

# Radical Scavenging and Endogenous Defence System Inducing Activities of 5-[(4-Chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol: A Novel Antioxidant

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## Shehzadi, *et al.*: Broad Spectrum Radical Scavenging Effect of 5-[(4-Chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol

5-[(4-Chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol owing to the presence of –SH group is expected to have a significant reducing potential, which could be translated into antioxidant potential and to prove this, the present study explored the antioxidant activity and binding pattern of this compound to oxidative stress-related protein targets. The antioxidant properties were determined using *in vitro* methods with ascorbic acid and butylated hydroxytoluene as standards while interactions of 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol with protein tyrosine kinase 2- $\beta$  and glutathione reductase were determined using online software, Mcule 1-Click Docking, 3DLigandSite and COACH. The antioxidant activity of 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol was found to be 89.30 $\pm$ 0.013, 81.20 $\pm$ 0.002, 80.52 $\pm$ 0.016, 54.81 $\pm$ 0.007, 52.87 $\pm$ 0.008, 34.44 $\pm$ 0.019 and 19.91 $\pm$ 0.014 % in hydrogen peroxide scavenging assay, 2,2-diphenyl-1-picrylhydrazyl radical scavenging assay, phosphomolybdenum assay, nitric oxide scavenging assay, reducing power assay, ferric thiocyanate assay and  $\beta$ -carotene bleaching assay, respectively. In all these assays, EC<sub>50</sub> of 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol ranged from 0.32-0.93 mg/ml. The docking results indicated excellent binding to protein tyrosine kinase 2- $\beta$  and glutathione reductase, maximum for the latter. The results of the present study revealed that 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol has the propensity to abrogate oxidation by inducing endogenous defence system and preventing radical chain reactions, hence might be considered a potential antioxidant for further investigations.

**Key words:** Antioxidant, molecular docking, glutathione reductase, protein tyrosine kinase 2- $\beta$ , 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol

Free radicals (FRs), species having unpaired electron in their outermost orbit play a pivotal role in the origin of life, physiological adaptation and biological evolution<sup>[1]</sup>. Many cellular biochemical processes, signal transduction, gene transcription and regulation of soluble guanylate are governed by FRs. Nitric oxide free radical (NO $\cdot$ ) is a very important signalling species that regulates relaxation and proliferation of vascular smooth muscles, leukocytes adhesion, platelets aggregation, angiogenesis, thrombosis, vascular tone and haemodynamics<sup>[2]</sup>.

Mammalian cells being constantly fortified with oxygen produce oxygen free radicals (OFR), hydroxyl (OH $\cdot$ ) and superoxide (O<sub>2</sub> $\cdot^-$ ), and reactive oxygen species (ROS), NO $\cdot$ , hypochlorous acid, hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) and

peroxynitrite as a by-product of aerobic metabolism<sup>[3,4]</sup>. Since, the rate of production and removal of FRs is balanced by reactions involving enzymes such as catalase, glutathione peroxidase, glutathione reductase and superoxide dismutase, and non-enzymatic processes such as thioredoxin, thiols and disulphide-bonding, these species do not necessarily present any threat to the body under normal physiological conditions<sup>[5]</sup>. Nevertheless, an imbalanced defence or

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Accepted 15 October 2018

Revised 08 April 2018

Received 27 May 2017

Indian J Pharm Sci 2018;80(6):1125-1135

overproduction, under-elimination and assimilation of FRs from the exogenous sources (electromagnetic radiation, pollution and smoking) can cause serious penalty to the biomolecules leading to inflammation, tissue injury and ultimately cell death. Long term oxidative damage is known to be the culprit cause of aging, cancer, neural and cardiovascular disorders, and liver diseases<sup>[6-12]</sup>. Protection against injury due to the FRs is possible only by ample intake of antioxidants, which delay or inhibit the oxidation of biomolecules by their reducing activity, free radical scavenging, complexation of pro-oxidant metals, and quenching of singlet oxygen. Such compounds not only prevent or postpone the onset of degenerative diseases but also improve the quality of life and save the cost of healthcare delivery.

5-[(4-Chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol (OXCPM, fig. 1) belongs to an important class of heterocyclic compounds due to the presence of nitrogen, oxygen and exocyclic sulphur atoms. Oxadiazole ring containing systems have remained the focus in the drug discovery programs because of their versatile nature, privileged structure (reactive pharmacophore), molecular modulating properties, ligand-binding site interactions and pharmacological profile<sup>[13]</sup>. This is especially true for the substituted 1,3,4-oxadiazoles that are reported to have a broad-spectrum pharmacological activities like antibacterial, antitubercular, antifungal, antiviral, anthelmintic, antioxidant, anticancer, central nervous system depressing, anticonvulsant, muscle relaxant, hypoglycaemic, analgesic, antiinflammatory and genotoxic<sup>[14-38]</sup>. In the recent past, OXCPM was synthesized and evaluated for drug-like characteristics using various *in silico* models<sup>[39,40]</sup>. Due to thiol group, the OXCPM is expected to have a significant reducing potential, which could translate into novel antioxidant effects and to prove this, the present study is attempted to explore the antioxidant activity and binding pattern of OXCPM to oxidative stress-related protein targets.

