chemical structure of (a) ZnO, (b) POT, and (c) DBSA (Vaseem, Umar et al. 2010, Jagadish and Pearton 2011).

Table 2. EDX data of POT-DBSA/ 5% ZnO

Elt	W%	A%
C	62.02	75.49
O	17.77	16.23
Al	1.11	0.60
S	14.56	6.64
Fe	0.68	0.18
Zn	3.87	0.86
	100.00	100.00

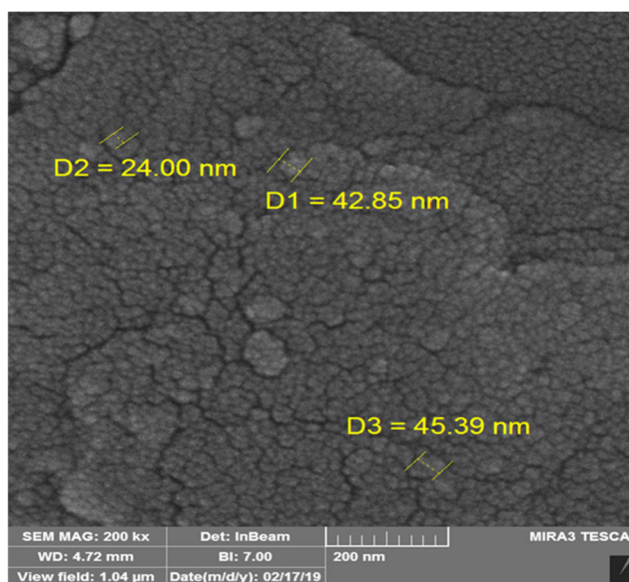


Figure 3. FESEM images of POT-DBSA/5% ZnO nanocomposite.

3.3 Determination the inhibition zone diameter

The inhibition zone diameters against *Staphylococcus aureus*, *Escherichia coli* and *Pseudomonas aerogmosa* bacteria by POT-DBSA/ZnO nanocomposite and POT-DBSA which were compared with POT taken separately and with standard drug. There are four types of antibacterial activities (R. Kavitha 2011): (i) the inhibition zone diameter $\gg 12$ mm resulting in high sensitive bacteria; (ii) moderate sensitive bacteria for the inhibition zone diameter in the range 9-12 mm; (iii) if the inhibition zone diameter is in the range 6-9 mm, then the sensitivity of bacteria is less; and (iv) bacteria becomes resistant if then inhibition zone diameter is < 6 mm. From Table 3 and photographic pictures shown in Figure 4, it can be indicated that the inhibition zone diameter is highest for POT-DBSA/ZnO nanocomposite and POT-DBSA; and that the inhibition zone diameter increases as the concentration of the nanocomposite increases.

From Table 3 the following points can be highlighted:

1. POT-DBSA/ZnO nanocomposite exhibits a better antibacterial activity than POT-DBSA, POT and Oxytetracyclin drug.
2. The inhibition zone diameter against *Pseudomonas aerogmosa* and *Escherichia coli* (gram negative) is relatively higher than against *Staphylococcus aureus* (gram positive) due to the difference in the cell wall between gram positive and gram-negative bacteria. The increase in the inhibition zone diameter may be due to the presence of ZnO NPs in POT-DBSA/ZnO nanocomposite because the antibacterial activity has been reported to be size dependent (Boomi, Prabu et al. 2014), then the bacterial death due to electromagnetic

attraction between the bacteria (negative charge) and ZnO nanocomposite (positive charge) (A. Thomas1 2014). In addition, the effect of DBSA in POT-DBSA may destruct bacterial enzymes by coordinating to the electron donating groups, thus DBSA causes deformities in bacterial cell wall to render them leaky (Shakir, Khan et al. 2014).

Table 3. The inhibition zone diameter of three types of bacteria by POT-DBSA/ZnO nanocomposite, POT-DBSA, POT and drug.

POT-DBSA/ZnO nanocomposite				POT-DBSA					
	Conc. $\mu\text{g/ml}$	Staph	E. coli	Psed.		Conc. $\mu\text{g/ml}$	Staph.	E. coli	Psed.
5	100	10	18	18	10	100	6	12	16
4	200	12	20	20	9	200	8	16	18
3	300	16	22	21	8	300	9	20	19
2	400	18	24	22	7	400	10	22	20
1	500	20	28	25	6	500	12	25	22
POT		10	17	10					
Oxytetracyclin		16	24	22					

