

Northumbria Research Link

Citation: Zolghadri, Samaneh, Bahrami, Asieh, Hassan Khan, Mahmud Tareq, Munoz, Jose, Garcia-Molina, Francisco, Garcia-Canovas, F. and Saboury, Ali Akbar (2019) A comprehensive review on tyrosinase inhibitors. *Journal of Enzyme Inhibition and Medicinal Chemistry*, 34 (1). pp. 279-309. ISSN 1475-6366

Published by: Taylor & Francis

URL: <https://doi.org/10.1080/14756366.2018.1545767>
<<https://doi.org/10.1080/14756366.2018.1545767>>

This version was downloaded from Northumbria Research Link:
<http://nrl.northumbria.ac.uk/37532/>

Northumbria University has developed Northumbria Research Link (NRL) to enable users to access the University's research output. Copyright © and moral rights for items on NRL are retained by the individual author(s) and/or other copyright owners. Single copies of full items can be reproduced, displayed or performed, and given to third parties in any format or medium for personal research or study, educational, or not-for-profit purposes without prior permission or charge, provided the authors, title and full bibliographic details are given, as well as a hyperlink and/or URL to the original metadata page. The content must not be changed in any way. Full items must not be sold commercially in any format or medium without formal permission of the copyright holder. The full policy is available online: <http://nrl.northumbria.ac.uk/policies.html>

This document may differ from the final, published version of the research and has been made available online in accordance with publisher policies. To read and/or cite from the published version of the research, please visit the publisher's website (a subscription may be required.)

www.northumbria.ac.uk/nrl





A comprehensive review on tyrosinase inhibitors

Samaneh Zolghadri, Asieh Bahrami, Mahmud Tareq Hassan Khan, J. Munoz-Munoz, F. Garcia-Molina, F. Garcia-Canovas & Ali Akbar Saboury

To cite this article: Samaneh Zolghadri, Asieh Bahrami, Mahmud Tareq Hassan Khan, J. Munoz-Munoz, F. Garcia-Molina, F. Garcia-Canovas & Ali Akbar Saboury (2019) A comprehensive review on tyrosinase inhibitors, Journal of Enzyme Inhibition and Medicinal Chemistry, 34:1, 279-309, DOI: [10.1080/14756366.2018.1545767](https://doi.org/10.1080/14756366.2018.1545767)

To link to this article: <https://doi.org/10.1080/14756366.2018.1545767>



© 2018 The Author(s). Published by Informa UK Limited, trading as Taylor & Francis Group.



Published online: 03 Jan 2019.



Submit your article to this journal [↗](#)



Article views: 57



View Crossmark data [↗](#)

A comprehensive review on tyrosinase inhibitors

Samaneh Zolghadri^a, Asieh Bahrami^a, Mahmud Tareq Hassan Khan^b, J. Munoz-Munoz^c, F. Garcia-Molina^d, F. Garcia-Canovas^d and Ali Akbar Saboury^e

^aDepartment of Biology, Jahrom Branch, Islamic Azad University, Jahrom, Iran; ^bAura Dynamics, Tromsø, Norway; ^cGroup of Microbiology, Department of Applied Sciences, Northumbria University at Newcastle, Newcastle Upon Tyne, UK; ^dGENZ-Group of Research on Enzymology, Department of Biochemistry and Molecular Biology-A, Regional Campus of International Excellence "Campus Mare Nostrum", University of Murcia, Espinardo, Murcia, Spain; ^eInstitute of Biochemistry and Biophysics, University of Tehran, Tehran, Iran

ABSTRACT

Tyrosinase is a multi-copper enzyme which is widely distributed in different organisms and plays an important role in the melanogenesis and enzymatic browning. Therefore, its inhibitors can be attractive in cosmetics and medicinal industries as depigmentation agents and also in food and agriculture industries as antibrowning compounds. For this purpose, many natural, semi-synthetic and synthetic inhibitors have been developed by different screening methods to date. This review has focused on the tyrosinase inhibitors discovered from all sources and biochemically characterised in the last four decades.

ARTICLE HISTORY

Received 18 September 2018
Revised 4 November 2018
Accepted 5 November 2018

KEYWORDS

Tyrosinase; inhibitor; depigmentation agents; antibrowning compounds

Introduction

Browning of fruits, fungi and vegetables and hyperpigmentation in human skin are two common undesirable phenomena. Tyrosinase is the main enzyme recognised as responsible for this enzymatic browning and melanogenesis in mammals^{1,2}. This encouraged researchers and scientists to focus on the identification, isolation, synthesis and characterisation of new potent tyrosinase inhibitors for various application in the food³, cosmetics⁴ and medicinal industries. However, very few inhibitors are qualified for clinical use and skin-whitening agents. Moreover, as the clinical and industrial demands for tyrosinase inhibitors increase, *in vitro* assays and improved screening techniques are also undergoing rapid development for *in vitro* high-throughput screening tyrosinase inhibitors and putative skin-whitening agents⁵. In other words, sensitive and correct assay methods for screening and development of effective tyrosinase inhibitors are of great importance. For this purpose, several spectrophotometric^{6–10}, chromatographic^{11–17}, electrophoretic^{18–22}, radiometric^{23,24} and electrochemical^{25–27} assays have been applied and developed by researchers so far. Recently, a novel fluorescent biosensor²⁸ and tyrosinase-based thin-layer chromatography-autography have been suggested for tyrosinase inhibitor screening²⁹.

Additionally, further improvements of *in vitro* detection methods for rapidly screening tyrosinase inhibitors may be achieved through using virtual screening³⁰ and construction of quantitative structure–activity relationship (QSAR) models of inhibitors^{31,32}. Thus, a combination of bioinformatics simulation and biological *in vitro* analysis will be useful to understand the functional mechanisms of the tested compounds^{9,21,27,33–48}. Lately, Gao et al. have performed a virtual screening from Traditional Chinese medicine (TCM) and predicted tyrosinase inhibition by 3D QSAR pharmacophore models⁴⁹. For more information about successful utilisation

of computational tools like QSAR-based and ligand-based virtual screening, a review published by Khan in 2012 organised and summarised novel and potent inhibitors of the enzyme⁵⁰. Furthermore, with regard to tyrosinase inhibition importance, several other reviews have presented the organisation of tyrosinase inhibitors from natural, semi- and full synthetic sources^{1,51–62}.

The present review also focuses on the tyrosinase inhibitors discovered from all sources, including synthetic compounds, extracts and active ingredients of natural products, virtual screening and structure-based molecular docking studies published in the last four decades. We hope that the knowledge offered in this review serves as an updated comprehensive database contributing to the development of new safe and efficient anti-tyrosinase agents for the prevention of browning in plant-derived foods, seafood and hyperpigmentation treatments.

The role of tyrosinase in the melanin biosynthesis

Melanins, the main pigment primarily responsible in the skin, hair and eyes pigmentation of human, are produced by melanocytes through melanogenesis. Melanogenesis and skin pigmentation are the most important photoprotective factor in response to ultraviolet radiation damaging from the sun and skin photocarcinogenesis. The abnormal loss of melanin and depigmentation can be a serious facial esthetic and dermatological problem among human⁶³. On the contrary, the increased melanin synthesis and accumulation of these pigments occur in many types of skin disorders, including Acanthosis nigricans, Cervical Poikiloderma, melasma, Periorbital hyperpigmentation, Lentiginos, neurodegeneration associated with Parkinson's disease and skin cancer risk^{64–66}. Although melanogenesis is a complicated process represented by numerous enzymatic and chemical reactions, the enzymes such as tyrosinase and other tyrosinase-related proteins

(TYRP1 and TYRP2) have a critical role in melanin synthesis. Tyrosinase is a multifunctional copper-containing metalloenzyme with dinuclear copper ions, which plays as a rate-limiting enzyme in the synthesis of melanin (Figure 1)^{52,67}. Also, tyrosinase constitutes the primary cause for undesired browning of fruits and vegetables as well as diseases resulting from overproduction of melanin. Therefore, controlling the activity of enzyme by tyrosinase inhibitors is an essential endeavor for treating hypopigmentary disorders of mammals and enzymatic browning of fruits and fungi. To date, numerous effective inhibitors are identified and developed for using in the medical and cosmetic products, as well as food bioprocessing and agricultural industries and environmental industries. However, in medicine, tyrosinase inhibitors are a class of important clinical antimelanoma drugs but only a few compounds are known to serve as effective and safe tyrosinase inhibitors.

Mushroom tyrosinase properties

Tyrosinases have been isolated and purified from different sources such as some plants, animals and microorganisms. Although many of them (such as human) have been sequenced, only few of them have been characterised. Recently, a novel tyrosinase produced by Sahara soil actinobacteria have been isolated and biochemically characterised with the aim to identify novel enzymes with exclusive features for biotechnological applications^{68–80}. However, among different sources of tyrosinase, mushroom tyrosinase from *Agaricus bisporus* is a major and cheap source of tyrosinase with high similarity and homology compared to human tyrosinase⁷⁸. Because of these good properties, the structural, functional and biochemical characteristics of mushroom tyrosinase have been studied extensively as a model system for screening of tyrosinase inhibitors and melanogenic studies, enzyme-catalysed reactions and enzyme-inhibitor structural studies so far^{78,81–90}. Tyrosinase from *Agaricus bisporus* is a 120 kDa tetramer with two different

subunits, heavy and light⁹¹, which was the first isolated by Bourquelot and Bertrand⁹² in 1895. It has three domains and two copper binding sites which bind to six histidine residues and interact with molecular oxygen in the tyrosinase active site. Also, a disulfide linkage stabilise its structure⁹³. Recently, a 50 kDa tyrosinase isoform from *Agaricus bisporus* (H-subunit) have been purified with a high specific tyrosinase activity of more than 38,000 U/mg⁹⁴.

Reaction mechanism

Tyrosinase (EC 1.14.18.1) has two activities in its catalytic cycle, see Figure 2^{95,96}, a monophenolase activity where it hydroxylates monophenols (e.g. L-tyrosine) to *o*-diphenols (e.g. L-dopa) and a diphenolase activity where tyrosinase oxidises *o*-diphenols to *o*-quinones (*o*-dopaquinone). At the same time of these enzymatic reactions, there are different chemical reactions coupled where two molecules of *o*-dopaquinone react their-selves generating an *o*-diphenol molecule (L-dopa) and a dopachrome molecule.

Diphenolase activity can be independently studied, when tyrosinase reacts with an *o*-diphenol (see Figure 2). The form met-tyrosinase (E_m) binds the *o*-diphenol (D) originating the complex E_mD . This complex oxidises the *o*-diphenols transforming it to *o*-quinone and the enzyme is converted into the form deoxy-tyrosinase (E_d). E_d has a very big affinity for the molecular oxygen originating the form oxy-tyrosinase (E_{ox}), which binds another *o*-diphenol molecule and originating the complex $E_{ox}D$. After that, the *o*-diphenol is oxidised again to *o*-quinone and the form E_m is formed again completing the catalytic cycle. However, after these enzymatic reactions, two *o*-quinone molecules (e.g. *o*-dopaquinone) react generating dopachrome and regenerating a molecule of *o*-diphenol.

As mentioned before, we can independently study the diphenolase activity. However, it is not applicable for the monophenolase activity, see Figure 2, because the chemical reactions of

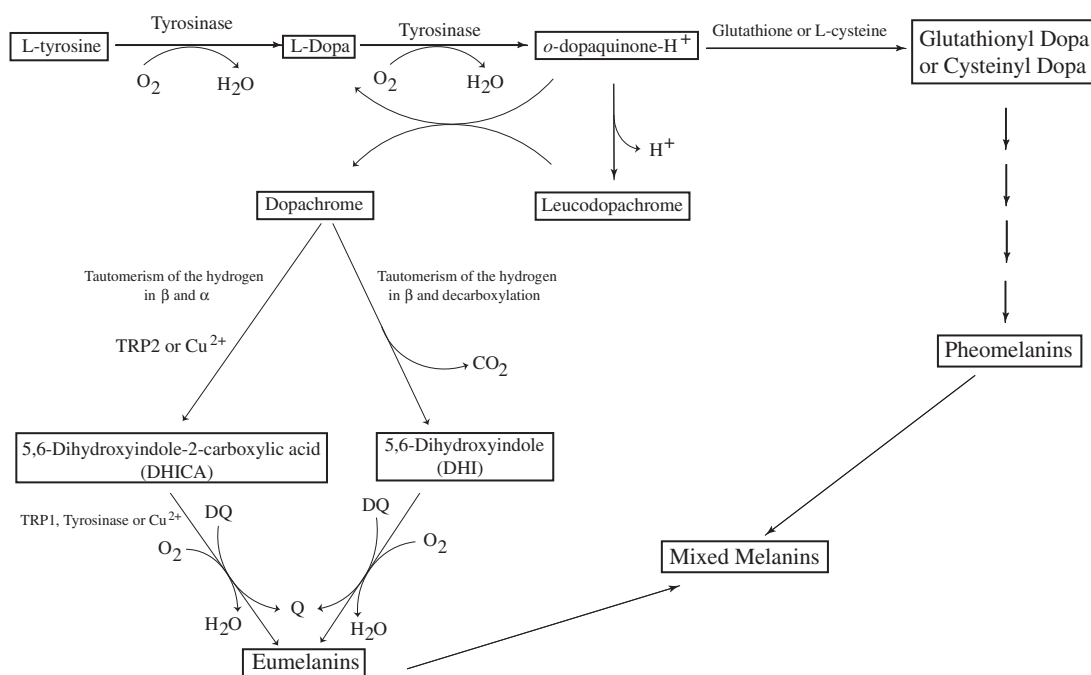


Figure 1. Scheme of the biosynthetic pathway of eumelanins and pheomelanins. The activities of tyrosinase are indicated in the scheme. Moreover, the enzyme can oxidize DHICA to its *o*-quinone directly, or it can oxidize DHICA and DHI indirectly via the formation of *o*-dopaquinone. TRP2 (dopachrome tautomerase) or Cu^{2+} can participate in the evolution of dopachrome to DHICA. The oxidation of DHICA can be catalyzed by TRP1, (DHICA oxidase), tyrosinase or Cu^{2+} . When glutathione or L-cysteine attack *o*-dopaquinone, glutathione-dopa or cysteinyl-dopa adducts are formed and these later evolve to pheomelanins⁶⁷.