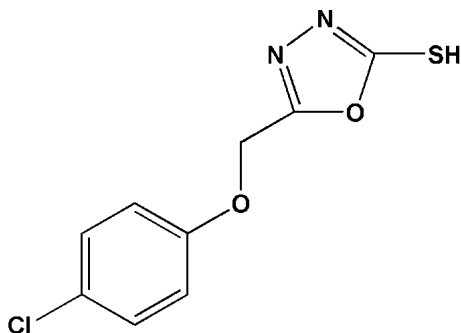


Fig. 1: Chemical structure of 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol

## MATERIAL AND METHODS

The chemicals used and their sources are as follows. Phosphate-buffered saline tablets (PBS404.100, BioShop Canada Inc., Burlington), butylated hydroxytoluene (BHT), Tween-40, acetic acid (100 %), hydrochloric acid, trichloroacetic acid, ferric chloride, disodium hydrogen orthophosphate (BDH Laboratory Supplies, Poole, England),  $\beta$ -carotene (BC), potassium ferricyanide, sodium hydroxide, ammonium heptamolybdate, potassium dihydrogen phosphate, sodium dihydrogen phosphate (Merck KGaA, Darmstadt, Germany), sodium nitroprusside dihydrate, sulfanilic acid, sulphuric acid, ammonium thiocyanate (Riedel-deHaen AG. D-3016, Seize, Germany), ascorbic acid (AA), linoleic acid (99 %),  $\alpha, \alpha$ -diphenyl- $\beta$ -picrylhydrazyl (DPPH), chloroform,  $H_2O_2$  (30 % v/v) by Sigma-Aldrich Chemie, Steinheim, Germany, naphthylethylenediamine dichloride (MP Biomedicals Inc., Illkirch, France) and ferrous chloride (Uni-Chem Chemical Reagents, Karachi, Pakistan) were procured from the local chemical market. The solvents used in the study included methanol (HPLC grade, Tedia Company Inc., Fairfield, USA) and in-house prepared double-distilled water (DDW). OXCPM was received as a gift sample from Department of Chemistry, Government College University, Lahore, Pakistan.

The equipment used were, refrigerator centrifuge (2-16KC, Sigma Laborzentrifugen, Germany), sonicator (DSA50-SK1-1.8L, Germany), thermostatic water bath (M 270, Memmert, Germany), thermostatic oven (U10, Memmert, Germany), pH meter (Hanna Instruments, Romania), rotary evaporator (HEI-VAP series, Heidolph, Germany), hot plate (S37-988, Griffin and George Ltd., Britain) and ultra-low chiller (MOF U32V) and incubator (MIR-153) by Sanyo Electric Co. Ltd., Japan. The absorbance of the reaction mixtures were recorded in 1 cm matching quartz cell using a UV/Vis spectrophotometer (T70<sup>+</sup>, PG Instruments, UK).

### Preparation of standard and sample solutions:

The stock solutions of AA, BHT and OXCPM of the strength 1.00 mg/ml were prepared in methanol. A range of working solutions of OXCPM (0.10-1.00 mg/ml) was prepared by diluting the stock solution with methanol.

### DPPH radical scavenging assay:

This assay was performed using a method described by Manzocco *et al.* with slight modifications<sup>[41]</sup>. Briefly, an

aliquot of 0.20 ml of the sample (0.10-1.00 mg/ml) was mixed with 2 ml of 0.50 mM solution of DPPH radical (DPPH $\cdot$ ) in methanol. The mixture was incubated in the dark at room temperature for 30 min and the absorbance was recorded at 517 nm against methanol as a blank. AA and BHT were used as the standards and the test tubes containing methanol in the place of the sample served as a control. The % radical scavenging activity was calculated using the following Eqn., % DPPH $\cdot$  scavenging activity =  $((A_c - A_s)/A_c) \times 100$ , where,  $A_c$  and  $A_s$  denote the absorbance of control and sample, respectively.

#### NO $\cdot$ scavenging assay:

The method of Marcocci *et al.*, with some modifications was used to evaluate the ability of the OXCPM to scavenge NO $\cdot$ .<sup>[42]</sup> A reaction mixture containing 3 ml sodium nitroprusside (10 mM) in PBS (pH 7.40) and 1 ml sample (0.10-1.00 mg/ml) was incubated at 25 $^\circ$  for 150 min under a visible polychromatic light source (100 W tungsten lamp). Finally, 0.50 ml of the incubated solution was mixed with equal volume of Griess reagent (mixture of equal volumes of sulfanilic acid, 0.33 % in 20 % glacial acetic acid and 0.10 % w/v naphthylethylenediamine dichloride) and incubated for further 30 min. The absorbance of the resulting pink-colored mixture was recorded at 546 nm against DDW as a blank. AA and BHT were used as the standards and the test tubes containing methanol in the place of OXCPM served as a control. The amount of NO $\cdot$  scavenged was calculated using the following Eqn., % NO $\cdot$  scavenging activity =  $[(A_0 - A_1)/A_0] \times 100$ , where,  $A_0$  and  $A_1$  denote the absorbance of mixture before and after reaction with Griess reagent.

#### H $_2$ O $_2$ scavenging assay:

The H $_2$ O $_2$  scavenging assay was performed as described earlier<sup>[43]</sup>. The reaction mixture containing 1 ml of OXCPM (0.10-1.00 mg/ml), 3.40 ml phosphate buffer (50 mM, pH 7.40) and 0.60 ml H $_2$ O $_2$  solution (40 mM, prepared in buffer) was incubated at room temperature. The absorbance was recorded at 240 nm at 0 min and after every 10 min for 40 min. A similar reaction mixture containing the buffer in place of H $_2$ O $_2$  was prepared for background subtraction. AA and BHT were used as the standards and the test tubes containing methanol in the place of OXCPM served as a control. The % H $_2$ O $_2$  scavenging effect was calculated using the following Eqn., % H $_2$ O $_2$  scavenging activity =  $(A_c - A_s)/A_c \times 100$ .

