Evaluation of novel pyrimidine derivatives as a new class of mushroom tyrosinase inhibitor

S Shohreh Mirmortazavi 1
Mahdieh Farvandi 1
Hossein Ghafouri 1
Asadollah Mohammadi 2
Mostafa Shourian 1

1Department of Biology, Faculty of Sciences, University of Guilan, Rasht, Iran; 2Department of Chemistry, Faculty of Sciences, University of Guilan, Rasht, Iran

Background and aim: Tyrosinase (EC 1.14.18.1) is responsible for enzymatic browning in fruits and vegetables. Its inhibitors may be applied to efficiently treat hyperpigmentation and are widely used in pharmaceutical and cosmetic products, food supplements and insecticides. Previous studies have shown that heterocyclic compounds with an amino group can inhibit tyrosinase activity. The present study aims to evaluate the inhibitory effect of some novel 2,6-diamino-4-chloropyrimidine derivatives (1a-e) and 2,4,6-triaminopyrimidine (2a–e) including bioactive aniline moiety on the activity of the mushroom tyrosinase.

Methods: In practice, the azo salt was initially synthesized from aniline derivatives and combined subsequently with the 2,4,6-triaminopyrimidine and 2,6-diamino-4-chloropyrimidine followed by crystallization. The structures of resulting compounds were confirmed by FT-IR, 13C NMR, and 1H NMR. The derivatives (0–100 µM) were evaluated for their inhibitory effect on tyrosinase activity using L-3,4-dihydroxyphenylalanine (L-DOPA) as substrate.

Results: All compounds showed inhibitory effects against the activity of the enzyme. About 23.72–55.08% inhibition was observed in the presence of 30 µM of each compound. The IC50 values of the synthesized compounds were measured, and their inhibition properties were also visualized by zymography. Based on the results, the compounds 1a-e and 2a-e showed moderate inhibitory activities. Notably, pyrimidine derivatives 1a (IC50=24.68) and 1d (IC50=24.45) also exhibited similar inhibitory activities when compared with the positive control, kojic acid (IC50=25.24 µM). Kinetic studies indicated that the type of inhibition was noncompetitive.

Conclusion: All results suggest that pyrimidine derivatives, especially 1d and 1a, can be considered as safe and efficient tyrosinase inhibitors.

Keywords: mushroom tyrosinase, inhibitor, pyrimidine derivatives, noncompetitive inhibition

Introduction

Tyrosinase is a copper-dependent enzyme from the oxidoreductase family considered as the rate-limiting enzyme in the melanin production pathway. Melanin is a dark brown to black pigment that effectively absorbs light. It is found in plants and fungi as well as in hair, skin, and iris of the eye in human and animals.1,2 Tyrosinase is one of the multifunctional enzymes that catalyze two different reactions resulting from the binding of dioxygen to two copper atoms in the strongly conserved active center: the hydroxylation of monophenols (tyrosine) to o-diphenols (monophenolase activity) and the oxidation of o-diphenols to o-quinones (diphenolase activity). The process ultimately leads to the production of melanin through several reactions from o-quinones.3 This type of...
Tyrosinase activity is due to four possible oxidation states (deoxy-, oxy-, met- and deact-tyrosinase). At the met state, both copper atoms present in the active site are in Cu (II) state. In the case of the deoxy state, copper atoms in Cu (I) can be attached to molecular oxygen. At the oxy state, the molecular oxygen in peroxo conformation is surrounded by both copper atoms in a Cu (II) state, in which case it can be linked to phenols and catechols, as substrate, converting them into ortho-quinones by different oxidative cycles. The deact state is an inactivated form of tyrosinase with one of the copper atoms in the Cu (0) state and the other in the Cu (II) state.  

Given this background, while pigmentation under normal conditions protects skin against harmful UV injury, melanin increment due to the tyrosinase hyperactivity can give rise to hyperpigmentation disorders, which is followed by freckles, wrinkles, age spots, skin irritation, postinflammatory hyperpigmentation, melano derma and melasma in extreme cases that are particularly common among women.  