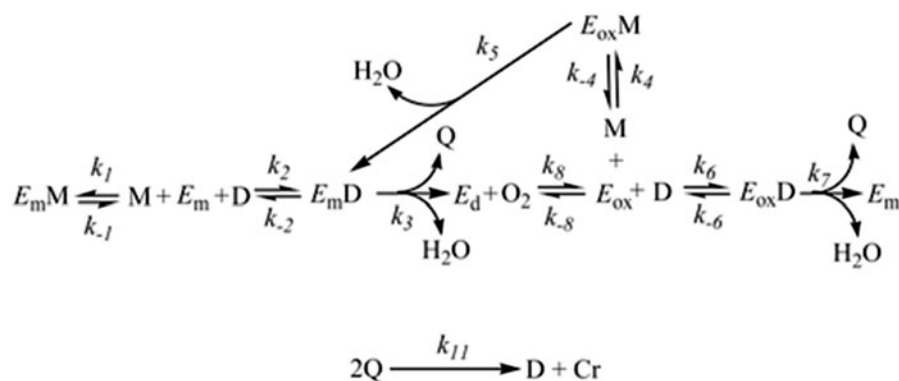


Figure 2. Monophenolase and diphenolase activities of Tyrosinase. E_mM , met-tyrosinase/monophenol complex; M , monophenol; D , *o*-diphenol; E_m , met-tyrosinase; E_mD , met-tyrosinase/*o*-diphenol complex; E_d , deoxy-tyrosinase; O_2 , molecular oxygen; E_{ox} , oxy-tyrosinase; $E_{ox}D$, oxy-tyrosinase/*o*-diphenol complex; $E_{ox}M$, oxy-tyrosinase/monophenol complex; Q , *o*-quinone; Cr , Dopachrome.

diphenolase activity have to occur at the same time of monophenolase activity. Tyrosinase shows the monophenolase activity with a lag period. This period is the time that the enzyme requires to accumulate a quantity of *o*-diphenol in reaction medium and is proportional to the quantity of monophenol used. Figure 2 shows the new complexes appeared in the monophenolase activity: $E_{ox}M$ (oxy-tyrosinase bound to monophenol) and E_mM (met-tyrosinase bound to monophenols). $E_{ox}M$ is active and is transformed into E_mD , which is an intermediate of the catalytic cycle⁹⁵. *o*-Quinones formed by these two oxidation cycle spontaneously react with each other to form oligomers⁹⁷.

Tyrosinase inhibition

Due to the critical role of tyrosinase in the melanogenesis and browning process, several investigations have been reported for the identification of tyrosinase inhibitor from both natural (fungi, bacteria, plants) and synthetic sources so far. General speaking, tyrosinase inhibitors are examined in the presence of a monophenolic substrate such as tyrosine or a diphenolic substrate such as *L*-dopa, and activity is assessed based on dopachrome formation.

Inhibition mechanism

Among different types of compounds such as specific tyrosinase inactivators and inhibitors, *o*-dopaquinone scavengers, alternative enzyme substrates, nonspecific enzyme inactivators and denaturants, only specific tyrosinase inactivators and reversible inhibitors actually bind to the enzyme as true inhibitors and really inhibit its activity:

- Specific tyrosinase inactivators. They are called suicide inactivators or mechanism-based inhibitors. This group of compounds can be considered very interested from a pharmacological point of view, in hyperpigmentation processes (Figure 3)⁹⁸.

To explain the suicide inactivation of tyrosinase, mainly two mechanisms have been proposed^{98,99}. Accordingly, Haghbeen et al. have suggested that the conformational changes, triggered by the substrate then mediated by the solvent molecules, in the tertiary and quaternary structures of tyrosinase, might be the real reason for the suicide inactivation¹⁰⁰. On the other hand, however, based on reports, it was found that acetylation of tyrosine residues with *N*-acetylimidazole protects mushroom tyrosinase from the suicide inactivation in the presence of its catecholic substrate,

4-[(4-methylbenzo) azo]-1,2-benzenediol without any major impact on the secondary structure of enzyme¹⁰¹.

The studies about the kinetics of suicide inactivation of tyrosinase have been carried out with several *o*-diphenolic substrates¹⁰², ascorbic acid¹⁰³, *L*- and *D*-dopa¹⁰⁴ and with different aminophenols and *o*-diamines¹⁰⁵. The authors have established that the suicide inactivation could occur after the transference of a proton to the peroxide group on the active site of oxy-tyrosinase^{98,106}, also it has been proposed that the monophenols do not inactivate the enzyme^{107,108}.

The chemical structure of the different substrates is diverse, but the process always requires a step of oxidation/reduction: *o*-diphenols^{102,104}, ascorbic acid¹⁰³, aminophenols and *o*-diamines¹⁰⁵, hydroxyhydroquinone¹⁰⁹, tetrahydrobiopterines¹¹⁰, tetrahydrofolic acid¹¹¹ and NADH¹¹².

- Generally, the mode of inhibition by "true inhibitors" is one of these four types: competitive, uncompetitive, mixed type (competitive/uncompetitive), and noncompetitive. A competitive inhibitor can bind to a free enzyme and prevents substrate binding to the enzyme active site. Regarding the property that tyrosinase is a metalloenzyme, copper chelators such as many aromatic acids, phenolic and poly-phenolic compounds, a few non-aromatic compounds, can inhibit tyrosinase competitively by mimicking the substrate of tyrosinase^{52,60}. Recently, it was found that *D*-tyrosine negatively regulates melanin synthesis by inhibiting tyrosinase activity, competitively¹¹³. In addition, *L*-tyrosine has been shown as an inhibitor¹¹⁴.

In contrast, an uncompetitive inhibitor can bind only to the enzyme-substrate complex and a mixed (competitive and uncompetitive mixed) inhibitor can bind to both forms of free enzyme and enzyme-substrate complex. Finally, noncompetitive inhibitors bind to a free enzyme and an enzyme-substrate complex with the same equilibrium constant¹¹⁵. Non-competitive and mixed-inhibition are frequent modes observed in the kinetics studies on mushroom tyrosinase activities. Phthalic acid and cinnamic acid hydroxypyridinone derivatives¹¹⁶ are two examples of mixed type inhibitors of mono-phenolase activity¹¹⁷. Also, some compounds such as phthalic acid⁴⁶ and terephthalic acid¹¹⁸, *D*-(-)-arabinose¹¹⁹, brazilein¹²⁰, thymol analogs¹²¹ were demonstrated as mixed-type effector examples of di-phenolase activity. Furthermore, other compounds such as bi-pyridine derivatives¹²², two thiazazole derivatives⁴⁴ barbarin¹²³, chlorocinnamic acids¹²⁴, propanoic acid¹²⁵, some *N*-(mono- or dihydroxybenzyl)-*N*-nitrosohydroxylamines¹²⁶

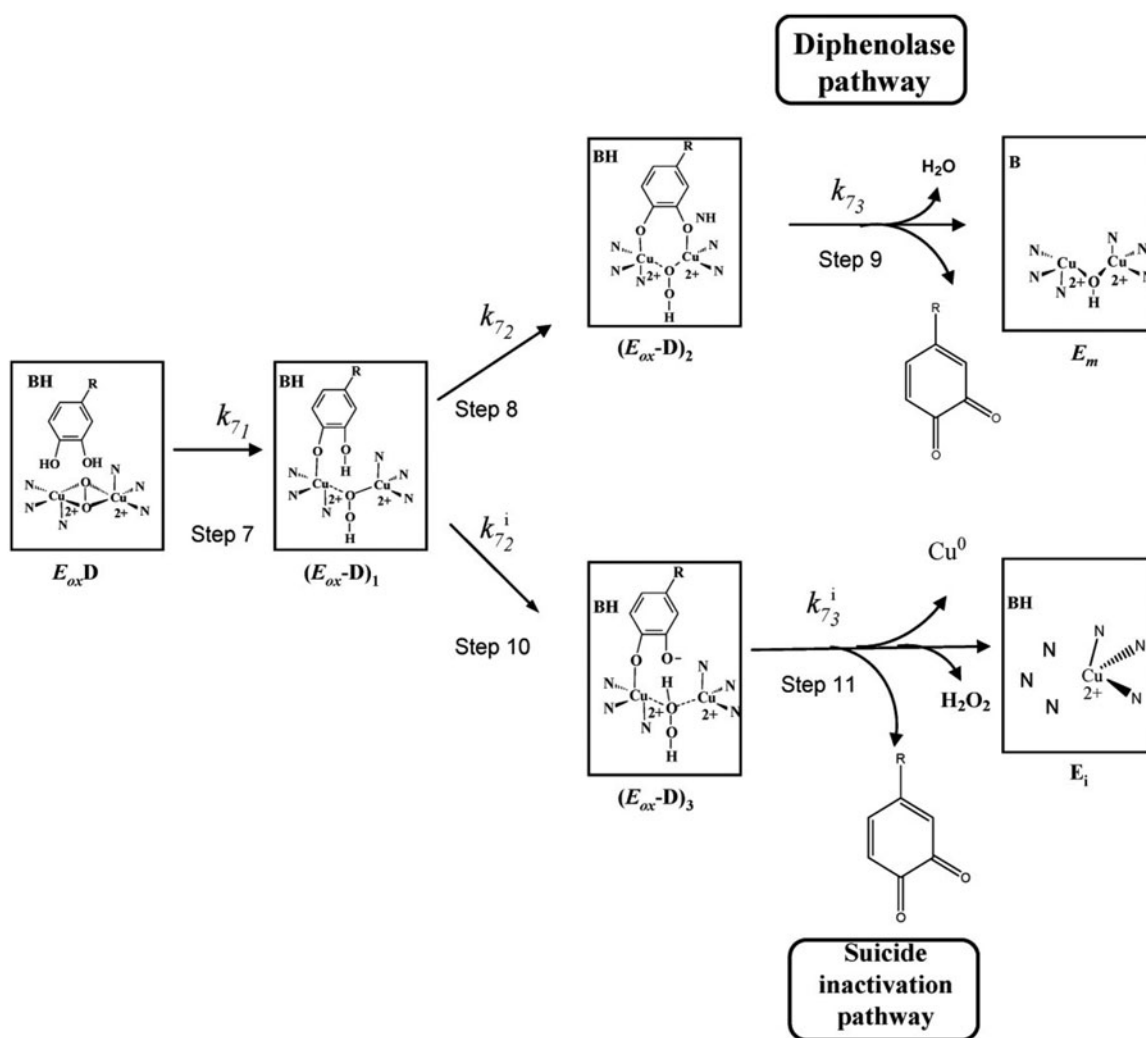


Figure 3. Detail of the structural mechanism proposed to explain the suicide inactivation of tyrosinase during its action on *o*-diphenols. E_m , met-tyrosinase; E_{ox} , oxy-tyrosinase; $E_{ox}D$, oxy-tyrosinase/*o*-diphenol complex; $(E_{ox}D)_1$, oxy-tyrosinase/*o*-diphenol complex axially bound to a Cu atom; $(E_{ox}D)_2$, oxy-tyrosinase/*o*-diphenol complex axially bound to the two Cu atoms; $(E_{ox}D)_3$, oxy-tyrosinase/*o*-diphenol complex axially bound to one Cu atom and the deprotonated hydroxyl group of C-3; E_i , inactive form of tyrosinase. A general view of this scheme is shown in Ref ⁹⁸.

and *p*-alkylbenzaldehydes¹²⁷ inhibited catecholase activity of mushroom tyrosinase uncompetitively. Some derivatives of thiazoles are examples for noncompetitive tyrosinase inhibition¹²⁸.

In addition to determining the inhibition mechanism, inhibitory strength which is expressed as the IC_{50} value (the concentration of inhibitor at which 50% of your target is inhibited) should be calculated in the enzyme kinetics studies and inhibitor screening to compare the inhibitory strength of an inhibitor with others. However, the IC_{50} values may be incomparable due to the varied assay conditions (different substrate concentrations, incubation time, and different sources of tyrosinase) but a positive control can be used for this purpose⁵². Although, some researchers have not calculated IC_{50} and have not applied a positive control in their studies but, fortunately, in most studies conducted for screening new tyrosinase inhibitors, the popular whitening agents, such as kojic acid, arbutin or hydroquinone, were used as a positive control¹²⁹ at the same time. However, among different types of mushroom tyrosinase inhibitors, some inhibitors such as hydroquinone⁴⁹ arbutin, kojic acid^{15,49}, azelaic acid, L-ascorbic acid, ellagic acid and tranexamic acid have been reported as skin-whitening agents in the cosmetic industry but there are a few reports failed to confirm their effect as an agent to lighten skin in clinical trials despite the safety of this compound⁵.

Recently, Mann et al., have compared the inhibitory effects of hydroquinone, arbutin and kojic acid by human tyrosinase and mushroom tyrosinase. They have found hydroquinone and arbutin and kojic acid ($IC_{50} > 500 \mu\text{mol/L}$) weakly inhibits human tyrosinase. In contrast, a resorcynyl-thiazole derivative, thiamidol, is a most potent inhibitor of human tyrosinase (IC_{50} of $1.1 \mu\text{mol/L}$) but inhibits mushroom tyrosinase weakly ($IC_{50} = 108 \mu\text{mol/L}$)¹³⁰. Also, deoxyarbutin, a novel reversible tyrosinase inhibitor with effective *in vivo* skin lightening potency, have been reported due to its increased skin penetration and binding affinity to human tyrosinase¹³¹. In another research, Sugimoto et al. have investigated a comparison of inhibitory effects of alpha-arbutin and arbutin with human tyrosinase and they have found α -arbutin is stronger than arbutin¹³².

Natural tyrosinase inhibitor sources

Natural sources including plants, bacteria and fungi have recently become of increasing interest for their antityrosinase activity by producing bioactive compounds. A number of researchers prefer to identify inhibitors from natural sources due to their less toxicity and better bioavailability, especially for food, cosmetic and medicinal applications.