#### Reducing power assay:

The assay was performed using a method reported previously<sup>[44]</sup>. Briefly, a reaction mixture containing 2.50 ml of phosphate buffer (20 mM, pH 6.60), 2.50 ml of potassium ferricyanide (1 % w/v) and 1 ml of sample (0.10-1.00 mg/ml) or standards (AA and BHT) or control (methanol) was incubated at 50 $^\circ$  for 20 min followed by addition of trichloroacetic acid (2.50 ml, 10 % w/v). The resulting mixture was centrifuged at 3000 rpm for 10 min and the supernatant (2.50 ml) was collected and mixed with equal volume of distilled water. Finally, the reaction was stopped by addition of ferric chloride (0.50 ml, 0.10 % w/v) and the resulting colored complex was analysed at 700 nm against DDW as the blank. Increase in absorbance of the reaction mixture was taken as a direct measure of the reducing power of the sample<sup>[45]</sup>. The percent reducing power of the OXCPM was calculated using the following Eqn., % reducing power =  $(A_s - A_c)/A_s \times 100$ .

#### Phosphomolybdenum assay:

Phosphomolybdenum assay was performed as described by Prieto *et al.*<sup>[46]</sup>. Briefly, 0.10 ml of OXCPM (0.10-1.00 mg/ml) or standards (AA and BHT) or control (methanol) was treated with 1 ml of the molybdate reagent (0.60 M sulfuric acid, 28 mM sodium phosphate and 4 mM ammonium molybdate in distilled water). The tubes were incubated in a thermostatic water bath at 95 $^\circ$  for 90 min, cooled to room temperature and absorbance of the mixture was recorded at 695 nm against a suitable blank. Increase in the absorption of the reaction mixture indicated the antioxidant activity. The percent antioxidant activity of the OXCPM was calculated using the following Eqn., % antioxidant activity =  $(A_s - A_c)/A_s \times 100$ .

#### BC bleaching assay:

BC bleaching assay was performed by the methods of Velioglu *et al.*<sup>[47]</sup> and Lu and Foo<sup>[48]</sup>. Briefly, BC (100.00 mg) was dissolved in 50 ml chloroform (2.00 mg/ml) and added into a flask containing 1.00 g linoleic acid and 10.00 g Tween-40. The organic solvent was completely evaporated from the mixture using a rotary evaporator at 40 $^\circ$  followed by addition of 50 ml PBS to form a stable emulsion. Aliquots (4.70 ml) of the emulsion were added in test tubes containing 0.20 ml of sample (0.10-1.00 mg/ml), incubated at 50 $^\circ$  in a thermostatic water bath, and the absorbance of the reaction mixture was measured immediately ( $t=0$ ) and after every 30 min up to 90

min at 470 nm. The blank consisted of an emulsion prepared in similar manner as mentioned above but devoid of BC. The percent antioxidant activity was calculated using the following Eqn., % antioxidant activity =  $1 - [(A_{T_0} - A_{T_{90}}) / (A_{C_0} - A_{C_{90}}) + (A_{S_0} - A_{S_{90}})] \times 100$ , where,  $A_{C_0}$  and  $A_{C_{90}}$ ,  $A_{T_0}$  and  $A_{T_{90}}$ , and  $A_{S_0}$  and  $A_{S_{90}}$  are the absorbance values of the control (methanol), test sample and standards (AA and BHT), at zero time and after incubation for 90 min, respectively.

### Ferric thiocyanate assay:

The method described by Kikuzaki *et al.*<sup>[49]</sup> was employed to determine the effect of OXCPM on lipid peroxidation. In the screw cap vials, a reaction mixture containing 4 ml of the test concentration of OXCPM (0.10-1.00 mg/ml), 4.10 ml of 2.51 % linoleic acid (in methanol), 8 ml of phosphate buffer (50 mM, pH 7.00) and 3.90 ml of DDW was incubated at 40° in the dark. Aliquots of 0.10 ml of the reaction mixture were transferred to the test tubes containing 9.70 ml methanol (100 %), 0.10 ml 30 % aqueous ammonium thiocyanate and 0.10 ml 20 mM ferrous chloride in 3.50 % hydrochloric acid. The absorbance of the resulting mixture was measured at 500 nm after 3 min of addition of ferrous chloride. The measurements were made at 6 h and after every 24 h until the absorbance of the control reached its maximum. AA and BHT were used as positive control, and the mixture containing methanol in place of the sample served as a negative control.

### Statistical analyses:

Each assay was performed in replicates and the results were presented as mean  $\pm$  standard deviation (SD). The data were analysed using one-way ANOVA and difference between groups was determined by post-hoc multiple comparison, Tukey's HSD, using SPSS version 12.00. A p-value below 0.05 was considered as significant difference.

### Binding interactions of OXCPM with oxidative stress-related protein targets:

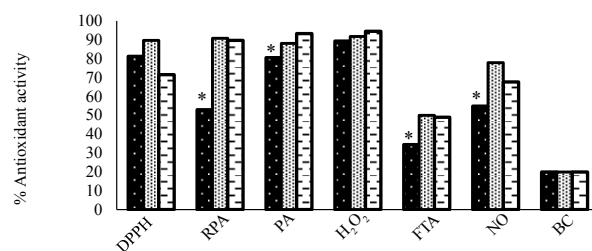
The binding of the ligand (OXCPM) with two oxidative stress-related targets, protein tyrosine kinase 2- $\beta$  (PTK-2 $\beta$ ) and glutathione reductase (GR) was evaluated using online docking software, 1-Click Docking by Mcule, Inc., USA. The OXCPM was entered as a SMILE in the input section of the software and refined in the drawer. Afterwards, the protein target was selected from the library and docking was performed. The results were obtained as "scores" of OXCPM binding affinity with

the selected targets. Moreover, the pose of the binding of the OXCPM in/on the protein surface was visualized and saved to determine the best binding conformation. The docking results were saved as "pdb" and binding site composition and average distance between the ligand and amino acid residue was determined, using two online software, 3DLigandSite by Structural Bioinformatics Group, Imperial College London and COACH by Zhang Lab, University of Michigan<sup>[50,51]</sup>.