Tyrosinase is regarded as a key agent in the risk of malignant melanoma cancer and might play a significant role in neuromelanin production responsible for the neurodegeneration associated with Parkinson’s and Huntington’s diseases.  

Tyrosinase has also been confirmed to be one of the most important causes of browning in vegetables and fruits during post-harvest handling, and storage, leading to quick degradation.  

Finally, tyrosinase has a different biochemical process in insects, including wound healing, sclerotization, melanin synthesis, and defensive encapsulation. Therefore, the production of useful and safe inhibitors of tyrosinase as insecticides and skin lightening agents is important for the agricultural, cosmetic and medical industries. While there have been many efforts to discover effective inhibitors, only a few compounds have been used commercially for therapeutic and cosmetic uses; these include kojic acid, tropolone, arbutin and 1-phenyl-2-thiourea (PTU).  

Tyrosinase can oxidize several aromatic amines and o-aminophenols as substrates, which are analogs of monophenols and o-diphenols, respectively.  

4-Aminophenylalanine, 4-aminotoluene, 4-aminoacetanilide, and several o-aminophenolic derivatives of benzoic acid served as substrates for the enzyme.  

Because of the widespread presence of a pyrimidine ring in biological and organic compounds such as nucleic bases, vitamins, enzymes, chlorophyll, hemoglobin, and hormones, and, moreover, the broad spectrum of pharmaceutical properties subsequently has attracted much attention in drug industries. Pyrimidine and its derivatives with an active heterocyclic structure having two nitrogen atoms in the ring are highly regarded as interesting bioactive compounds in medicinal chemistry. These heterocyclic compounds have anticancer, antiviral, anti-inflammatory, antibacterial, and anti-parkinson properties.  

Considering that several aromatic amines can play a role as tyrosinase substrates, and the fact that amines and anilines are its inhibitors, we decided to synthesize new derivatives based on pyrimidine and aniline derivatives. In this study, for the first time, we synthesized a series of novel azo-pyrimidine dyes and evaluated the inhibitory activities of these low-cost and easy-to-prepare compounds against mushroom tyrosinase based on dopaquinone production by an enzyme assay with L-dopa as substrate. The IC50 was obtained and compared with kojic acid as positive control. Also, zymography was used to show the inhibitory effects of these compounds on tyrosinase activity. The kinetic analysis revealed the Ki value and inhibition mechanism, and finally, the docking study of these compounds was performed by the auto dock software and Swiss dock web server.

**Materials and equipment**  

All chemicals and reagents used in this study were purchased from Sigma-Aldrich Co., St Louis, MO, USA and Merck Chemical companies and used without further purification. Silica gel thin layer chromatography (TLC) using n-hexane and ethyl acetate as mobile phase was used to monitor the progress of reactions. The solutions were prepared using distilled water. Melting points (°C) of the synthesized compounds were recorded on Electrothermal IA9000 equipment (Cole-Parmer, Stone, UK) and are uncorrected. The UV-visible absorption spectra were recorded using a Pharmacia Biotech Spectrophotometer in the range of 200–800 nm. Fourier transform infrared (FT-IR) spectra were recorded on a Shimadzu 8400 FT-IR spectrophotometer (Shimadzu, Kyoto, Japan).  

1H NMR and 13C NMR spectra were obtained on an FT-NMR (400 MHz) Bruker apparatus (Bruker AXS Inc., Madison, WI, USA), the chemical shifts were expressed in δ ppm using TMS as an internal standard and J values in Hz.

**Synthesis of pyrimidine-based azo dyes (1a-e and 2a-e)**  

Pyrimidine-based azo dyes (1a-e and 2a-e) were synthesized in high yield via diazotization and coupling reactions as shown in Figure 1. One mmol of diazonium salts from donor- and acceptor-substituted anilines was prepared according to a method previously described. To 1 mmol of solution containing coupling reagent (2,4,6-triaminopyrimidine or 2,6-diamino-4-chloropyrimidine) was added dropwise from diazonium salt from
the previous step over 30 min with stirring at 0–5°C. The stoichiometry ratio was 1:1. The pH of the solution was slowly adjusted to 7 with HCl (0.5 M). The reaction was continued by addition of 20 ml distilled water to precipitate. The precipitate was then isolated by recrystallization from dimethylformamide (DMF)/H2O at 80°C. Structures of the obtained azo dyes were confirmed by FT-IR and NMR (1H and 13C) spectroscopy.