Plants

It is well known that phenolic compounds are the largest group of phytochemicals found in plants, which are mainly the factors responsible for the activities in plant extracts⁵². Tyrosinase inhibitory activity of many plant extracts was carried out to find new sources of anti-tyrosinase compounds. For example, anti-tyrosinase activities of the following plants have been reported by various researchers: *Asphodelus microcarpus*¹³³, *Morus nigra* L.¹³⁴, *Greyia radlkoferi* Szyszyl⁴⁵, *Limonium tetragonum*¹³⁵, *Arctostaphylos uva-ursi*¹³⁶, *Pleurotus ferulae*¹³⁷, *Agastache rugosa* Kuntze fermented with *Lactobacillus rhamnosus* and *Lactobacillus paracasei*¹³⁸, *Artemisia aucheri* Boiss¹³⁹, *Cassia tora*¹⁴⁰, *S. brevibracteata* subsp¹⁴¹, *Rhodiola crenulata*, *Alpinia officinarum* Hance and *Zanthoxylum bungeanum* Maxim¹⁴², *Mangifera indica*¹⁴³, *Podocarpus falcatus*¹⁴⁴, *Momordica charantia*¹⁴², *Cymbopogon citrates*¹⁴⁵, *Greyia flanaganii* (IC₅₀ = 32.62 µg/ml)¹⁴⁶, *Vitis vinifera* Leaf extracts (IC₅₀ = 3.84 mg/ml)¹⁴⁷ and *Inula britannica* L.¹⁴⁶. Also, tyrosinase inhibitory activity of 91 native plants from central Argentina was carried out by Chiari et al.^{138,147}. Their results approved the inhibitory activity of these extracts against tyrosinase: *Achyrocline satureioides*, *Artemisia verlotiorum*, *Cotoneaster glaucophylla*, *Dalea elegans*, *Flourensia campestris*, *Jodina rhombifolia*, *Kageneckia lanceolata*, *Lepechinia floribunda*, *Lepechinia meyenii*, *Lithrea molleoides*, *Porlieria microphylla*, *Pterocaulon alopecuroides*, *Ruprechtia apetala*, *Senna aphylla*, *Sida rhombifolia*, *Solanum argentinum*, *Tagetes minuta*, and *Thalictrum decipiens*. Besides, plants from the *Moraceae* family including genera *Morus* species, *Artocarpus*, *Maclura* (*Cudrania*), *Broussonetia*, *Milicia* (*Chlorophora*), and *Ficus* have shown *in vitro* tyrosinase inhibition¹⁴⁸. Also, ethanolic and methanolic extracts of some other plants such as *Ardisia elliptica* Thunb¹⁴⁹, *Phyllanthus acidus* (L.) Skeels, *Rhinacanthus nasutus* L. Kurz (IC₅₀ value of 271.50 µg/ml), *Arbutus andrachne* L. (IC₅₀ = 1 mg/mL)¹⁵⁰, *Withania somnifera* L. Dunal and *Solanum nigrum* L. berries¹⁵¹, *Pulmonaria officinalis* and *Centarium umbellatum*¹⁵² and Camel's foot creeper leaves (*Bauhinia vahlii*)¹⁵³ significantly inhibited tyrosinase activity, too. Quispe et al. have screened tyrosinase inhibitory properties of Peruvian medicinal plants. Among these plant extracts, *Hypericum laricifolium* Juss, *Taraxacum officinale* F.H.Wigg. (IC₅₀ value of 290.4 µg/ml), and *Muehlenbeckia vulcanica* Meisn (IC₅₀ value of 280.1 µg/ml) showed the greatest anti-tyrosinase activity¹⁵⁴. Furthermore, tyrosinase inhibitory activity of mangrove plants in Micronesia¹⁵⁵, Korean indigenous plants¹⁵⁶, plants from Brazilian Cerrado¹⁵⁷, five traditional medicinal plants from Iran¹⁵⁸, ethanol extracts from medicinal and edible plants cultivated in Okinawa¹⁵⁹, seashore plants¹⁶⁰, some tropical plants¹⁶¹ and Bangladeshi indigenous medicinal plants¹⁶², have been investigated by various researchers. Bonesi et al. have reported recent trends in the discovery of tyrosinase inhibitors from plant sources¹⁶³.

Fungi and bacteria

Fungi from different genera such as *Aspergillus* sp.¹⁶⁴, *Trichoderma* sp.¹⁶⁵, *Paecilomyces* sp.¹⁶⁶, *Phellinus linteus*¹⁶⁷, *Daedalea dickinsii*¹⁶⁸, *Dictyophora indusiata*¹⁶⁹ along with a liquid culture of *Neolentinus lepideus*¹⁷⁰ have been reported as a source of novel tyrosinase inhibitor by producing bioactive compounds. Also, there have been several reports on tyrosinase inhibitors from some marine fungi species such as *Myrothecium* sp. isolated from algae¹⁷¹ and *Pestalotiopsis* sp. Z233¹⁷². Also, there are several reports on tyrosinase inhibition by bacterial species and their metabolites.

Among them, *Streptomyces* sp., such as *S. hiroshimensis* TI-C3 isolated from soil¹⁷³, an actinobacterium named *Streptomyces swartbergensis* sp. Nov.¹⁷⁴ and *Streptomyces roseolilacinus* NBRC 12815¹⁷⁵ are potential bacterial sources of tyrosinase inhibitors. Moreover, some tyrosinase inhibitors have been reported from a gram-negative marine bacterium *Thalassotalea* sp. Pp2-459¹⁷⁶ and a toxic strain of the cyanobacterium, *Oscillatoria agardhii*¹⁷⁷. Interestingly, some probiotics such as *Lactobacillus* sp.¹⁷⁸ which are used in the fermentation process have been investigated as natural tyrosinase inhibitor sources. Based on the studies, it has been confirmed that the physiological activities of fermented extracts are considerably higher than those of unfermented extracts and their cytotoxic activity is lower as compared to unfermented extracts¹⁷⁹. Recently, tyrosinase inhibitory four different lactic acid bacteria (LAB) strains isolated from dairy cow feces have been proved by Ji et al.¹⁸⁰.

Finally, in an updated review by Fernandes from reported findings, tyrosinase inhibitors produced by microorganisms have been summarised⁶¹. This review shows that diverse tyrosinase inhibitors isolated from plant sources and fungi are mostly phenolic compounds, steroids, and alkaloids structurally comparable with each other. In contrast, tyrosinase inhibitors from bacteria comprise a smaller group of alkaloids, macrolides, and polyphenols, which competitively inhibit the enzyme⁶¹.

Inhibitors from natural, semisynthetic and synthetic sources

Simple phenols

Phenolic compounds which are characterised by having at least one aromatic ring and one (or more) hydroxyl group can be classified based on the number and arrangement of their carbon atoms. These compounds are commonly found to be conjugated to sugars and organic acids. Phenolics range from simple to large and complex tannins and derived polyphenols due to their molecular-weight and number of aromatic-rings¹⁸⁰.

The simple phenols such as hydroquinone^{181,182} and its derivatives^{183,184}, deoxyarbutin^{185,186} and its derivatives¹⁸⁷, 4-(6-Hydroxy-2-naphthyl)-1,3-benzendiol, resorcinol (or resorcin)¹⁸⁸ and 4-n-butylresorcinol¹⁸⁹, vanillin¹⁹⁰ and its derivatives^{191,192} have been reported in the scientific literature as possible phenolic inhibitors of the tyrosinase (Figure 4). Chen et al. have found the alkyhydroquinone 10'(Z)-heptadecenylhydroquinone, isolated from the sap of the lacquer tree *Rhus succedanea*, can inhibit the

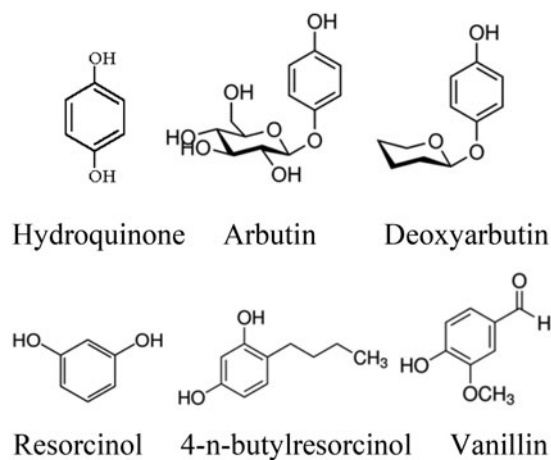


Figure 4. Chemical structures of some simple phenolic compounds.

activity of tyrosinase and suppress melanin production in animal cells. The IC_{50} of this compound (37 μ M) is less than hydroquinone (70 μ M) as a known inhibitor of tyrosinase. They have suggested that the potent inhibitory effect of this derivative on tyrosinase activity is likely due to its heptadecenyl chain, which facilitates the oxidation of the hydroquinone ring^{183,184}.

Isotachioside, a methoxy-hydroquinone-1-O-beta-D-glucopyranoside isolated from *Isotachis japonica* and *Protea neriifolia* and its glycoside derivatives (glucoside, xyloside, cellobioside, and maltoside) are categorised as analogs of arbutin. However, isotachioside and arbutin could not be determined as potent inhibitor. But, glucoside, xyloside, cellobioside and maltoside derivatives, missing methyl and benzoyl groups, acted as tyrosinase inhibitors with IC_{50} s of 417, 852, 623 and 657 μ M, respectively. Among these novel inhibitors, glucoside derivative (IC_{50} = 417 μ M) was the most potent, indicating that the structural combination of resorcinol and glucose was significant for inducing the inhibitory effect¹⁹³.

Hydroquinone and some of its known derivatives, including α and β -arbutin, are described as both a tyrosinase inhibitor and a substrate^{194,195}. Deoxyarbutin and its second-generation derivatives have been proposed as promising agents to ameliorate hyperpigmented lesions or lighten skin due to less toxicity at their effective inhibitory dose^{185,186}.

Monophenolic compounds such as L-tyrosine, L- α -methyl-tyrosine and tyramine are substrates of tyrosinase. *o*-Quinone evolves in the medium of reaction accumulating *o*-diphenol and this accumulation provokes that met-tyrosinase (E_m) is transformed into oxy-tyrosinase (E_{ox}), which is the active form of the tyrosinase for monophenols and *o*-diphenols. Therefore, tyrosinase is active with monophenols such as: umbelliferone¹⁹⁶, hydroquinone^{197,198}, *p*-hydroxybenzyl alcohol¹⁹⁹, 4-hexylresorcinol²⁰⁰, oxyresveratrol²⁰¹, 4-*n*-butylresorcinol²⁰², resorcinols²⁰³, α and β -arbutin¹⁹⁵ and *p*-coumaric acid^{204,205} when we add the following reagents to medium of reaction: hydrogen peroxide (transforms E_m to E_{ox}), an *o*-diphenol or a reducing agent such as ascorbic acid transforming E_m to E_d which, with molecular oxygen, is transformed into E_{ox} . A particular case is deoxyarbutin, which acts as a substrate of tyrosinase even if any reagent is not added to the medium of reaction²⁰⁶. Taking into consideration all the previous comments, several methods have been developed to discriminate between true inhibitors and alternative substrates of the enzyme^{98,207}.

Polyphenols

Plants produce a large diverse class of polyphenols including phenolic acids, flavonoids, stilbenes and lignans^{208,209}. A large number of these compounds have been reported as a weak or potent inhibitor of tyrosinase from natural^{210–215} and synthetic^{216–219} sources.

Flavonoids

Among polyphenolic compounds, some of the flavonoid derivatives mostly found in herbal plants, fruits and synthetic sources have been revealed to be the potent inhibitors of tyrosinase^{133,211,220–225}. There is a significant correlation between the inhibitory potency of flavonoids on mushroom tyrosinase and melanin synthesis in melanocytes²²⁶. In searching effective tyrosinase inhibitors from natural products, many flavonoid compounds have been isolated and evaluated for their inhibitory activity on mushroom tyrosinase from different natural sources such as *Trifolium nigrescens* Subsp. *Petrisavi*²²⁷, mung bean (*Vigna radiata* L.)²²⁸, *calamondin* peel²²⁹,

*Morus yunnanensis*²³⁰, *Bhagwa* and *Arakta cultivar*²³¹, *Tibouchina semidecandra* L.²³², *Maackia faurie*²³², *Pleurotus ostreatus*²³³, *Potentilla bifurca*²³⁴, *Alpinia officinarum*²³⁵, roots of *Morus lhou*²³⁶, *Garcinia subelliptica*¹⁶⁰, *Artocapus altilis*¹⁹⁰, *Myrsine africana*²³⁷, *Pulsatilla cernua*²³⁸, *Salvia miltiorrhiza-Carthamus tinctorius* (Danshen-Honghua, DH) herbal pair²³⁹ and other various medicinal plants²⁴⁰.

Generally, major flavonoids (Figure 5) are classified into several main classes: flavones, flavonols, isoflavones, flavanones, flavanols and anthocyanidins. Minor flavonoids included: dihydroflavones, flavan-3,4-diols, coumarins, chalcones, dihydrochalcones and aurones²⁴¹. Also, prenylated and vinylylated flavonoids, such as flavonoid glycosides, are other subclasses of flavonoids. Some flavonoid glycosides such as myricetin 3-galactoside and quercetin 3-O- β -galactopyranoside from *Limonium tetragonum*¹³³ and 3',5'-di-C- β glucopyranosylphloretin from unripe calamondin peel (IC_{50} = 0.87 mg/ml)²²⁹, have been investigated for their inhibitory activities on tyrosinase. Moreover, the inhibitory activities of some other prenylated and vinylylated flavonoids, such as kuwanon C, papyriflavonol A, sanggenon D and sophoflavescenol, and sanggenon D (IC_{50} = 7.3 μ M) against tyrosinase, have been approved by Lee et al.²⁴². However, according to their findings, the prenylation with isoprenyl group or the vinylation of some flavonoid molecules does not enhance their tyrosinase inhibitory activity²⁴². Interestingly, it has even demonstrated that deglycosylation of some flavonoid glycosides by far-infrared irradiation can be improved tyrosinase inhibitory activity²⁴³. In a survey from reported findings (2008–2013), Orhan et al. reviewed many examples of tyrosinase inhibitors with flavonoid structure²²⁰. In the following, some tyrosinase inhibitors from various flavonoid classes have been mentioned and discussed.

Flavones and dihydroflavones. The most common flavones are luteolin, apigenin, baicalein, chrysin and their glycosides (e.g. apigenitrin, vitexin, and baicalin)²⁰⁹. Furthermore, nobiletin and tangeretin are the polymethoxylated flavones²⁴⁴. Nguyen et al. have investigated the presence of apigenin and nobiletin from the methanolic extract of the heartwood of *Artocapus altilis* with 11 other phenolic compounds for their inhibitory activities on tyrosinase¹⁹⁰. In another research, Shang et al. have found a derivative of flavone, namely 7,8,4-trihydroxyflavone which inhibits diphenolase activity of tyrosinase with an IC_{50} value of $10.31 \pm 0.41 \mu$ M and a noncompetitive manner with a K_i of $9.50 \pm 0.40 \mu$ M. The quenching analysis of tyrosinase by this compound showed a static mechanism and a single binding site with a binding constant of $7.50 \pm 1.20 \times 10^4 \text{ M}^{-1}$ at 298 K. Based on the thermodynamics parameters, the binding process involved hydrogen bonds and van der Waals forces. Also, docking simulation illustrated hydrogen bonds between this compound and the residues His244 and Met280 of active site²⁴⁵.