## RESULTS AND DISCUSSION

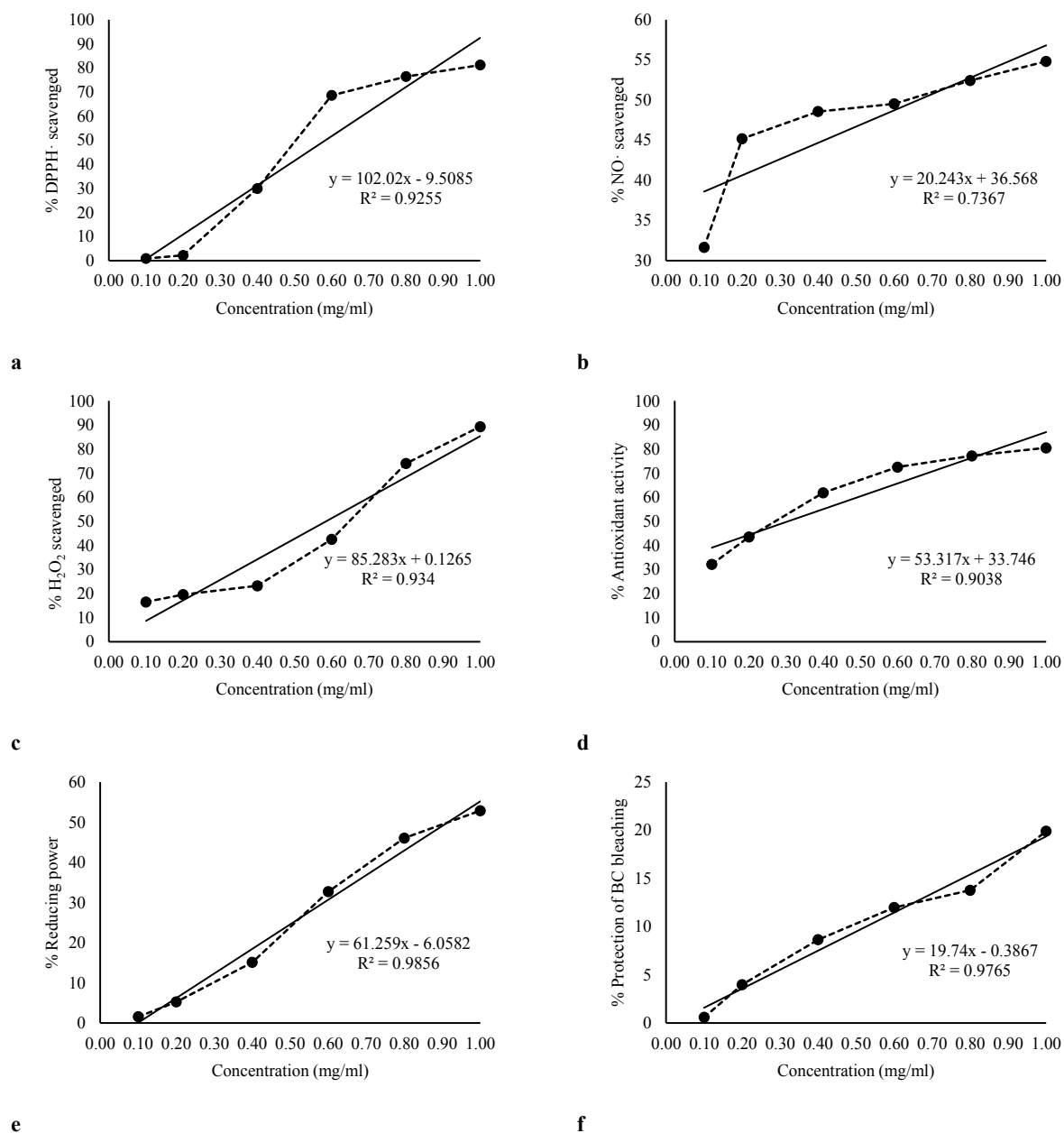
The results of antioxidant activities of OXCPM and the standards (AA and BHT), at similar concentration level (1.00 mg/ml), determined using seven *in vitro* methods are shown in fig. 2. The OXCPM demonstrated promising antioxidant activity in all the employed models in the order as; H<sub>2</sub>O<sub>2</sub> scavenging assay (89.30 $\pm$ 0.013 %) > DPPH· scavenging assay (81.20 $\pm$ 0.002 %) > phosphomolybdenum assay (80.52 $\pm$ 0.016 %) > NO· scavenging assay (54.81 $\pm$ 0.007 %) > reducing power assay (52.87 $\pm$ 0.008 %) > ferric thiocyanate assay (34.44 $\pm$ 0.019 %) > BC bleaching assay (19.91 $\pm$ 0.014 %). It is noteworthy that antioxidant effects of OXCPM were comparable to the standards in the DPPH· and H<sub>2</sub>O<sub>2</sub> scavenging, and BC bleaching assays. However, in all the other models, the activity of the OXCPM was markedly lower than the standards (p<0.05).