**Compound 1a (6-chloro-5-((2-nitrophenyl) diazenyl) pyrimidine-2,4-diamine)**

Red solid; yield 79%; melting point (MP) 301–303°C. FT-IR (KBr, cm−1): 3,467 (NH), 3,308 (NH), 3,138 (NH), 1,632 (C=O), 1,658 (C=C), 1,598 (C=N), 1,502 (N=N). 1H NMR (400 MHz, DMSO-d6, 298 K), δ (ppm): 7.58–7.62 (m, 2H), 7.76 (s, 1H, NH), 7.79–7.85 (m, 2H), 8.01 (s, 1H, NH), 9.27 (s, 2H, NH), 10.19 (s, 1H, NH). 13C NMR (DMSO-d6), δ (ppm): 118.1, 120.4, 125.1, 129.7, 134.2, 144.7, 146.1, 156.2, 161.9, 166.1.

**Compound 1b (6-chloro-5-((2-nitrophenyl) diazenyl) pyrimidine-2,4-diamine)**

Orange solid; yield 82%; MP 305–307°C. FT-IR (KBr, cm−1): 3,502 (NH), 3,327 (NH), 3,186 (NH), 1,632 (C=O), 1,509 (C=N). 1H NMR (400 MHz, DMSO-d6, 298 K), δ (ppm): 7.55 (s, 1H, NH), 7.73 (s, 1H, NH), 7.90 (d, 2H, J = 8.8 Hz, Ar-H), 8.35 (d, 2H, J = 7.6 Hz, Ar-H), 8.39 (s, 1H, NH), 9.41 (s, 1H, NH). 13C NMR (DMSO-d6), δ (ppm): 120.1, 122.5, 125.4, 147.0, 156.4, 161.8, 162.8, 166.2.

**Compound 1c (6-chloro-5-(p-tolyldiazenyl) pyrimidine-2,4-diamine)**

Light brown solid; yield 84%; MP 234–236°C. FT-IR (KBr, cm−1): 3,385 (NH), 3,312 (NH), 3,205 (NH), 3,028 (=CH), 2,919 (CH), 1,643 (C=C), 1,601 (C=N), 1,551 (N=N). 1H NMR (400 MHz, DMSO-d6, 298 K), δ (ppm): 2.38 (s, 3H, CH3), 7.29 (s, 1H, NH), 7.33 (d, 2H, J = 8.0 Hz, Ar-H), 7.51 (s, 1H, NH), 7.71 (d, 2H, J = 8.4 Hz, Ar-H), 8.08 (s, 1H, NH), 9.28 (s, 1H, NH). 13C NMR (DMSO-d6), δ (ppm): 20.4, 118.4, 121.9, 130.3, 139.7, 150.8, 156.5, 161.5, 164.6.

**Compound 1d (N-(4-((2, 4-diamino-6-chloropyrimidin-5-yl) diazenyl) phenyl) acetamide)**

Dark red solid; yield 74%; MP 286–288°C. FT-IR (KBr, cm−1): 3,425 (NH), 3,390 (NH), 3,127 (NH), 1,691 (C=O), 1,658 (C=C), 1,598 (C=N), 1,502 (N=N). 1H NMR (400 MHz, DMSO-d6, 298 K), δ (ppm): 2.09 (s, 3H, CH3), 7.24 (s, 1H, NH), 7.34 (s, 1H, NH), 7.75 (m, 4H, Ar-H), 8.05 (s, 1H, NH), 9.27 (s, 2H, NH), 10.19 (s, 1H, NH). 13C NMR (DMSO-d6), δ (ppm): 24.5, 118.9, 122.7, 141.0, 148.3, 150.8, 156.5, 161.4, 162.8, 164.3, 169.0.