In addition, several hydroxyflavones including baicalein, 6-hydroxyapigenin, 6-hydroxygalangin and 6-hydroxykaempferol²⁴⁶ and tricrin (5,7,4'-trihydroxy-3',5'-dimethoxyflavone)²⁴⁷ have been demonstrated as inhibitors of diphenolase activity of tyrosinase. The mechanism of inhibition by baicalein (IC_{50} = 0.11 mM) indicated a mix-type (K_i of 0.17 mM, α = 0.56). A single binding site with a binding constant of $2.78 \times 10^5 \text{ M}^{-1}$ was obtained from the quenching fluorescence analysis for this compound. Thermodynamic parameters suggested spontaneous binding through hydrogen bonding and van der Waals forces. Furthermore, circular dichroism spectra indicated a reduction in the content of α -helix from 32.67% to 29.00% due to this binding. Docking simulations also indicated that baicalein mainly bound tyrosinase via its Met280 residue²⁴⁸. While, tricrin was found as a noncompetitive inhibitor of tyrosinase with good efficacy

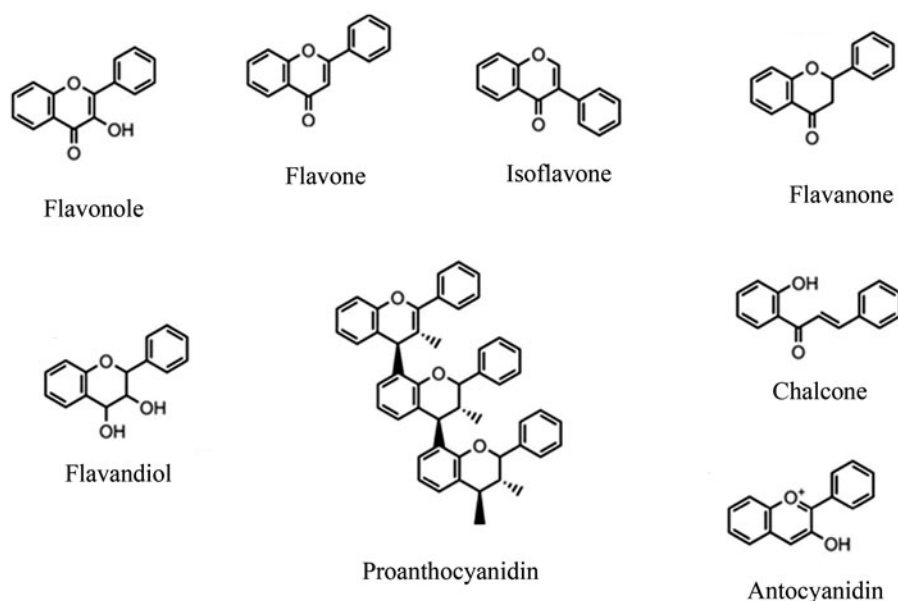


Figure 5. Structure of the main classes of flavonoids.

compared to its control. Based on circular dichroism spectra, the interactions between triclin and tyrosinase did not change the secondary structure. Fluorescence quenching revealed that the interaction of triclin with residues in the hydrophobic pocket of tyrosinase is stabilised by hydrophobic interactions and hydrogen bonding. Also, docking results implied that the stereospecific effects of triclin on substrates or products and flexible conformation alterations of tyrosinase produced by weak interactions between triclin and this enzyme are the possible inhibitory mechanisms of this compound²⁴⁷.

Another flavone named morusone from the twigs of *Morus alba* L. ($IC_{50} = 290.00 \pm 7.90 \mu\text{M}$)²⁴⁹, a new bioflavone 4''',5,5'',7,7''-pentahydroxy-3',3'''-dimethoxy-3-O- β -D-glucosyl-3',4'-O-biflavone from *Trifolium nigrescens* Subsp. Petrisavi²²⁷, along with apigenin, flavone glucoside vitexin ($IC_{50} = 6.3 \text{ mg/ml}$) and a C-glycosylflavone isovitexin ($IC_{50} = 5.6 \text{ mg/ml}$) from *Vigna radiatae* L. extracts exhibited significant tyrosinase inhibition activities²²⁸. Also, inhibitory effects of five flavones including mormin ($IC_{50} = 0.088 \text{ mM}$), cyclomorusin ($IC_{50} = 0.092 \text{ mM}$), morusin ($IC_{50} = 0.250 \text{ mM}$), kuwanon C ($IC_{50} = 0.135 \text{ mM}$) and norartocarpetin ($IC_{50} = 1.2 \mu\text{M}$) isolated from the stem barks of *Morus lhou* (S.) Koidz, have been investigated by Ryu et al. The mechanism of inhibition indicated that mormin, cyclomorusin, kuwanon C and norartocarpetin inhibited tyrosinase competitively²⁵⁰.

Flavonoles. Myricetin, kaempferol, quercetin, morin, isorhamnetin, galangin and their glycosides (e.g. rutin, quercitrin, and astragalin) are the predominant flavonols most commonly found as O-glycosides²⁰⁹. So far, several flavonols such as kaempferol from *Hypericum laricifolium* Juss¹⁵⁴ and *Crocus sativus* L.²⁵¹, quercetin from *Olea europaea* L.²⁵², quercetin-4'-O-beta-D-glucoside from *Potentilla bifurca*²⁵³, quercetin-3-O-(6-O-malonyl)- β -D-glucopyranoside and kaempferol-3-O-(6-O-malonyl)- β -D-glucopyranoside from mulberry leaves²⁵³, galangin from *Alpinia officinarum*²³⁵, morin²⁵⁴ and (\pm) 2,3-cis-dihydromorin ($IC_{50} = 31.1 \mu\text{M}$), 2,3-trans-dihydromorin ($IC_{50} = 21.1 \mu\text{M}$) from *Cudrania cochinchinensis*²⁵⁵, were identified as tyrosinase inhibitors.

Based on kinetics studies, morin reversibly inhibited tyrosinase through a multi-phase kinetic process and bind to tyrosinase at a single binding site mainly by hydrogen bonds and van der Waals

forces. It inhibited tyrosinase reversibly in a competitive manner with $K_i = 4.03 \pm 0.26 \text{ mM}$ and the binding of morin to tyrosinase-induced rearrangement and conformational changes of the enzyme²⁵⁴. Furthermore, it was reported that three flavonols including galangin²³⁵, kaempferol²⁵¹ and quercetin inhibit the oxidation of L-DOPA catalysed by mushroom tyrosinase and presumably this inhibitory activity comes from their copper chelating ability. While their corresponding flavones, chrysin, apigenin and luteolin, are not identified as copper chelator, Kubo et al. believed that the chelation mechanism by flavonols may be attributed to the free 3-hydroxyl group²⁵¹. Interestingly, quercetin behaves as a cofactor and does not inhibit monophenolase activity. In contrast, galangin inhibits monophenolase activity and does not act as a cofactor, and kaempferol neither acts as a cofactor nor inhibits monophenolase activity. However, inhibiting of diphenolase activity by chelating copper in the enzyme is the common feature of these three flavonols¹⁶⁰.

Recently, 8-prenylkaempferol as a competitive tyrosinase inhibitor along with Kushenol A (noncompetitive) isolated from *Sophora flavescens*²⁵⁶, have been investigated with IC_{50} values less than $10 \mu\text{M}$. Finally, based on the literature review, many flavonol inhibitors are usually competitive inhibitors due to the 3-hydroxy-4-keto moiety of the flavonol structure, which chelates the copper in the active site²⁵¹. Also, among all these compounds, quercetin-4'-O-beta-D-glucoside with a IC_{50} value of $1.9 \mu\text{M}$ is revealed stronger tyrosinase inhibition than their positive control, kojic acid²³⁶. While the other flavonol inhibitors listed above are very weak inhibitors and have little potential as skin whitening or food antibrowning.

Isoflavones. Isoflavones such as daidzein, genistein, glycitein, formononetin, and their glycosides (e.g. genistin, daidzin) mostly are detected in the medicinal herbs²⁰⁹. Park et al. have investigated tyrosinase inhibition activities of some natural o-dihydroxyisoflavone derivatives with variable hydroxyl substituent at the aromatic ring of isoflavone isolated from five-year-old Korean fermented soybean paste. They have demonstrated that two derivatives 7,8,4'-trihydroxyisoflavone and 7,3',4'-trihydroxyisoflavone inhibit tyrosinase by IC_{50} value of $11.21 \pm 0.8 \mu\text{M}$ and $5.23 \pm 0.6 \mu\text{M}$, respectively, whereas very low inhibition activity was obtained for

6,7,4'-trihydroxyisoflavone, daidzein, glycitein and genistein²⁵⁷. Also, 6,7,4'-trihydroxyisoflavone was identified as a potent competitive inhibitor of monophenolase activity of tyrosinase by Chang et al., with an IC_{50} value of $9.2 \mu\text{M}$, which is six times potent than kojic acid²⁵⁸. But, its analogs, glycitein, daidzein, and genistein showed little anti-tyrosinase activity. Therefore, they have suggested that C-6 and C-7 hydroxyl groups of the isoflavone skeleton might play an important role in the tyrosinase inhibitory activity. Furthermore, two other isoflavone metabolites, 7,8,4'-trihydroxyisoflavone and 5,7,8,4'-tetrahydroxyisoflavone isolated from soygerm koji, were investigated by Chang et al.²⁵⁹. These compounds inhibited both monophenolase and diphenolase activities with an irreversible inhibition manner. Interestingly, by using HPLC analysis and kinetic studies, they have found that 7,8,4'-trihydroxyisoflavone and 5,7,8,4'-tetrahydroxyisoflavone are potent suicide substrates of mushroom tyrosinase. It may be concluded that the hydroxyl groups at both the C7 and C8 positions could completely change the inhibitory mechanism of the isoflavones from the reversible competitive to the irreversible suicide form⁵².

Recently, a noncompetitive inhibitor, glabridin ($IC_{50} = 0.43 \mu\text{M}$), isolated from the root of *Glycyrrhiza glabra* Linn, has exhibited excellent inhibitory effects on tyrosinase. The quenching analysis of tyrosinase by glabridin showed a static mechanism²⁶⁰. Notably, a drug delivery system by using glabridin microsphere-loaded gel as a new approach for hyperpigmentation disorders have been proposed by Deshmukh et al.²⁶¹. In another research, Jirawattanapong et al. have identified a synthetic glabridin, 3'',4''-dihydroglabridin, with higher activity than glabridin ($IC_{50} = 11.40 \mu\text{M}$) against tyrosinase. They have suggested the more effective interaction with the enzyme may be due to more conformational flexibility of this compound that has occurred by the 4-substituted resorcinol skeleton and the lacking of double bond between carbon atom 3'' and 4'' in its structure²⁶². Also, Nerya et al. have reported that another isoflavone, glabrene, in the licorice extract can inhibit both monophenolase and diphenolase tyrosinase activities²⁶³. In the study reported by Heo et al., two new isoflavones desmodianone H and uncianone B have been identified as novel tyrosinase inhibitors. However, uncianone B has higher anti-tyrosinase rate than desmodianone H²⁶⁴. Glyasperin C from *Glycyrrhiza glabra* is another kind of isoflavone identified as tyrosinase inhibitor²⁶⁵. Furthermore, some other isoflavones, formononetin, genistein, daidzein, texasin, tectorigenin, odoratin and mirkoin isolated from the stems of *Maackia fauriei*, have been investigated by Kim et al. for their tyrosinase inhibition activity. Based on their results, among these flavonoids, mirkoin ($IC_{50} = 5 \mu\text{M}$) revealed stronger tyrosinase inhibition than the positive control, kojic acid and inhibited tyrosinase reversibly in a competitive mode²³². Recently, two isoflavonoids lupinalbin ($IC_{50} = 39.7 \pm 1.5 \mu\text{M}$), and 2'-hydroxygenistein-7-O-gentibioside ($IC_{50} = 50.0 \pm 3.7 \mu\text{M}$) from *Apios americana* were identified as competitive inhibitors, with K_i values of $10.3 \pm 0.8 \mu\text{M}$ and $44.2 \pm 1.7 \mu\text{M}$, respectively²⁶⁶.

Flavanones. Flavanones such as naringenin, hesperetin, eriodictyol and their glycosides (e.g. naringin, hesperidin, and liquiritin) and flavanonols (taxifolin) are mainly found in citrus fruits and the medicinal herbs²⁰⁹. A copper chelator flavanone named hesperetin inhibits tyrosinase reversibly and competitively. Based on the ANS-binding fluorescence analysis, hesperetin disrupted of tyrosinase structure by hydrophobic interactions. In addition, hesperetin chelates a copper ion coordinating with 3 histidine residues (HIS61, HIS85, and HIS259) within the active site pocket of the

enzyme due to docking simulation results²⁶⁷. In another study, Chiari et al. have illustrated tyrosinase inhibitory activity of a 6-isoprenoid-substituted flavanone isolated from *Dalea elegans*²⁶⁸. Also, Steppogenin is a natural flavanone with a strong tyrosinase inhibitory activity ($IC_{50} = 0.98 \pm 0.01 \mu\text{M}$), from *Morus alba* L²⁴⁹. Recently, a new isoprenylated sanggenon-type flavanone, nigrasin K, along with some other analogs including sanggenon M, C and O, chalconoracin, sorocein H and kuwanon J isolated from the twigs of *Morus nigra* have been identified as potent tyrosinase inhibitors by Hu et al.²⁶⁹. Among these natural inhibitors, sanggenon D revealed stronger tyrosinase inhibition than the positive control, kojic acid or arbutin.

Flavanols and flavan-3,4-diols. Flavan-3-ols are the most complex subclass of flavonoids ranging from the simple monomers (+)-catechin and its isomer (–)-epicatechin to the oligomeric and polymeric proanthocyanidins, which are also known as condensed tannins. Flavanols, such as catechin, epicatechin, epi-gallocatechin, epicatechin gallate (ECG), epigallocatechin gallate (EGCG) and proanthocyanidins are widespread in the medicinal herbs and higher plants^{231,270}. *Alphitonia neocaledonica* (Rhamnaceae) is an endemic tree of New Caledonia, which has been identified as an anti-tyrosinase source due to the presence of tannins and gallo catechin²²⁸. Moreover, a catechin compound isolated from the ethanol extract of *Distylium racemosum* branches, with IC_{50} value of $30.2 \mu\text{g/mL}$, showed higher tyrosinase inhibition activity than arbutin as a positive control²⁷¹. Also, a proanthocyanidins from *Clausena lansium* demonstrated potent mushroom tyrosinase inhibition in a mixed competitive manner and illustrated strong inhibition of the melanogenic activity of B16 cells. The IC_{50} values for the monophenolase and diphenolase activities were 23.6 ± 1.2 and $7.0 \pm 0.2 \mu\text{g/mL}$, respectively. Furthermore, from the inhibition mechanism of this compound, it can be concluded that a chelation between the hydroxyl group on the B ring of the proanthocyanidins and dicopper ions of the enzyme has been occurred³⁹.