The antioxidant activity of the OXCPM in a concentration range (0.10-1.00 mg/ml) indicated a linear rise in the response on increasing the concentration. The dose-dependent antioxidant effects of the OXCPM by various *in vitro* methods and the respective linear



**Fig. 2: Antioxidant activities of equivalent amounts of AA, BHT and OXCPM in *in vitro* assays**

Comparative activities of ascorbic acid (AA), butylated hydroxytoluene (BHT) and 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol (OXCPM) in seven *in vitro* assays. \*Indicates a significant difference in the activity of OXCPM compared to the standards, ■ 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol; ▨ ascorbic acid; ▤ BHT. RPA: Reducing power activity; PA: Phosphomolybdenum activity; H<sub>2</sub>O<sub>2</sub>: H<sub>2</sub>O<sub>2</sub> scavenging activity; FTA: ferric thiocyanate activity; NO: NO radical scavenging activity; BC: BC bleaching activity



**Fig. 3: Concentration-dependent activity of OXCPM**

**a. DPPH· scavenging assay, b. NO· scavenging assay, c. H<sub>2</sub>O<sub>2</sub> scavenging assay, d. reducing power assay, e. phosphomolybdenum assay, f. β-carotene (BC) bleaching assay.** All data points are mean±standard deviation (n=3). OXCPM is 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol

regression equations are shown in figs. 3a-f. The median effective concentration (EC<sub>50</sub>) of the OXCPM, in different *in vitro* antioxidant assays, calculated from the respective linear regression equations are given in Table 1.

The kinetics of OXCPM and the standards in producing the antioxidant effects was evaluated using the DPPH· scavenging, H<sub>2</sub>O<sub>2</sub> scavenging, BC bleaching and ferric thiocyanate assays. In the DPPH· scavenging assay, a sharp transition in the colour of the free radical solution, from purple to yellow, was observed upon

addition of 0.20 ml of AA, whereas, the rate of reaction of OXCPM and BHT with the radical was a bit slower and they took 20 and 45 min, respectively, to reach the steady state absorbance. This colour transition kinetics classified AA, OXCPM and BHT as rapid, intermediate and slow antioxidants, respectively. For the H<sub>2</sub>O<sub>2</sub>, the scavenging effect of OXCPM started within 10 min of incubation and the ROS was completely scavenged by all the studied concentrations after 20 min of incubation. Similar reaction kinetics were observed in the case of AA and BHT.

**TABLE 1: EC<sub>50</sub> VALUES OF OXCPM IN VARIOUS ANTIOXIDANT ASSAYS**

Antioxidant assays	Median effective concentration (EC <sub>50</sub> , mg/ml)
DPPH· scavenging assay	0.58
NO· scavenging assay	0.67
H <sub>2</sub> O <sub>2</sub> scavenging assay	0.59
Phosphomolybdenum assay	0.93
Reducing power assay	0.32
BC bleaching assay	2.56

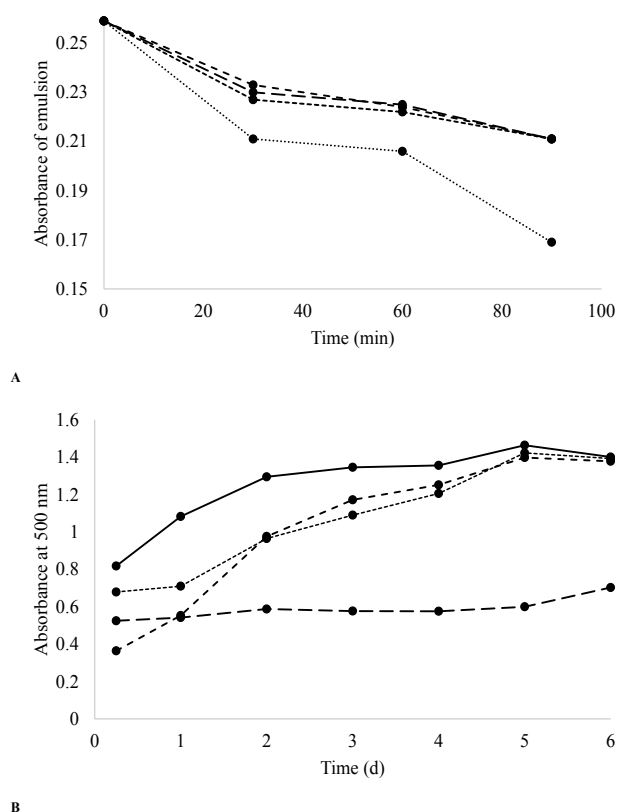
OXCPM is 5-[(4-chlorophenoxy) methyl]-1,3,4-oxadiazole-2-thiol. EC<sub>50</sub> is median effective concentration

The kinetics of the OXCPM and the standards in bleaching BC is shown in fig. 4A. A sharp decline in absorbance of the control was observed after heating for 20 min and continued until end of the study time period. However, the reaction mixtures containing OXCPM effectively inhibited the bleaching of BC emulsion and the absorbance of the emulsion was hence greater than the control. After 90 min incubation, the absorbance values of the emulsions containing OXCPM were not statistically different from those of BHT (0.211±0.014 and 0.211±0.009, respectively) and AA (0.211±0.014 and 0.211±0.010, respectively). The % antioxidant activity calculated using the formula reported by Jayaprakash *et al.* was found to be 41.97<sup>[45]</sup>.