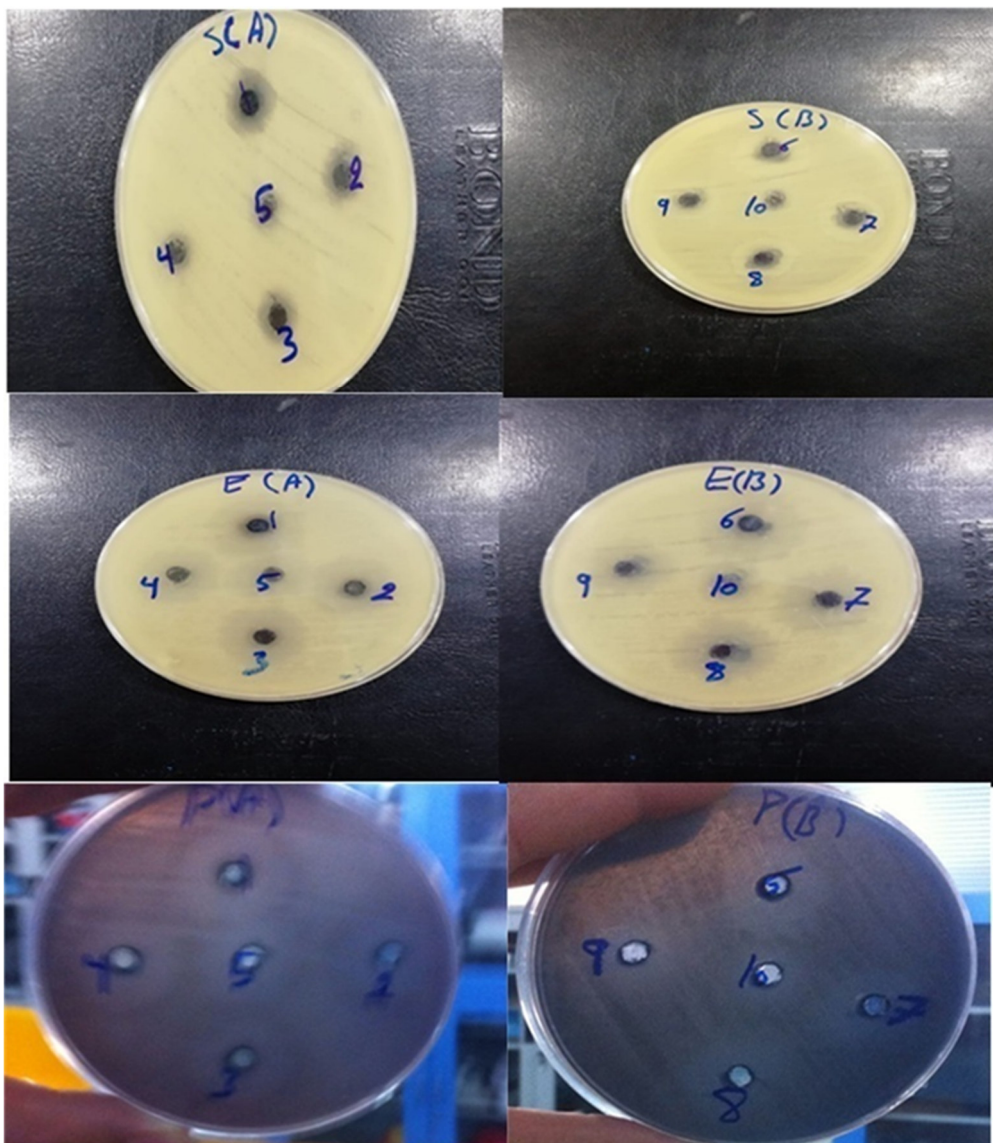


Figure4. Photographic the inhibition zone diameter of (A) POT-DBSA/ZnO nanocomposite and (B) POT-DBSA against (S) Staphylococcus aureus bacteria, (E) Escherichia coli bacteria and (P) Pseudomonas aeruginosa bacteria.

4. Conclusion

POT- DBSA/5% ZnO were successfully prepared using in-situ polymerization of (o-toluidine) monomer in the presence of 5% weight ratio of ZnO NPs. Characteristic bands of ZnO and POT-DBSA were identified by FTIR and confirming the interaction between ZnO and polymer matrix. FESEM observations reveal rock salt structure embedded within the polymeric structure, signifying that ZnO NPs are fully covered by POT, with an average diameter of POT-DBSA/ZnO grains around 37 nm. The comparative study of biological application (antibacterial activities) of POT, POT-DBSA, POT-DBSA/ZnO nanocomposite and oxytetracyclin drug have been investigated. The results confirm that POT-DBSA/ZnO nanocomposite exhibits enhanced antibacterial activity against E.coli and Pseud. aerogmosa Gram negative and Staph. aureus Gram positive bacteria, as well that the inhibition growth of bacteria increases when POT-DBSA are coordinated with ZnO NPs. The antibacterial properties could make POT-DBSA/ZnO nanocomposite potentially useful as novel drug, thereby future in-vivo measurements are underway.

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