Another investigation revealed that procyanidin-type proanthocyanidins, purified from cherimoya (*Annona squamosa*) pericarp could powerfully inhibit the activities of monophenolase and diphenolase of tyrosinase, competitively²⁷². In addition, Kim et al. have demonstrated that (+)-catechin-aldehyde polycondensates inhibit the L-tyrosine hydroxylation and L-DOPA oxidation by chelation to the active site of tyrosinase²⁷³. Recently, another tyrosinase inhibitor from this class, condensed tannins (mixtures of procyanidins, propelargonidins, prodelphinidins) and their acyl derivatives (galloyl and p-hydroxybenzoate) from *Longan Bark* indicated the reversible and mixed (competitive is dominant) inhibition of tyrosinase²⁷⁴.

Anthocyanidins. Anthocyanins, including anthocyanidins (e.g. cyanidin, delphinidin, malvidin, peonidin, pelargonidin, etc.) and their glycosides, are widely distributed in the medicinal herbs²¹⁷. It seems that there is a significant relationship between anthocyanin content with anti-human and anti-mushroom tyrosinase activities²⁷⁵.

Curcuminoids. Two phenolic compounds, namely curcumin and desmethoxycurcumin have been isolated from the methanolic extract of the heartwood of *Artocarpus altilis* and showed more potent tyrosinase inhibitory activities than the positive control kojic acid¹⁹⁰. Also, a curcumin included in Chouji and Yakuchi extracts inhibited the enzyme competitively¹⁹². In addition, some synthetic curcumin derivative compounds^{217,276} and its analogs possessing *m*-diphenols and *o*-diphenols have been

investigated as potent inhibitors of mushroom tyrosinase²¹⁶. Based on the results, 4-hydroxyl groups in curcumin analogs containing 4-hydroxyl-substituted phenolic rings with C-2/C-4- or C-3/C-4-dihydroxyl-substituted diphenolic rings make them more active than kojic acid²¹⁷.

Coumarins. In search of tyrosinase inhibitors, the inhibitory effects of several coumarin derivatives (Figure 6)²⁷⁷⁻²⁷⁹ such as 3-aryl and 3-heteroaryl coumarins²⁸⁰, esculetin²⁸¹, coumarinolignoid 8'-epi-cleomiscosin²⁸², umbelliferone and their analogs²⁸³, phenyl coumarins²⁸⁴, hydroxycoumarins^{285,286}, thiophosphonic acid diamides, diazaphosphinanes coumarin derivatives²⁸⁷, cardol-coumarin derivatives²⁸⁸ and coumarin-resveratrol hybrids²⁸⁹, were evaluated on tyrosinase activity.

Interestingly, among hydroxycoumarins, the 3-hydroxycoumarin²⁸⁶ and 7-hydroxycoumarin showed potent activity for the tyrosinase inhibition²⁷⁸, while the 4-hydroxycoumarin is not an inhibitor²⁸⁶. Also, 2-(1-(coumarin-3-yl)-ethylidene)

hydrazinecarbothioamide and 2-(1-(6-chlorocoumarin-3-yl)ethylidene)-hydrazinecarbothioamide demonstrated an irreversible inhibition of tyrosinase²⁷⁷. Recently, in the screening of natural products for the development of cosmetic ingredients, two major compounds, *trans*-N-coumaroyltyramine (IC₅₀ = 40.6 μM) and *cis*-N-coumaroyltyramine (IC₅₀ = 36.4 μM) from *Humulus japonicus* showed potent tyrosinase inhibition²⁹⁰.

Chalcones and dihydrochalcones. Chalcones (butein, phloretin, sappan-chalcone, carthamin, etc.), or 1,3-diphenyl-2-propen-1-ones, are one of the most important classes of flavonoids. Chalcone-containing plants have been used for a long time in traditional medicine²⁰⁹. Based on the reports, some natural and synthetic chalcones and their derivatives are identified as new potent depigmentation agents and tyrosinase inhibitors (Figure 7). So far, natural chalcones isoliquiritigenin (2',4'-trihydroxychalcone) and glabrene from licorice roots²⁸³, 2,4,2',4'-hydroxycalcone and three of its analogs with 3'-substituted resorcinol moieties

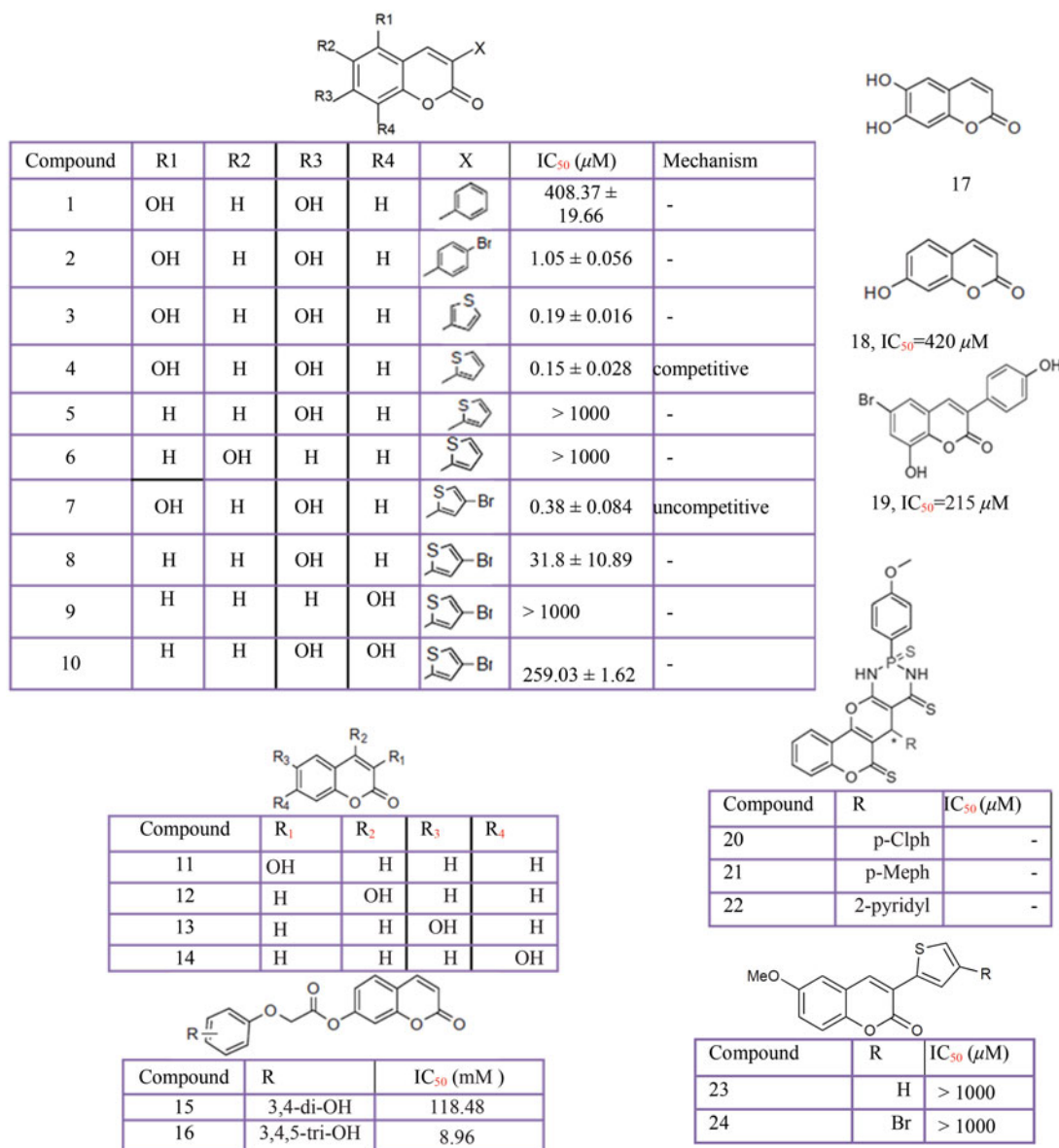
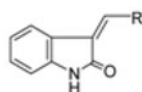
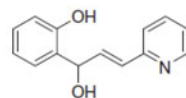


Figure 6. Inhibitory effects of the coumarins derivatives against mushroom tyrosinase activity: 3-aryl and 3-heteroaryl coumarins (1-10, 23-24), 3-hydroxycoumarin (11), 4-hydroxycoumarin (12), 6-hydroxycoumarin (13), 7-hydroxycoumarin (14), umbelliferone analogs (15-16), Esculetin (17) umbelliferone (18), 3-phenyl coumarins with bromo substituent (19), thiophosphonic acid diamides (20-22).

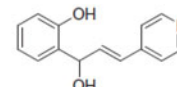


| Compound | R | R ₁ | R ₂ | R ₃ | IC ₅₀ (μM) | | Mechanism |
|----------|---|----------------|------------------|----------------------------------|-----------------------|--------|-------------|
| | | | | | L-Tyrosine | L-DOPA | |
| 1 | | H | H | H | 77.07 | 85.33 | competitive |
| 2 | | H | H | OCH ₃ | 152.88 | 145.17 | competitive |
| 3 | | H | H | N(CH ₃) ₂ | 256.70 | 241.89 | competitive |
| 4 | | H | OCH ₃ | H | 182.46 | 180.25 | competitive |
| 5 | | H | OCH ₃ | OCH ₃ | 63.37 | 59.71 | competitive |
| 6 | | ---- | ---- | ---- | 223.56 | 232.32 | competitive |
| 7 | | ---- | ---- | ---- | 95.98 | 99.10 | competitive |
| 8 | | ---- | ---- | ---- | 110.77 | 107.2 | competitive |

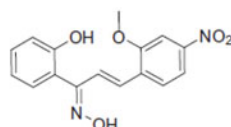
| Compound | R | IC ₅₀ (μM) |
|----------|---|-----------------------|
| | | L-Tyrosine |
| 9 | H | 0.21 |
| 10 | | 0.82 |
| 11 | | 4.62 |
| 12 | | 0.17 |



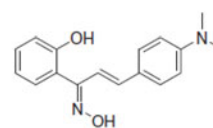
13



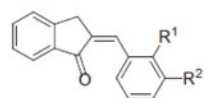
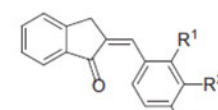
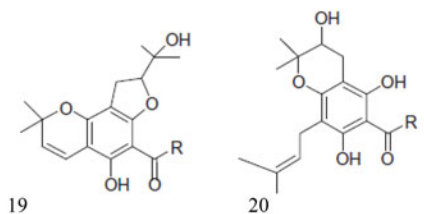
14



15

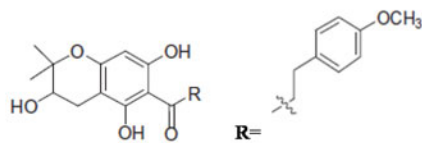


16

17 R¹=OH, R²=H, R³=OH18 R¹=OH, R²=OH, R³=H

19

20



21

R=

22

| Compound | IC ₅₀ (μM) |
|----------|-----------------------|
| | L-Tyrosine |
| 13 | 1.70 |
| 14 | 2.30 |
| 15 | 4.77 |
| 16 | 7.89 |
| 17 | 12.3 |
| 18 | 8.2 |

Figure 7. Tyrosinase Inhibition Activity of chalcone derivatives inhibitors: Oxindole-based chalcone (1–8), chalcones isolated from *Morus australis* (9–12) azachalcones (13–14), oxime based chalcone series (15,16) 2,3-dihydro-1H-inden-1-one chalcone-like derivatives (17,18), Dihydrochalcones from *Flemingia philippinensis* (19–21), chalcone (22).

from *Morus australis* (Figure 6, 19–22)²⁹¹, 2,4,2',4'-tetrahydroxy-3-(3-methyl-2-butenyl)-chalcone from *Morus nigra*²⁹², vulpinoideol B from *Carex vulpinoidea* seeds²⁹³, dihydrochalcones from *Flemingia philippinensis*²¹⁰, 2,4,2',4'-tetrahydroxychalcone (IC₅₀ = 0.07 ± 0.02 μM) and morachalcone A (IC₅₀ = 0.08 ± 0.02 μM) from *Morus alba* L.²⁴⁹ and

bavachinin from *Psoralea corylifolia*²¹ have been presented as tyrosinase inhibitors.

Also, tyrosinase inhibitory effects of several synthetic chalcones and their derivatives were evaluated by various researchers. Oxindole-based chalcones²⁹⁴, 1-(2-cyclohexylmethoxy-

6-hydroxy-phenyl)-3-(4-hydroxymethyl-phenyl) propenone derivative²⁹⁵, isoxazole chalcone derivatives²⁹⁶, some azachalcones and their oximes^{297,298}, 2,4,2',4'-tetrahydroxychalcone and its two derivatives (1,3,5-tris-(2,4-dihydroxy-phenyl) pentane-1,5-dione and 7,2',4'-trihydroxyflavanone)²⁹⁹, 2',4',6'-trihydroxychalcones³⁰⁰, naphthyl chalcones³⁰¹ and chalcone thiosemicarbazide derivatives³⁰² have been identified as a new class of tyrosinase inhibitors. Interestingly, the most important factors in the efficacy of a chalcone are the location of the hydroxyl groups on both aromatic rings and the number of these hydroxyls and the presence of a catechol moiety don't correlate with increasing tyrosinase inhibition potency³⁰³.

Aurones. Okombi et al. have identified Z-benzylidenebenzofuran-3(2H)-one and analogs as human tyrosinase inhibitors. However, they found that aurones are weak inhibitors, but their derivatives with two or three hydroxyl groups preferably at 4,6 and 4' positions make them significant tyrosinase inhibitors. For example, the most potent aurone, 4,6,4'-trihydroxyaurone induces 75% inhibition at 0.1 mM concentration and is highly effective compared to kojic acid³⁰⁴. In addition to synthetic compounds, several natural compounds such as (2'R)-2',3'-dihydro-2'-(1-hydroxy-1-methylethyl)-2,6'-bibenzofuran-6,4'-diol³⁰⁵ and 2-arylbenzofurans isolated from *Morus notabilis*³⁰⁶ and *Morus yunnanensis*²³⁰, benzofuran flavonoids such as mulberrofuran G (MG) and albanol B (AB) isolated from *Morus* sp³⁰⁷ and macourins E isolated from *Morus macroua* (IC₅₀ = 0.39 μM) are potent tyrosinase inhibitors among aurones³⁰⁸.