The kinetics of the inhibition of lipid peroxidation by OXCPM and the standards AA and BHT are shown in fig. 4B. The effects of the sample and the standards on lipid peroxidation level were investigated for six consecutive days. Throughout the study period, a gradual increase in the absorbance of the control was observed with maximum at 5<sup>th</sup> d of incubation. At 6<sup>th</sup> d, a drop in the absorbance of the red-colored complex was observed in the control test tubes. The absorbance of the OXCPM at 5<sup>th</sup> and 6<sup>th</sup> d was found markedly lower than the absorbance of control but higher than AA and BHT. The order of absorbance was found to be as control>OXCPM>AA>BHT. The absorbance values for all the concentration range of the sample (0.10-1.00 mg/ml) and AA were also the highest at 5<sup>th</sup> d and decreased afterwards. The decrease in absorbance might be attributed to the formation of malondialdehyde products from oxidation of linoleic acid, however, this was not observed in the case of BHT. The percent inhibition of lipid peroxidation by OXCPM, AA and BHT at 5<sup>th</sup> d was found to be 2.93±0.034, 4.64±0.014 and 59.07±0.016 %, respectively. The statistical analysis indicated that the activity of OXCPM was not significantly different from that of AA. However, at similar concentrations,

the activity of BHT was significantly higher than OXCPM and AA. It was noted that OXCPM and AA showed maximum response (34.44±0.019 and 55.50±0.026 %, respectively) after 24 and 6 h of incubation, respectively, and then a gradual decline in their activity was observed as compared to BHT, which showed maximum inhibition at 5<sup>th</sup> d. The results for both the standards were found to be consistent to the previously published report<sup>[52]</sup>. Hence, it is clear from the result of the study that OXCPM has comparable lipid peroxidation protection than that of AA but the both were less active than BHT.

OXCPM interacted with the protein targets, PTK-2β and GR, at the default binding sites having dimensions such as GR 1gsn (X, 61.0056; Y, -18.9883; Z, 51.4463), GR 1xan (X, 82.1538; Y, -5.7497; Z, 36.095), GR 3grs (X, 60.8014; Y, 51.3499; Z, 18.9135), GR 3grt (X, 63.1107; Y, 37.3065; Z, 20.1816), PTK-2β 3et7 (X, 22.7878; Y, 65.5996; Z, 18.6962), PTK-2β 3fzp (X, 14.893; Y, -10.8816; Z, 7.0870), PTK-2β 3fzs (X, -3.4674; Y,



**Fig. 4: Kinetics of BC bleaching and inhibition of lipid peroxidation by AA, BHT and OXCPM**

**A. β-carotene (BC) bleaching assay and B. inhibition of lipid peroxidation in ferric thiocyanate assay.** All points represent mean±standard deviation of 3 determinations. A. ....●..... control, ---●--- 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol (OXCPM, 1.00 mg/ml), —●— butylated hydroxytoluene (BHT) and -●- ascorbic acid (AA). B —●— control, —●— BHT, -●- AA and .....●..... OXCPM

-3.2746; Z, 12.4054), PTK-2 $\beta$  3fzt (X, -4.9729; Y, -1.8823; Z, 11.6155), PTK-2 $\beta$  3fzr (X, 23.0432; Y, 66.4355; Z, 18.3507) and PTK-2 $\beta$  3h3c (X, 16.7807; Y, -10.0077; Z, 4.3928). The scores of OXCPM binding to these sites indicated that the OXCPM possessed good affinity towards both the targets, PTK-2 $\beta$  and GR, with maximum towards the latter (Table 2). The binding poses of OXCPM relative to the best binding score are shown in fig. 5. These results indicate that OXCPM gives the best conformation with GR 1gsn and GR 3grs.

The composition of binding pockets and average distance between amino acid residue and the ligand was determined for two protein targets for which OXCPM gave best scores and conformations. For GR 3grs, mammoth scores for the ligand cluster included (average InE: 51.2, min LnE: 49.6, max LnE: 51.3), and composition of the binding pocket and average distance between amino acid residue and the ligand was ILE9(0.39), GLY10(0.01), GLY11(0.05), GLY12(0.03), SER13(0.00), GLY14(0.00), GLY15(0.01), VAL32(0.31), GLU33(0.00), SER34(0.00), HIS35(0.07), LYS36(0.34), GLY39(0.00), THR40(0.00), CYS41(0.00), VAL44(0.30), GLY45(0.18), CYS46(0.06), LYS49(0.00), GLY111(0.06), HIS112(0.02), ALA113(0.00), ALA138(0.12), THR139(0.01), GLY140(0.01), GLY141(0.04), SER160(0.04), TYR180(0.00), ILE181(0.58), ARG274(0.24), LEU281(0.32), GLY313(0.09), ASP314(0.00), VAL315(0.36), LEU320(0.03), LEU321(0.00), THR322(0.00), PRO323(0.05) and ALA325(0.57). For GR 1gsn, mammoth scores for the ligand cluster included (average InE: 50.9, min LnE: 49.3, max LnE: 51.0), and composition of the binding pocket and average distance between

amino acid residue and the ligand was ILE9(0.44), GLY10(0.02), GLY11(0.03), GLY12(0.02), SER13(0.00), GLY14(0.00), GLY15(0.02), VAL32(0.39), GLU33(0.00), SER34(0.00), HIS35(0.02), LYS36(0.32), GLY39(0.02), THR40(0.00), CYS41(0.00), VAL44(0.30), GLY45(0.33), LYS49(0.00), GLY111(0.03), HIS112(0.00), ALA113(0.00), ALA138(0.05), THR139(0.00), GLY140(0.02), GLY141(0.02), SER160(0.12), TYR180(0.00), ILE181(0.64), ARG274(0.10), LEU281(0.38), GLY313(0.01), ASP314(0.00), VAL315(0.32), LEU320(0.08), LEU321(0.02), THR322(0.00), ALA325(0.56) and PHE355(0.65)<sup>[50,51]</sup>.