**Compound 1e (6-chloro-5-((4-methoxyphenyl) diazenyl) pyrimidine-2,4-diamine)**

Dark red solid; yield 84%; MP 236–238°C. FT-IR (KBr, cm−1): 3,366 (NH), 3,313 (NH), 3,201 (NH), 3,002 (=CH), 2,907 (CH), 1,648 (C=C), 1,602 (C=N), 1,550 (N=N). 1H NMR (400 MHz, DMSO-d6, 298 K), δ (ppm): 3.35 (s, 3H, CH3), 7.0 (s, 1H, NH), 7.08 (d, 2H, J = 8.8 Hz, Ar-H), 7.25 (s, 1H, NH), 7.79 (d, 2H, J = 8.8 Hz, Ar-H), 8.01 (s, 1H, NH), 9.23 (s, 1H, NH). 13C NMR
Compound 2a N-(4-((2,4,6-triaminopyrimidin-5-yl) diazenyl) phenyl) acetamide

Red solid; yield 75%; MP 284–286°C. FT-IR (KBr, cm⁻¹): 3,389 (NH), 3,174 (NH), 2,960 (=CH), 2,929 (CH), 1,650 (C=O), 1,605 (C=N), 1,501 (N=N). ¹H NMR (400 MHz, DMSO-d₆, 298 K), δ (ppm): 2.09 (s, 3H, CH₃), 7.06 (s, 2H, NH), 7.61 (s, 2H, NH), 7.67 (d, 2H, J=8.0 Hz, Ar-H), 7.85 (d, 2H, J=8.0 Hz, Ar-H), 8.66 (s, 2H, NH). ¹³C NMR (DMSO-d₆), δ (ppm): 24.5, 108.2, 119.6, 122.1, 122.4, 139.9, 148.5, 168.9.

Compound 2b (5-p-tolyldiazenyl) pyrimidine-2,4,6-triamine

Orange solid; yield 73%; MP 307–309°C. FT-IR (KBr, cm⁻¹): 3,428 (NH), 3,316 (NH), 3,110 (NO₂), 1,508 (N=N). ¹H NMR (400 MHz, DMSO-d₆, 298 K), δ (ppm): 2.35 (s, 3H, CH₃), 6.45 (s, 2H, NH), 7.14 (s, 2H, NH), 7.23 (d, 2H, J=7.6 Hz, Ar-H), 7.70 (d, 2H, J=6.8 Hz, Ar-H), 7.97 (s, 2H, NH), 119.6, 122.1, 124.4, 137.2, 151.3, 162.9, 162.9, 163.

Compound 2c (5-((4-nitrophenyl) diazenyl) pyrimidine-2,4,6-triamine)

Orange solid; yield 83%; MP 250–252°C. FT-IR (KBr, cm⁻¹): 3,428 (NH), 3,338 (NH), 3,150 (NH), 2,958 (=CH), 2,925 (CH), 1,660 (C=O), 1,599 (C=C), 1,501 (N=N). ¹H NMR (400 MHz, DMSO-d₆, 298 K), δ (ppm): 2.93 (s, 3H, CH₃), 7.01 (d, 2H, J=9.2 Hz, Ar-H), 7.46 (t, 1H, J=7.6 Hz, Ar-H), 7.95 (s, 2H, NH), 7.97 (d, 2H, J=8.4 Hz, Ar-H), 9.05 (s, 2H, NH), 123.8, 124.7, 127.6, 133.8, 145.1, 145.7.

Compound 2d (5-((4-methoxyphenyl) diazenyl) pyrimidine-2,4,6-triamine)

Yellow solid; yield 75%; MP 328–330°C. FT-IR (KBr, cm⁻¹): 3,463 (NH), 3,379 (NH), 3,110 (NH), 1,513 (N=N). ¹H NMR (400 MHz, DMSO-d₆, 298 K), δ (ppm): 7.27 (s, 2H, NH), 7.46 (t, 1H, J=7.6, Ar-H), 7.71 (t, 1H, J=7.6 Hz, Ar-H), 7.95 (s, 2H, NH), 7.97 (d, 1H, J=8.0 Hz, Ar-H), 8.28 (d, 1H, J=8.4 Hz, Ar-H), 9.39 (s, 2H, NH). ¹³C NMR (DMSO-d₆), δ (ppm): 111.3, 118.4, 121.3, 123.8, 124.7, 127.6, 133.8, 145.1, 145.7.