Phenolic acids

Phenolic acids are divided into hydroxybenzoates and hydroxycinnamates. The most common hydroxycinnamates are *p*-coumaric, caffeic and ferulic acids. So far, *p*-hydroxybenzoic acid, chlorogenic acid (the ester of caffeic acid), vanilic acid (4-hydroxy-3-methoxybenzoic acid) and protocatechuic acid (a dihydroxybenzoic acid) from *Hypericum laricifolium* Juss¹⁵⁴, protocatechualdehyde (IC₅₀ = 0.40 μg/mL) from *Phellinus linteus*¹⁷⁵, benzoic acid propyl gallate³⁰⁹, orsellinic acid (2,4-dihydroxy-6-methylbenzoic acid) and orsellinates (2,4-dihydroxy-6-methyl benzoates)³¹⁰, *p*-coumaric acid from *ginseng* leaves³¹¹, *m*-coumaric acid³¹², *p*-coumarate³¹³ and its derivatives from leaves of *Breynia officinalis*¹⁸⁴ caffeic acid and its *n*-nonyl ester³¹⁴, ferulic acid from *Spiranthes sinensis*²²⁴, 4-Hydroxy cinnamic acid³¹⁵, synthetic hydroxycinnamoyl phenylalanyl/prolyl hydroxamic acid derivatives³¹⁶, and seven hydroxycinnamoyl derivatives in green coffee beans³¹⁷ have been investigated for their tyrosinase inhibition activity. Among these, propyl gallate is a reversible and mixed-type inhibitor on diphenolase activity of tyrosinase with $K_{iS} = 2.135$ mM and $K_j = 0.661$ mM³⁰⁹. Furthermore, *n*-butyl, *iso*-propyl, *sec*-butyl, *n*-pentyl, *n*-hexyl and *n*-octyl orsellinates (uncompetitive, with an inhibition constant of 0.99 mM) behaved as inhibitors at 0.50 mM, whereas methyl, ethyl, *n*-propyl, *tert*-butyl, and *n*-cetyl orsellinates acted as tyrosinase activators. Thus, tyrosinase inhibition increased with chain elongation, suggesting that the enzyme site can accept an eight-carbon alkyl chain³¹⁰.

In addition to these compounds, 3-phenylbenzoic acid (3-PBA) was revealed to be the most potent inhibitor against monophenolase (noncompetitive, IC₅₀ = 6.97 μM) and diphenolase (mixed type inhibition, IC₅₀ = 36.3 μM) activity of mushroom tyrosinase. Also, Oyama et al. have found that some modification such as esterification can abrogate this inhibitory activity of tyrosinase³¹⁸.

Stillbenes

Resveratrol is the most common stilbene. Several stilbenes derivatives from natural and synthetic sources (Figure 8) have been investigated for their tyrosinase inhibition activity including: resveratrol from *Morus alba*³¹⁹, *Pleurotus ferulae*¹³⁵, *vitis viniferae caulis*³²⁰, Carignan grape juice³²¹ *Artocarpus gomezianus*³²² and *Streptomyces avermitilis* MA4680³²³ and also, its derivatives from Dipterocarpaceae plants³²⁴ and synthetic sources³²⁵, oxyresveratrol³²⁶ from *Morus australis*³²⁷, *Morus alba* L (IC₅₀ = 0.10 ± 0.01 μM)²⁴⁹ and *Cudrania cochinchinensis* (IC₅₀ = 2.33 μM)²⁵⁵, azoresveratrol and its derivatives such as (E)-2-((2,4-dihydroxyphenyl) diazenyl) phenyl 4 methylbenzenesulfonate³²⁸ and azo-oxyresveratrol³²⁹, *trans*-resveratrol from *Streptomyces avermitilis* MA4680³¹³, a resveratrol dimer named gnetin C, from melinjo (*Gnetum gnemon*)³³⁰. Also, several hydroxystilbene compounds from synthetic and semisynthetic sources^{331,332} and from the extract of *Veratrum patulum*³³³, along with synthetic glycosides of resveratrol, pterostilbene, and pinostilbene³³⁴, synthetic *trans*-stilbene derivatives³³⁵, azastilbene analogs³³⁶, a newly synthesised stilbene 5-(6-hydroxy-2-naphthyl)-1,2,3-benzenetriol³³⁷, coumarin-resveratrol hybrids²⁹⁰, synthetic polyphenolic deoxybenzoins²¹⁸, hydroxy substituted 2-phenyl-naphthalenes³³⁸ and 4-(6-hydroxy-2-naphthyl)-1,3-benzenediol³³⁹ have been studied for their inhibition activity against tyrosinase. However, based on the enzymatic assays, resveratrol did not inhibit the diphenolase activity of tyrosinase, but L-tyrosine oxidation by tyrosinase was suppressed in presence of 100 μM resveratrol. Interestingly, after the 30 min of preincubation of tyrosinase and resveratrol, both monophenolase and diphenolase activities of tyrosinase were significantly suppressed. Furthermore, this effect was reduced with the addition of L-cysteine, which indicated suicide inhibition mechanism of resveratrol³⁴⁰. Also, oxyresveratrol²⁰¹ is identified as a tyrosinase substrate like hydroquinone, arbutin, caffeic acid and some other inhibitors. In addition to these studies on resveratrol, Fachinetti et al., have demonstrated that the incorporation of resveratrol into nanostructured lipid carriers allowed an enhanced tyrosinase inhibitory activity³⁴¹.

Lignans

Lignans are complex and diverse structures, which are formed from three primary precursors. So far, lignans and lignan glycosides isolated from exocarp of *Castanea henryi*³⁴², *Marrubium velutinum* and *Marrubium cylleneum*³⁴³, *Pinellia ternate*³⁴⁴ and *Crataegus pinnatifida*³⁴⁵ have been evaluated for their tyrosinase inhibitory potentials. However, these compounds mostly displayed a moderate mushroom tyrosinase inhibitory activity.

Terpenoid derivatives

Carvacrol is a monoterpenoid phenol. To date, some carvacrol derivatives³⁴⁶ from synthetic sources, bakuchiol, a terpene phenol from *Psoralea corylifolia*²¹, iridoid glucosides (another type of monoterpenoids) from *Wulfenia carinthiaca* Jacq³⁴⁷ and two new bis-iridoids, namely 7-O-caffeoyl-sylvestroside I and 7-O-(*p*-coumaroyl)-sylvestroside I isolated from *Scabiosa stellata*³⁴⁸ have been investigated for their anti-tyrosinase activities. Among these terpenoid derivatives, Cheng et al. have demonstrated that bakuchiol is a potent inhibitor by applying capillary electrophoresis with reliable online immobilised enzyme microreactor²¹. Also, carvacrol derivatives such as 2-[2-methyl-5-(propan-2-yl)phenoxy]-2-oxoethyl(2E)-3-(2,4-dihydroxyphenyl)prop-2-enoate showed

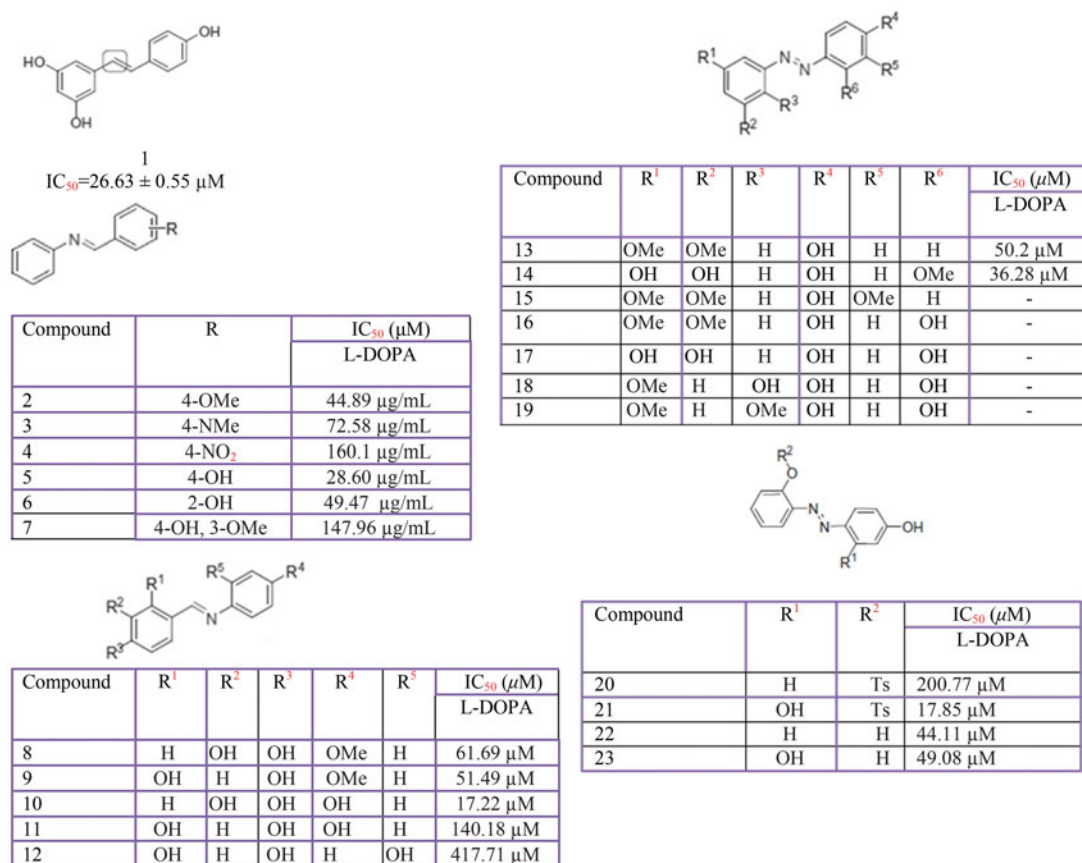


Figure 8. Resveratrol (3,5,4-trihydroxy-trans-stilbene) (1), and its analogs (2–23).

excellent tyrosinase inhibitory activity by a noncompetitive manner with K_i value 0.05 μM and $IC_{50} = 0.0167 \mu M$ ³⁴⁹.

Quinone derivatives

The quinones are a class of small molecules that are mostly derived from aromatic compounds such as benzene or naphthalene. Among these compounds, Aloin, an anthraquinone-C-glycoside from *Aloe vera*³⁴⁹, anthraquinones from *Polygonum cuspidatum*³⁵⁰ and tanshinone IIA ($IC_{50} = 1214 \mu M$) have been verified as tyrosinase inhibitors²³⁹.

Phenyl derivatives

Several biphenyl derivatives³⁵¹ (Figure 9) such as 4,4'-dihydroxybiphenyl³⁵², biphenyl ester derivatives³⁴⁰, biphenyl construction from flavan-3-ol substrates³⁵³, hydroxylated biphenyls²⁶, functionalised bis-biphenyl substituted thiazolidinones³⁶, phenylbenzoic acid derivatives³⁵⁴, phenylethylamide and phenylmethylamide derivatives³⁵⁵, hydroxy substituted 2-phenyl-naphthalenes³¹⁸, 4-hydroxyphenyl beta-D-oligoxylsides³⁵⁶, benzenethiol or phenylthiol³⁵⁷, 2-((1Z)-(2-(2,4-dinitrophenyl)hydrazin-1-ylidene)methyl)phenol³⁵⁸ and 4-[(4-hydroxyphenyl)azo]-benzenesulfonamide³⁵⁹, have been identified as tyrosinase inhibitors.

Pyridine, Piperidine, pyridinone and hydroxypyridinone derivatives

Some hydroxypyridinone derivatives³⁶⁰, 3-hydroxypyridine-4-one derivatives³⁶¹ hydroxypyridinone-L-phenylalanine³⁶² and

pyridinones³⁶³ have been characterised for their antityrosinase activity (Figure 10). Among these inhibitors, one mixed-type inhibitor from hydroxypyridinone-L-phenylalanine conjugates named ((S)-(5-(benzyloxy)-1-octyl-4-oxo-1,4-dihydropyridin-2-yl)methyl 2-amino-3-phenylpropanoate) showed potent inhibitory effect with IC_{50} values of 12.6 and 4.0 μM for monophenolase and diphenolase activities, respectively³⁶².

Thiosemicarbazones, Thiosemicarbazide and other Thio derivatives

Several kinds of thiosemicarbazone derivatives^{38,34,364–376} has been investigated as possible tyrosinase inhibitors (Figure 11). Furthermore, some benzaldehyde derivatives of thiosemicarbazone such as chlorobenzaldehyde thiosemicarbazones³⁶³, *p*-hydroxy and *p*-methoxy benzaldehyde thiosemicarbazone³⁶² along with *p*-methoxybenzaldehyde thiosemicarbazone and 4-dimethylaminobenzaldehyde-thiosemicarbazone and 4-dimethylaminobenzaldehyde-N-phenyl-thiosemicarbazone³⁷⁷ were evaluated for their inhibitory activities on mushroom tyrosinase.