Oxidation is a chemical reaction, which involves the transfer of electrons from one molecule to the other (an oxidizing agent). The importance of oxidation reactions in the body is extensively recognized. All living organisms depend on the oxidative metabolism for their survival and this dependency results in generation of oxygen FRs and ROS. FRs, owing to their instability and high reactivity, undergo oxidative chain reactions, thus producing more radical species. In situations, when the oxidation exceeds the antioxidant defence mechanisms of the body secondary to a loss of balance between them, oxidative stress occurs<sup>[53]</sup>. The prevention against oxidative stress is possible through the use of antioxidants.

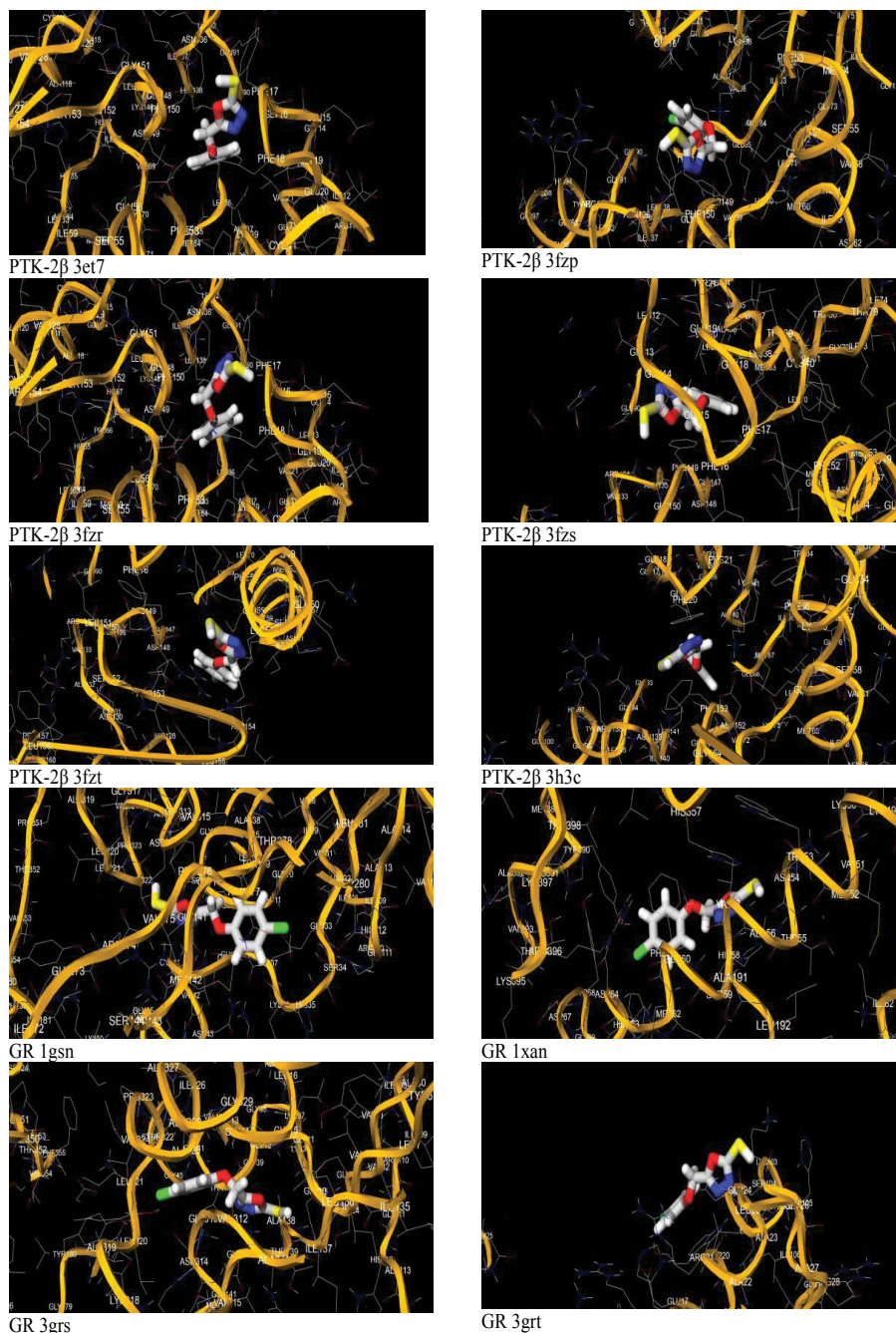
Antioxidants are the substances which slow down, prevent or terminate oxidative chain reactions being oxidized themselves, hence, are chemically reducing agents (thiols, acids or phenols) or conjugated systems. Despite controversies, use of antioxidant-based formulations has increased in the last three decades in the hope of promoting health and preventing chronic diseases<sup>[54]</sup>. However, incorporation of common synthetic antioxidants e.g. butylated hydroxyanisole, BHT, tertiary-butyl hydroquinone and propyl gallate in nutritional supplements, cosmetics and food products has been suspected to be associated with potential health hazards e.g. carcinogenesis, mutagenicity and liver injuries<sup>[55]</sup>. Hence, this situation creates a necessity for scrutinizing safe alternative synthetic antioxidants with broad-spectrum actions.

OXCPM contains a thiol group bonded to the aromatic oxadiazole ring at position 2. The labile hydrogen group attached to sulphur makes the molecule susceptible to oxidation, thereby acting as reducing agent. Keeping in view the structural information, the present study was designed to evaluate the antioxidant potential of OXCPM using seven *in vitro* methods detailed above.