Mushroom tyrosinase inhibition assay (IC₅₀)

Tyrosinase was assayed in the presence of synthesized products using a previously described method with some modification. Briefly, reaction mixtures (total volume of 100 μL) with 70 μL of phosphate buffer (pH 6.8, 100 mM) and 0, 10, 30, 60, 100 μM synthesized compounds dissolved in DMSO (final concentration 7% v/v) were first prepared. This was followed by the addition of 3 μL mushroom tyrosinase (0.0003 mg/mL), mixing well and pre-incubation for 5 min at 37°C. The final concentration of DMSO (7% v/v) did not interfere with the enzyme activity. Twenty μL of L-DOPA (10 mM) were then added as the substrate and incubated for 30 min at 37°C. Subsequently, the mixed solution was placed in a 96-well plate, and the absorbance of dopachrome was measured at 490 nm by ELISA reader. The half maximal inhibitory concentration of each tested compound (IC₅₀) was determined by interpolation of the dose–response curves. All the experiments were carried out in triplicate for each compound, and concentration and values were expressed as the means of three experiments before further calculation. Kojic acid and phosphate buffer were used as a reference inhibitor and negative control, respectively. The percent inhibition of the enzyme reaction was calculated as follows:

\[
\text{Inhibition rate (\%)} = \left(\frac{B - S}{B}\right) \times 100
\]

where B and S are the absorbance values for the blank and sample, respectively.

Determination of the inhibition type and the inhibition constants

In order to obtain kinetic parameters, the activity of tyrosinase was measured at different concentrations of substrate (0, 0.1, 0.3, 0.75, 1.25 and 2 mM) in the absence and presence of 30 μM concentration of inhibitors. The protocol was the same as discussed in mushroom tyrosinase inhibition assay. Using the value of enzyme activity obtained from the assay in different concentrations of the substrate and inhibitor, Michaelis–Menten and Lineweaver–Burk
plots were obtained. Km and Vmax were calculated from the Lineweaver–Burk plot linear equation and the inhibition constants were determined by the secondary plots of 1/Vmax versus the concentration of the inhibitor.

**Zymogram analysis**
The effects of synthetic compounds on tyrosinase activity were also visualized by native polyacrylamide gel electrophoresis (PAGE). Synthesized compounds with the enzyme and the appropriate amount of phosphate buffer (pH 6.8, 100 mM) were pre-incubated for 10 min and resolved in 12% PAGE. Preparation of samples was carried out with a sample buffer without mercaptoethanol without heat, and the electrophoresis was performed at 4°C. After electrophoresis, the gel was washed twice with phosphate buffer at room temperature. An appropriate amount of L-DOPA (10 mM) was later added in fresh buffer. The brown color bands of the product appeared after enzymatic activity with gentle shaking at 37°C for 1 h.

**Docking**
The interaction of compound 1d as a ligand with tyrosinase was evaluated by molecular docking. The PDB code of the enzyme from the RCSB Protein Data Bank (http://www.rcsb.org) was obtained as 2Y9W. The REDUCE program were fixed missing hydrogen of the tyrosinase crystal structure. The blind molecular docking was performed by the AutoDock program. The necessary input files were prepared by MGLTools, and the exhaustiveness parameter was set to 1,000. The structure of the ligand was built and optimized by Hyper-Chem program version 8 using a semi-empirical method (PM3 force filed). The images were created using Chimera 1.13.1 based on binding modes and scored using their FullFitness and clustered. Also, the molecular interactions between 1d and tyrosinase was predicted by SwissDock web service (http://www.swissdock.ch).