Based on the findings, the appropriate functionalisation of thiosemicarbazone may be improved the inhibitory activity of these inhibitors. Dong et al. believe that the sterically bulky group at the C-4 position of the thiophene ring contributes to this activity. For example, the 4-functionalisation thiophene-2-carbaldehyde thiosemicarbazone with a methoxyacetyl group³⁶⁸ or introducing benzene ring to the 4-functionalised ester group³⁶⁷ enhanced inhibitory activity of thiophene-2-carbaldehyde thiosemicarbazone. However, 5-functionalisation decreased its inhibitory activity. Also, Soares et al., have demonstrated thiosemicarbazones

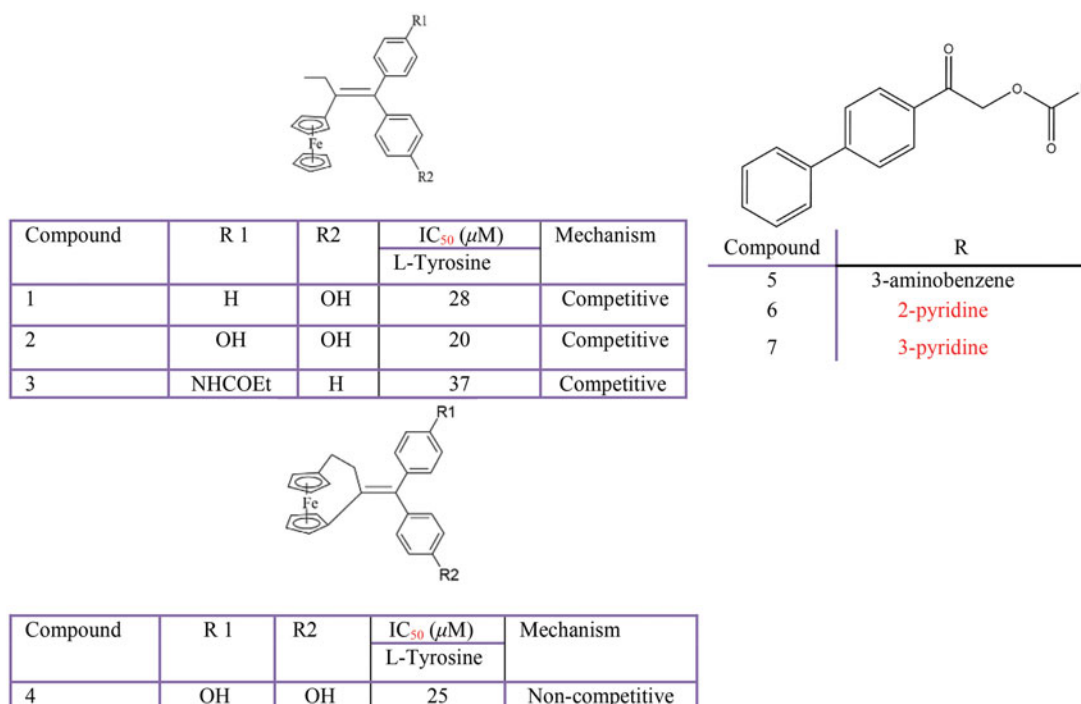


Figure 9. Some phenyl derivatives: aryl butane (1–4), biphenyle ester (5–7).

Thio-1, Thio-2, Thio-3 and Thio-4 substituted with oxygenate moieties, displayed better inhibitory activity (IC₅₀ 0.42, 0.35, 0.36 and 0.44 mM, respectively) than Thio-5, Thio-6, Thio-7 and Thio-8³⁴.

In addition to thiosemicarbazone derivatives, thiosemicarbazide and its derivatives^{378–381}, 5-benzylidene(thio)barbiturate-beta-D-glycosides³⁸², *n*-alkyl³⁸³, *p*-phenylene-bis, phenyl³⁸⁴, benzyl, *p*-xylylene-bis and *p*-pyridine dithiocarbamate sodium salts³⁸⁵, diethyldithiocarbamate, phenylthiourea³⁸⁶ and other thiourea derivatives (Figure 12) such as methimazole, thiouracil, methylthiouracil, propylthiouracil, ambazone, and thioacetazone³⁸⁷ have been identified as tyrosinase inhibitors.

Azole and thiazolidine derivatives

So far, several azole derivatives (Figure 13) have been studied for their tyrosinase inhibitory activity³⁸⁸. The discovered new types of inhibitors included DL-3(5-benzazoly) alanines and alpha-methyl-dopa analogs³⁸⁹, aryl pyrazoles³⁹⁰, heterocyclic hybrids based on pyrazole and thiazolidinone scaffolds³⁹¹, 3,5-diaryl-4,5-dihydro-1H³⁹² and 3,5-diaryl pyrazole derivatives³⁹³, pyrazolo[4,3-*e*] [1,2,4]triazine sulfonamides and sildenafil^{394–396}, 1,3-oxazine-tetrazole³⁹⁷, indole-spliced thiadiazole³⁹⁸, benzimidazole-1,2,3-triazole hybrids³⁹⁹, 1,2,3-triazole-linked coumarinopyrazole conjugates⁴⁰⁰, isoxazolone derivatives⁴⁰¹ 5(4H)-oxazolone derivative⁴⁰², imidazolium ionic liquids⁴⁰³, thiazolyl resorcinols⁴⁰⁴ have demonstrated the inhibitory effect on tyrosinase. Furthermore, some thiazolidine derivatives have been evaluated for their tyrosinase inhibitory activity including azo-hydrazone tautomeric dyes substituted by thiazolidinone moiety⁴⁰⁵, (Z)-5-(2,4-dihydroxybenzylidene) thiazolidine-2,4-dione⁴⁰⁶, 5-(substituted benzylidene) thiazolidine-2,4-dione derivatives⁴⁰⁷, (2R,4R)-2-(2,4-dihydroxyphenyl)thiazolidine-4-carboxylic acid⁴⁰⁸, 2-(substituted phenyl) thiazolidine-4-carboxylic acid derivatives⁴⁰⁹ and (Z)-5-(3-hydroxy-4-methoxybenzylidene)-2-iminothiazolidin-4-one⁴¹⁰.

Kojic acid analogs

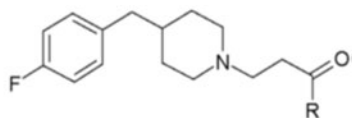
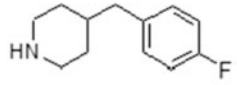
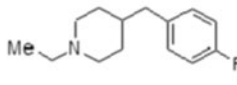
Kojic acid is a well-known tyrosinase inhibitor. When DL-DOPA, norepinephrine and dopamine are oxidised by tyrosinase, Kojic acid inhibits effectively the rate of formation of pigmented product(s) and of oxygen uptake⁴¹¹. Furthermore, several of its derivatives have demonstrated a potent tyrosinase inhibitory activity^{361,412–418}. Noh et al. have modified kojic acid with amino acids and screened their tyrosinase inhibitory activity. Among them, kojic acid-phenylalanine amide showed a strong non-competitive inhibition⁴¹⁷. Interestingly, some kojic acid derivatives despite their depigmenting activities did not display tyrosinase inhibitory activity⁴¹⁹.

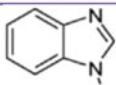
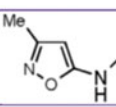
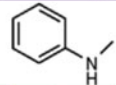
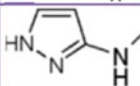
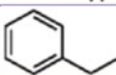
Recently, Xie et al. have reported a kojic acid analog namely 5-phenyl-3-[5-hydroxy-4-pyrone-2-yl-methylmercapto]-4-(2,4-dihydroxybenzylamino)-1,2,4-triazol as a potent competitive tyrosinase inhibitor with an IC₅₀ value of 1.35 ± 2.15 μM⁴¹². Tyrosinase inhibitory activity of some kojic acid derivatives is shown in Figure 14.

Benzaldehyde derivatives

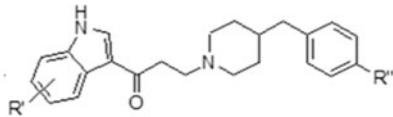
Benzaldehyde⁴²⁰ and its derivatives⁴²¹, hydroxy- or methoxy-substituted benzaldoximes and benzaldehyde-O-alkyloximes⁴²², piperonal or 4-(methylenedioxy) benzaldehyde mesoionic derivatives⁴²³, 4-hydroxybenzaldehyde derivatives⁴²⁴, anisaldehyde⁴²⁵ have been investigated for their inhibitory activities against tyrosinase (Figure 15).

Among these derivatives, 3,4-dihydroxybenzaldehyde-O-ethylloxime (IC₅₀ = 0.3 ± 0.1 μM) is of the same magnitude as one of the best tyrosinase known inhibitors tropolone (IC₅₀ = 0.13 ± 0.08 μM)⁴²². However, in benzaldehyde derivatives, the presence of the aldehyde group and the terminal methoxy group in C4 was found to play an important role in its inhibitory effect. But, due to their lower activity levels or serious side effects, unfortunately, most 4-substituted benzaldehyde derivatives cannot be considered for practical use⁴²¹.

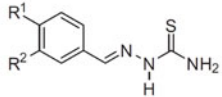
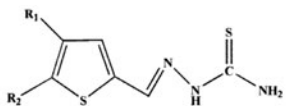
| Compound | R | IC ₅₀ (μM) | |
|----------|---|-----------------------|----------------|
| | | L-Tyrosine | L-DOPA |
| 1 |  | 113.34 ± 16.37 | 159.24 ± 18.65 |
| 2 |  | 28.55 ± 0.75 | 110.95 ± 18.55 |
| 3 |  | 28.85 ± 0.95 | 27.90 ± 0.99 |
| 4 |  | 5.87 ± 1.59 | 39.58 ± 9.61 |
| 5 |  | 13.98 ± 0.81 | 83 ± 6.54 |

| Compound | IC ₅₀ (μM) | |
|----------|-----------------------|----------------|
| | L-Tyrosine | L-DOPA |
| 14 | 43.70 ± 2.61 | 286.83 ± 10.52 |
| 15 | 32.17 ± 4.19 | 116.0 ± 17.69 |



| Compound | R' | R'' | IC ₅₀ (μM) | | Mechanism |
|----------|--------------------------------------|-----|-----------------------|----------------|-----------------|
| | | | L-Tyrosine | L-DOPA | |
| 6 | H | H | 175.00 ± 2.71 | 252.00 ± 3.60 | ---- |
| 7 | 5-OCH ₃ | H | 25.78 ± 1.80 | 197.45 ± 10.62 | ---- |
| 8 | 6-OCH ₃ | H | 67.25 ± 5.92 | 168.55 ± 14.01 | ---- |
| 9 | 5,6-(OCH ₃) ₂ | H | 57.24 ± 1.93 | 133.00 ± 2.32 | ---- |
| 10 | H | F | 9.45 ± 0.36 | 8.82 ± 0.38 | non-competitive |
| 11 | 5-OCH ₃ | F | 7.86 ± 3.02 | 14.15 ± 1.60 | non-competitive |
| 12 | 6-OCH ₃ | F | 9.40 ± 0.63 | 18.03 ± 0.91 | noncompetitive |
| 13 | 5,6-(OCH ₃) ₂ | F | 5.11 ± 0.36 | 7.56 ± 1.90 | mixed-type |

Figure 10. Inhibitory effects of some piperidine derivatives on mushroom tyrosinase activity. 4-(4-fluorobenzyl) piperidine derivatives (1–5) indole derivatives (6–13) amine (14) and N-ethyl (15).

| | R ¹ | R ² | IC ₅₀ (mM) |
|------------------|--------------------------------|------------------|-----------------------|
| 1 | HO | H | 0.42 |
| 2 | HO | OCH ₃ | 0.35 |
| 3 | OC ₂ H ₅ | H | 0.36 |
| 4 | OCH ₃ | H | 0.44 |
| 5 | H | H | 0.82 |
| 6 | Br | H | 1.42 |
| 7 | Cl | H | 0.84 |
| 8 | Pyridine 2 Carboxaldehyde | | 0.19 |
| Ascorbic acid | – | – | 0.26 |

| | R ₁ | R ₂ | IC ₅₀ (μM) |
|----|---|---|--------------------------|
| 9 | CH ₂ OH | H | 0.67 |
| 10 | CH ₂ OCH ₃ | H | 0.74 |
| 11 | CH ₂ OCH ₂ CH ₃ | H | 0.49 |
| 12 | CH ₂ OCH ₂ CH ₂ OH | H | 1.42 |
| 13 | H | CH ₂ OH | 3.21 |
| 14 | CH ₂ OCOCH ₃ | H | 0.34 |
| 15 | H | CH ₂ OCH ₂ CH ₂ OH | 1.77 |
| 16 | H | CH ₂ OCH ₃ | 3.26 |
| 17 | H | CH ₂ OCOCH ₃ | 1.71 |
| 18 | H | CH ₃ | 2.59 |
| 19 | H | NO ₂ | 3.53 |
| 20 | H | Ph | 0.67 |

Figure 11. Inhibitory effects of some thiosemicarbazone derivatives on the tyrosinase monophenolase activity.

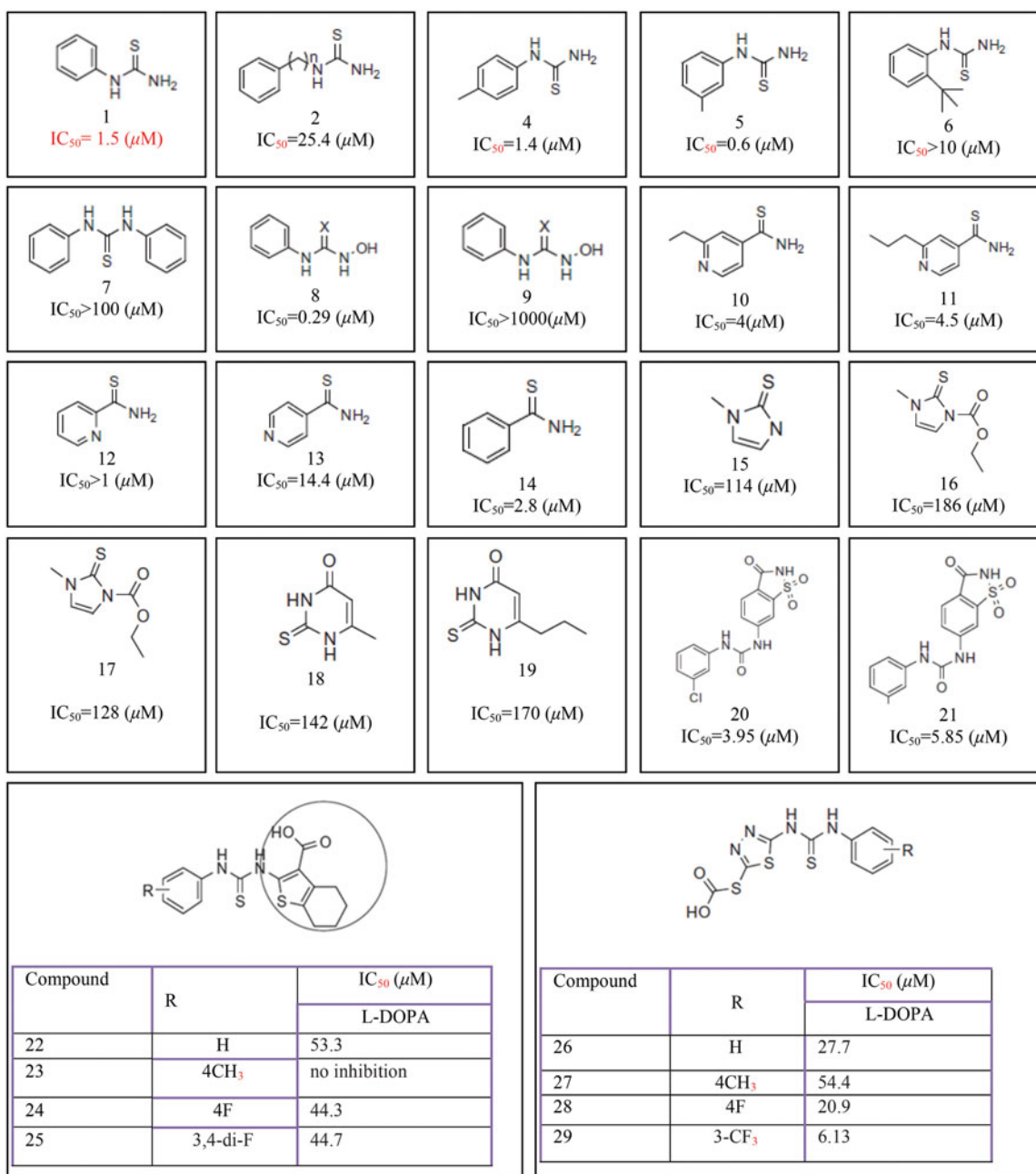


Figure 12. Thiourea derivatives (1–14), methimazole (15), carbimazole (16), thioracil (17), methylthioracil (18), propylthioracil (19), 6-(3-chlorophenylurenyl) saccharin (20), 6-(3-iodophenylthiourenyl) saccharin (21), 4,5,6,7-tetrahydro- 2-[[[(phenylamino)thioxomethyl]amino]-benzo[b]thiophene-3-carboxylic acid derivatives (22–25), 2-(1,3,4-thiadiazol-2-yl) thio acetic acid derivatives (26–29).