**TABLE 2: BINDING SCORES OF OXCPM, AA AND BHT AGAINST PROTEIN TARGETS**

Protein	PDB ID	Binding score		
		OXCPM	AA	BHT
Protein-tyrosine kinase 2-B	3et7	-5.8	-4.8	-6.7
	3fzp	-6.5	-4.9	-6.8
	3fzr	-6.0	-4.5	-6.3
	3fzs	-5.9	-4.6	-7.3
	3fzt	-6.6	-5.6	-7.2
	3h3c	-6.0	-5.1	-6.8
	1gsn	-6.9	-6.2	-5.5
Glutathione reductase	1xan	-5.0	-4.9	-6.7
	3grs	-7.0	-6.3	-5.7
	3grt	-5.1	-4.7	-5.9

Binding scores against various protein targets were generated using Mcule 1-Click Docking software. OXCPM is 5-[(4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol and AA is ascorbic acid and BHT is butylated hydroxytoluene



**Fig. 5: Best binding fits for OXCPM against oxidative stress-related endogenous targets**

**Protein tyrosine kinase 2-beta (PTK-2 $\beta$ ) and glutathione reductase (GR); carbon, hydrogen, nitrogen, oxygen, chlorine and sulphur are represented by silver, white, blue, red, green and yellow colors, respectively. 5-[4-chlorophenoxy)methyl]-1,3,4-oxadiazole-2-thiol (OXCPM)**

Preliminary findings of the antiradical activity of OXCPM by the DPPH assay indicated that OXCPM possessed excellent free radical scavenging potential. Later on, OXCPM was investigated for the potential to quench specific radicals involved in oxidative stress such as OH $\cdot$ , NO $\cdot$  and LOO $\cdot$ . OXCPM demonstrated good antioxidant activity against oxidative stress-related OFR and ROS by all adopted models, however, a non-significant difference was observed between the response of OXCPM and AA in preventing lipid

peroxidation (investigated using BC bleaching and ferric thiocyanate assay) and scavenging H<sub>2</sub>O<sub>2</sub>. The reaction kinetics by various models suggested that OXCPM was intermediate-fast antioxidant.

Though, the methods employed for evaluation of antioxidant activities gave a strong prediction of OXCPM's potential. However, the results from chemical methods cannot entirely be extrapolated to the living organisms. Several literature reports reveal that an antioxidant may not only exert its effect directly



by quenching the FRs but also by inducing endogenous defence mechanisms<sup>[56,57]</sup>. Hence, molecular docking studies were carried out to further support the findings and evaluate molecular level interaction of OXCPM with endogenous oxidative-stress-related targets.

The ligand-protein interactions were predicted using Mcule's 1-Click Docking software, which used AutoDock Vina (with default parameters) for docking calculations, which could facilitate the drug discovery program by providing useful information regarding binding score and orientation of a particular ligand to the selected target. The software stores best-four docking poses. An important advantage of using the software was that the identity of the molecule remains un-modified during docking i.e. InChI strings of the input ligand and output conformer are compared and in case of InChI mismatch, the results are discarded.

For docking studies, PTK-2 $\beta$  was selected since it is a custodian of cell regulatory pathways and signal transduction processes and during OS, functional changes in the protein lead to destructive events<sup>[53]</sup>. GR is another endogenous defence component, which copes the OS by catalysing the reduction of glutathione disulphide to the sulfhydryl glutathione<sup>[58]</sup>. The docking results obtained indicated that BHT exhibited highest binding scores for all the subclasses of PTK-2 $\beta$  followed by OXCPM and AA. Among the subclasses of PTK-2 $\beta$ , BHT showed maximum affinity towards PTK-2 $\beta$  3fzs and PTK-2 $\beta$  3fzt, OXCPM towards PTK-2 $\beta$  3fzt and PTK-2 $\beta$  3fzp, and AA towards PTK-2 $\beta$  3fzt. For glutathione reductase, maximum binding scores were obtained for OXCPM in all the subclasses except GR 1xan where BHT showed maximum affinity. OXCPM demonstrated high binding affinity for GR 3grs and 1gsn, BHT for GR 1xan, and AsA for GR 3grs and GR 1gsn. It is important to mention here that OXCPM and AA showed maximum binding affinities towards the same subclass of GR, but OXCPM had shown to better attach to the active site rather than AA. Hence, these results confirm the *in vitro* findings and indicate that the OXCPM is a putative antioxidant molecule.

The results of the present study indicated that OXCPM was a potential antioxidant that produced pleiotropic effects by scavenging a variety of OFR and ROS. Moreover, its strong binding affinities for PTK-2 $\beta$  and GR revealed that it possessed the ability to induce the endogenous defence system. Hence, it has the potential to be developed into an effective antioxidant drug. Nonetheless, more data revealing its safety and *in vivo*

antioxidant potential is needed.

### Acknowledgements:

One of the authors, NS thanks the authorities of the University of the Punjab, Lahore, Pakistan, for providing chemicals/solvents and laboratory facilities to conduct the study. Authors thank Dr S. Z. Siddiqui, Department of Chemistry, Government College University, Lahore, Pakistan for providing a gift sample of OXCPM.

### Conflict of interest:

There is no conflict of interest, financial or otherwise associated with this project.

### Financial support and sponsorship:

This research is a part of a project funded by the Higher Education Commission, Pakistan, under National Research Program for Universities, vide letter No.6777/Punjab/NRPU/R&D/HEC/2016, dated 16-05-2017.

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