The images were created using Python Molecule Viewer (PMV) and the program LigPlot v.1.0, which generate schematic two-dimensional representations of protein–ligand complexes from the input PDB file. For obtaining λ_max of all compounds was between 200 and 800 nm. Chemical properties of synthetic compounds are presented in Table 1.

**The inhibitory effects of synthetic compounds on the enzyme activity**
In this study, all compounds were evaluated for their inhibitory effect on tyrosinase activity according to the procedure mentioned in the materials and methods section, and kojic acid was used as the reference inhibitor. The results indicated that percent inhibition of all compounds at 30 μM concentration ranged from 23.72% (for 1b) to 55.08% (for 1e) as compared to kojic acid as a positive control with 57.27% inhibition at 30 μM (Figure 3). The half maximal inhibitory concentration (IC_{50}) of synthetic compounds on tyrosinase activity was determined from logarthmic concentration–inhibition curves (Figure 4). As shown in Table 2, compounds 1d and 1a revealed the most potent inhibitory effects on the mushroom tyrosinase activity with IC_{50} values of 24.45 and 24.69 μM, respectively. Therefore, it was found that our new compounds are more potent inhibitors of tyrosinase than the reference inhibitor, kojic acid (IC_{50}=25.24 μM). In addition, based on the results, it was found that compound 2d (IC_{50}=43.80 μM) exhibited the lowest inhibitory activity on mushroom tyrosinase among all synthetic compounds.

**Tyrosinase zymography**
The effects of synthetic compounds on tyrosinase activity were visualized by L-DOPA staining assay. Tyrosinase oxidation activity on the gel can be observed as a dark band to a lighter band after adding L-DOPA solution. Upon incubation with the synthetic compounds, the band intensity decreased for each compound as compared to the untreated control. This phenomenon suggests that all the compounds had an inhibition effect on tyrosinase. As shown in Figure 5, while the lightest bands corresponded to 1d and 1a compounds, compound 2d showed minimal inhibitory effect compared to the other compounds. These results confirm our findings from inhibitory activity measured by the spectrophotometric method.

**The type of tyrosinase inhibition by synthetic compounds**
The steady-state kinetic analysis of all compounds for inhibition of enzyme activity on different concentrations of L-DOPA (0.1 to 2 mM) was carried out to determine their inhibition type and inhibition constants (Table 2). Lines intersect on...
Lineweaver–Burk plots showed that all synthetic compounds were a noncompetitive inhibitor of the mushroom tyrosinase (Figure 6).

Docking
The binding of (1d) to tyrosinase (2Y9W) was computerized by molecular docking. The results of docking were well clustered around the conformer with the best score. According to Figure 7, the results of docking showed that hydrophobic interactions and polar–polar interaction between N3 of the ligand and carbonyl group of Lys5 play a major role in the binding of 1d to tyrosinase. Based on the results of the modeling, the dominant interaction is hydrophobic.

Discussion
Tyrosinase is a rate-limiting monooxygenase enzyme in the process of melanin production. Increasing the activity of this enzyme causes excessive production of melanin and, as a result, hyperpigmentation problems such as melasma and skin spots may occur. Therefore, the discovery

Table 1 Structures and properties of synthetic compounds

| No. | $\lambda_{\text{max}}$ (nm) | MP °C | MW (g/mol) | Molecular formula | No. | $\lambda_{\text{max}}$ (nm) | MP °C | MW (g/mol) | Molecular formula |
|-----|------------------|-------|-----------|-----------------|-----|------------------|-------|-----------|-----------------|
| 1a  | 402              | 301–303 | 293.67    | $\text{C}_{10}\text{H}_8\text{ClN}_2\text{O}_2$ | 2a  | 394              | 284–286 | 286.29    | $\text{C}_{12}\text{H}_7\text{N}_6\text{O}$ |
| 1b  | 423              | 305–307 | 293.67    | $\text{C}_{10}\text{H}_8\text{ClN}_2\text{O}_2$ | 2b  | 423              | 250–252 | 243.27    | $\text{C}_{11}\text{H}_7\text{N}_7$ |
| 1c  | 456              | 234–236 | 262.70    | $\text{C}_{11}\text{H}_7\text{ClN}_6$        | 2c  | 446              | 307–309 | 274.24    | $\text{C}_{10}\text{H}_8\text{N}_2\text{O}_2$ |
| 1d  | 402              | 286–288 | 305.7     | $\text{C}_{12}\text{H}_8\text{ClN}_2\text{O}$ | 2d  | 384              | 278–280 | 259.27    | $\text{C}_{11}\text{H}_7\text{N}_6$ |
| 1e  | 389              | 270–272 | 287.70    | $\text{C}_{12}\text{H}_8\text{ClN}_2\text{O}$ | 2e  | 402              | 322–324 | 274.24    | $\text{C}_{10}\text{H}_8\text{N}_2\text{O}$ |

Abbreviations: MP, melting point; MW, molecular weight.

Figure 2 $^1$HNMR spectrum of compound 1a in DMSO.

Figure 3 Percent inhibition curve for oxidation of L-DOPA catalyzed by mushroom tyrosinase at increasing concentrations of synthetic pyrimidine derivatives. Inhibitory activity of all synthetic compounds at 30 µM is 23.72–55.08%.
Figure 4 The IC$_{50}$ (µM) values of synthetic compounds against mushroom tyrosinase. L-DOPA was used as substrate and concentrations of synthetic compounds were 10, 30, 60, 100 µM. Absorbance of the reaction was taken at 490 nm. (IC$_{50}$: 2d>1c>2b>2c>1b>2e>1e>kojic acid>1a>1d).

Table 2 The kinetic constants and inhibition constants of all synthetic compounds on mushroom tyrosinase

| Compounds | Vmax  | Km   | αKi  | Ki    | Alpha(α) | IC$_{50}$(µM) | Inhibition type |
|-----------|-------|------|------|-------|-----------|---------------|----------------|
| Control   | 125.58| 0.41 | 6.079| 3.601 | 1.68      | 24.68         | Noncompetitive |
| 1a        | 21.15 | 0.63 | 7.11 | 6.559 | 1.08      | 37.191        | Noncompetitive |
| 1b        | 24.06 | 0.43 | 7.378| 5.871 | 1.25      | 42.04         | Noncompetitive |
| 1c        | 24.78 | 0.48 | 7.464| 7.837 | 0.95      | 24.4          | Noncompetitive |
| 1d        | 25.01 | 0.39 | 7.464| 7.837 | 0.95      | 28.01         | Noncompetitive |
| 1e        | 23.36 | 0.39 | 6.857| 7.05  | 0.97      | 29.77         | Noncompetitive |
| 2a        | 21.88 | 0.496| 6.33 | 4.98  | 1.27      | 41.4          | Noncompetitive |
| 2b        | 19.64 | 0.62 | 5.36 | 3.4   | 1.63      | 41.4          | Noncompetitive |
| 2c        | 23.95 | 0.57 | 7.07 | 4.62  | 1.52      | 40.62         | Noncompetitive |
| 2d        | 22.41 | 0.35 | 6.51 | 7.63  | 0.85      | 43.8          | Noncompetitive |
| 2e        | 23.1  | 0.46 | 6.76 | 5.66  | 1.19      | 34.06         | Noncompetitive |

Figure 5 The effect of the synthetized compounds (1a-e and 2a-e) on the enzyme activity estimated by zymography using L-DOPA staining (A) based on 2,6-diamino-4-chloropyrimidine and (B) based on 2,4,6-triaminopyrimidine. (A: 1d>1a>1e>1b>1c) (B: 2a>2e>2c>2b>2d).
and synthesis of safe and effective inhibitors for treating and preventing skin disorders are very important in the medicinal and cosmetic industries.2,27 On the other hand, this enzyme plays an important role in physiological processes in insects, including wound healing, sclerotization, and defensive encapsulation. Thus, its inhibitors could be useful as insecticides in the agricultural research activities.10,11 Tyrosinase is also responsible for the undesirable browning of fruits and vegetables during storage, which reduces the nutritional and commercial value of the products.8,9 Synthesis and detection of effective, harmless and affordable inhibitors is desirable in order to minimize these problems. Over recent years, there have been many studies of screening, discovery, design, and synthesis of novel tyrosinase inhibitors. Pyrimidine derivatives are active heterocyclic compounds with two nitrogen atoms in the ring and have attracted much attention in medicinal chemistry and pharmacology.15,16 They have been shown to have anticancer, antiviral, anti-inflammatory, antibacterial and antiparkinson properties.15 Considering the reports on the tyrosinase inhibition ability of aromatic amines and anilines, we were encouraged to synthesize some new derivatives based on pyrimidine and aniline derivatives.17 In this study, we synthesized and characterized ten novel azo-dyes including bioactive pyrimidine moiety and aniline derivatives. Their inhibitory activities against mushroom tyrosinase were then evaluated. Tyrosinase inhibition assay was carried out with L-DOPA as substrate and kojic acid as reference inhibitor. All synthetic compounds (1a-e and 2a-e) were able to inhibit mushroom tyrosinase activity in a concentration-dependent manner, as shown in Figure 3. The data revealed that compounds with
a Cl-halide group in the R position have more inhibitory activity than the compounds with an NH₂ group in the R position. The 1d (IC₅₀=24.45 µM) and 1a (IC₅₀=24.68 µM) compounds, possessing acetamide and NO₂ groups at position X, respectively, and with Cl at position R, are more potent inhibitors of mushroom tyrosinase than the other synthesized compounds and IC₅₀ values are slightly better than kojic acid (IC₅₀=25.24 µM), the reference tyrosinase inhibitor. The mechanisms of inhibition of mushroom tyrosinase by all synthetic compounds were further investigated by steady-state kinetic analysis. Inhibition data were then determined using Lineweaver–Burk double reciprocal plots. The outcome of compounds showed that the 1/V vs 1/[S] plot gave two straight lines with different slopes, but with one intersection on the horizontal axis, indicating that the values of Vmax decrease without changing the values of Km when the concentration of these compounds increased. These data suggest that of all synthetic compounds function as non-competitive inhibitors of tyrosinase (Table 2), hence all synthetic compounds bound both the free enzyme molecule and enzyme-substrate complex. Liu et al. synthesized several compounds on the basis of thiourea containing NH group in the ring and evaluated their inhibitory potencies on mushroom tyrosinase. According to their kinetic results, the thiouracil and methylthiouracil synthesized compounds with IC₅₀=128 and 142 µM, respectively, assigned as noncompetitive inhibitors. The SAR studies have shown that inhibitory activity of thiourea on tyrosinase enzyme activity must be derived from nitrogen and sulfur atoms. For these two compounds, it was observed that adding a methyl group in the meta position on the ring could reduce the inhibitory effect of the methylthiouracil compound. According to these results, in the present study, the effect of an NH group in the pyrimidine ring of synthesized compounds was investigated on the potency of inhibition. By comparing the results of the present study with Liu et al it was found that the presence of a methyl group decreases the inhibitory effect of synthetic compounds (2b and 1c). Also, Bae et al reported a series of azo-resveratrol and azoxyresveratrol compounds, containing an azo group in their structure. Among the synthesized compounds, 13b (R₁ = R₂ = OH, R₄ = OH, R₃ = R₅ = H; = R₆ = OMe) exhibited high tyrosinase inhibitory activity, with an IC₅₀ value of 36.28 µM. Therefore, in the present study, all compounds synthesized from base compounds TAP and DATP with 2-nitroaniline, paranitroaniline, paramethoxyaniline and para-aminoacetanilide containing the azo group. Considering the effect of ortho and para position on the benzene ring of synthetic compounds, it was found that 2e (IC₅₀=34.06) with a nitro group at position ortho had a more potent inhibitory effect on tyrosinase activity than 2c (IC₅₀=40.62) with a nitro group at the para position. Bae et al synthesized methoxyaniline derivatives and evaluated as a tyrosinase inhibitor. The theoretical studies conducted in this study identified the interaction of synthetic compounds with the key amino acids of tyrosinase (PDB =2Y9W).

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Author contributions

All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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