Carboxylic acids

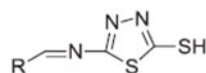
Inhibitory effects of pyruvic acid, acrylic acid, propanoic acid, 2-oxo-butanoic acid, and 2-oxo-octanoic acid¹²⁴, (S)- and (R)-6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acids⁴²⁶ have been investigated on tyrosinase activity.

Based on the findings investigated by Gheibi et al., aliphatic carboxylic acids have dual effects on the monophenolase and diphenolase activities of mushroom tyrosinase. They have found that optimal diphenolase activity of tyrosinase takes place in the presence of *n*-alkyl acids (pyruvic acid, acrylic acid, propanoic acid, 2-oxo-butanoic acid, and 2-oxo-octanoic acid). While, the monophenolase activity is inhibited by all types of *n*-alkyl acids. They have believed that there is a physical difference in the docking of *mono*- and *o*-diphenols to the tyrosinase active site. On the other hand, the binding of acids occurs through their carboxylate group

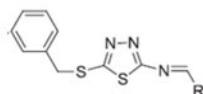
with one copper ion of the binuclear site. So these carboxylic acid compounds completely block the monophenolase reaction, by preventing monophenol binding to the oxyform of the enzyme¹²⁴.

Xanthate derivative

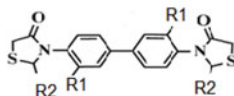
The inhibitory effect of some synthesised xanthates including C₃H₇OCS₂Na, C₄H₉OCS₂Na, C₅H₁₁OCS₂Na, C₂H₅OCS₂Na, and C₆H₁₃OCS₂Na have been examined for inhibition of both monophenolase and diphenolase activities of mushroom tyrosinase. Based on the reports, C₃H₇OCS₂Na and C₄H₉OCS₂Na showed a mixed inhibition pattern on monophenolase activity but C₅H₁₁OCS₂Na and C₆H₁₃OCS₂Na showed a competitive and C₂H₅OCS₂Na showed uncompetitive inhibition pattern. For diphenolase activity, C₃H₇OCS₂Na and C₂H₅OCS₂Na showed mixed



| Compound | R | IC ₅₀ (μM) | Mechanism |
|----------|-------------------------------|-----------------------|---------------|
| | | L-DOPA | |
| 1 | Phenyl | 49.246 ± 0.184 | ---- |
| 2 | 2-Fluorophenyl | 0.765 ± 0.077 | ---- |
| 3 | 3-Chlorophenyl | 4.753 ± 0.109 | ---- |
| 4 | 4-Chlorophenyl | 2.460 ± 0.155 | ---- |
| 5 | 3,4-Dichlorophenyl | 3.621 ± 0.142 | ---- |
| 6 | 5-Chloro-2-hydroxyphenyl | 1.718 ± 0.135 | ---- |
| 7 | 5-Bromo-2-hydroxyphenyl | 1.971 ± 0.131 | ---- |
| 8 | 4-Bu-phenyl | 1.390 ± 0.118 | ---- |
| 9 | 2-Nitrophenyl | 0.671 ± 0.093 | irreversible |
| 10 | 2-Methoxyphenyl | 0.733 ± 0.083 | ---- |
| 11 | 3,5-Dimethoxyphenyl | 5.353 ± 0.136 | ---- |
| 12 | 2-Hydroxyphenyl | 0.785 ± 0.064 | uncompetitive |
| 13 | 2,4-Dihydroxyphenyl | 0.255 ± 0.065 | uncompetitive |
| 14 | 4-Hydroxy-3-methoxyphenyl | 0.036 ± 0.002 | ---- |
| 15 | 3,5-Dimethoxy-4-hydroxyphenyl | 0.473 ± 0.096 | ---- |
| 16 | 2,4,6-Trimethoxyphenyl | 0.907 ± 0.142 | ---- |



| Compound | R | IC ₅₀ (μM) |
|----------|--------------------------|-----------------------|
| | | L-DOPA |
| 17 | 5-Chloro-2-hydroxyphenyl | 247.350 ± 3.782 |




| Compound | R1 | R2 | IC ₅₀ (μM) |
|----------|-----------------|----|-----------------------|
| | | | L-DOPA |
| 18 | H | | 21.61±0.11 |
| 19 | H | | |
| 20 | H | | 0.61±0.05 |
| 21 | H | | 2.41±0.32 |
| 22 | H | | 4.41±0.11 |
| 23 | H | | 342.52±0.17 |
| 24 | CH ₃ | | 7.71±0.12 |
| 25 | CH ₃ | | <500 |
| 26 | CH ₃ | | |
| 27 | CH ₃ | | 2.81±0.06 |
| 28 | CH ₃ | | |
| 29 | CH ₃ | | |

Figure 13. Thiaziazole derivatives: 1,3,4-thiadiazole derivatives (1–17) and thiazolidinones derivative (18–29).

inhibition but $C_4H_9OCS_2Na$ and $C_5H_{11}OCS_2Na$ and $C_6H_{13}OCS_2Na$ showed competitive inhibition⁴²⁷. According to their results, it seems that the lengthening of the hydrophobic tail of the xanthates leads to a decrease of the K_i values for monophenolase inhibition and an increase of the K_i values for diphenolase inhibition⁴²⁸.

Other tyrosinase inhibitors

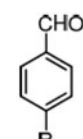
Except the inhibitors listed above, other compounds have also been registered for their tyrosinase inhibitory activity by different researchers such as: two Keggin-type polyoxometalates containing



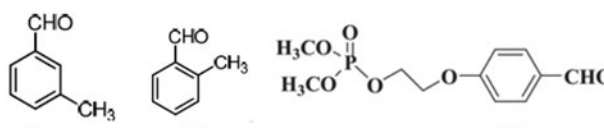
| Compound | R ₁ | R ₂ | IC ₅₀ (μM) | Mechanism |
|----------|---------------------------------|----------------|-----------------------|-------------|
| | | | L-DOPA | |
| 1 | CH ₃ | 2,4-di-OH | 5.80 ± 0.28 | ---- |
| 2 | CH ₃ | 2 OH | 10.20 ± 0.56 | ---- |
| 3 | CH ₂ CH ₃ | 2 OH | 7.60 ± 1.50 | ---- |
| 4 | H | 2 OH | 5.85 ± 0.60 | ---- |
| 5 | CH ₃ | 3-OH | 12.5 ± 1.08 | ---- |
| 6 | Ph | 2,4-di-OH | 1.35 ± 2.15 | competitive |
| 7 | 4-CH ₃ Ph | 2 OH | 3.80 ± 0.78 | ---- |
| 8 | 3-ClPh | 2 OH | 2.50 ± 0.89 | ---- |
| 9 | 4-ClPh | 2 OH | 1.71 ± 1.56 | ---- |
| 10 | Ph | 2 OH | 1.50 ± 1.25 | ---- |

| Compound | R ₁ | R ₂ | IC ₅₀ (μM) |
|------------|----------------|--------------------------|-----------------------|
| | | | L-DOPA |
| 11 | 2-ClPh | 3,4-di-OH | 5.20 ± 0.69 |
| 12 | H | 4-OH | 8.54 ± 2.38 |
| 13 | H | 2,4-di-OH | 9.60 ± 1.78 |
| 14 | H | 3-CH ₃ O-4-OH | 17.50 ± 2.75 |
| Kojic acid | | | 20.00 ± 1.08 |

Figure 14. Some kojic acid analogs: hydroxybenzaldehydebased kojic acid analogs (5-substituted-3-[5-hydroxy-4-pyrone-2-ylmethylmercapto]-4-arylmethylamino-1,2,4-triazole (1–10) and 5-substituted-3-[5-hydroxy-4-pyrone-2-yl-methylmercapto]-4-arylmethyleneamino-1,2,4-triazole (11–14).



| Compound | R | IC ₅₀ (μM) | Mechanism |
|----------|---|-----------------------|------------------------|
| | | L-DOPA | |
| 1 | H | 31.0 | Partial Noncompetitive |
| 2 | CH ₃ | 70.5 | Partial Noncompetitive |
| 3 | CH(CH ₃) ₂ | 376.7 | Partial Noncompetitive |
| 4 | OCH ₃ | 1200.9 | Partial Noncompetitive |
| 5 | O(CH ₂) ₂ CH ₃ | ---- | ---- |
| 6 | O(CH ₂) ₃ CH ₃ | ---- | ---- |
| 7 | CH ₂ (CH ₂) ₂ CH ₃ | 318.0 | Full Mixed |
| 8 | CH ₂ (CH ₂) ₃ CH ₃ | 112.0 | Full Mixed |
| 9 | C(CH ₃) ₃ | 455.7 | Full Mixed |
| 10 | CH ₂ CH ₃ | 133.4 | Partial Noncompetitive |
| 11 | C ₆ H ₅ | ---- | ---- |
| 12 | OCH ₂ CH ₃ | 3604.2 | Full Noncompetitive |



| Compound | IC ₅₀ (μM) | Mechanism |
|----------|-----------------------|------------------------|
| | L-DOPA | |
| 13 | 287.8 | Partial Noncompetitive |
| 14 | 940.5 | Partial Mixed |
| 15 | 59 | Noncompetitive |

Figure 15. Benzaldehyde derivatives: 4-substituted benzaldehyde (1–15).

glycine as potent inorganic reversible inhibitors⁴²⁹, cadmium ions with an IC₅₀ of 2.92 ± 0.16 mM⁴⁸ and rifampicin with an IC₅₀ = 90 ± 0.6 μM⁹ as reversible and noncompetitive inhibitors, ammonium tetrathiotungstate⁴³⁰, amoxicillin (IC₅₀ = 9.0 ± 1.8 mM)⁴³¹, mallotophilippen A and B⁴³² α-naphthol and β-naphthol⁴³³, red koji extracts (IC₅₀ of 5.57 mg/mL)⁴³⁴ and

alpha-hydrazinophloretic acid⁴³⁵ as competitive inhibitors and rottlerin as a mixed inhibitor⁴³². Furthermore, *n*-alkyl sulfates⁴³⁶, sericin extracted from tasar silk fiber waste⁴³⁷, 2-hydroxy-3-methylcyclopent-2-ene (IC₅₀ = 721.91 μg mL⁻¹) isolated from ribose-histidine Maillard reaction products⁴³⁸, three natural compounds from safflower⁴³⁹ and mimosine³⁸⁶ and ethylenediamine⁴⁴⁰ are other kinds of tyrosinase inhibitors.

Synergistic effects of tyrosinase inhibitors

Synergistic strategy for tyrosinase inhibitors is a useful strategy for the improvement of their inhibitory activities. Based on the findings, the mixtures of glabridin:resveratrol, glabridin:oxyresveratrol, resveratrol:oxyresveratrol, phenylethylresorcinol:resveratrol⁴⁴¹, oxyresveratrol:dioscin⁴⁴², aloesin:arbutin⁴⁴³, 4-methyl catechol:catechol⁴⁴⁴, 3-(2,4-dihydroxyphenyl)propionic acid:l-ascorbic acid⁴⁴⁵, dihydromyricetin:vitamin D3³⁷, linderanolide B combined with arbutin, 1-phenyl-2-thiourea or kojic acid⁴⁴⁶, have shown synergistic effect on tyrosinase. These studies may provide a scientific strategy for screening effective tyrosinase inhibitors.

Conclusion

Due to the vital role of tyrosinase in the enzymatic browning of food and depigmentation disorders in humans, its inhibitors have been considered by researchers, extensively. As mentioned above, natural sources such as plants and microorganisms and their effective compounds have wonderful potential as organic anti-tyrosinase sources.

However, the majority of the compounds identified from natural sources were isolated from plants but, recently, microorganisms are considered as potential sources of tyrosinase inhibitors. It is interesting that despite the diversity of natural inhibitors, a large number of tyrosinase inhibitors are phenolic-based structures. Many researchers have designed appropriate scaffold inspired by the structure of natural compounds and developed novel synthetic inhibitors. In this paper, many natural, semi-synthetic and synthetic inhibitors have been summarised and the inhibitory effects of these compounds on the tyrosinase activity are discussed.

Based on the results, phenolic compounds (simple phenols and polyphenols) and their derivatives and several compounds including terpenoid, phenyl, pyridine, piperidine, pyridinone, hydroxypyridinone, thiosemicarbazone, thiosemicarbazide, azole, thiazolidine, kojic acid, benzaldehyde and xanthate derivatives were characterised as potent tyrosinase inhibitors. The appropriate functionalisation of these inhibitors such as C-6 and C-7 hydroxyl groups of the isoflavone skeleton, 4-functionalisation thiophene-2-carbaldehyde thiosemicarbazone with a methoxyacetyl group and the aldehyde group and methoxy group in C4 of benzaldehyde derivatives may be improved the inhibitory activity of these inhibitors. Furthermore, in chalcone derivatives, the location of the hydroxyl groups on both aromatic rings and the number of hydroxyls is an important factor in the efficacy of a chalcone. In contrast, some modifications such as the prenylation or the vinylation of some flavonoid molecules do not enhance their tyrosinase inhibitory activity while deglycosylation of some flavonoid glycosides by far-infrared irradiation can be improved tyrosinase inhibitory activity. Interestingly, among different inhibitors, some compounds, especially hydroquinone and its known derivatives (α and β -arbutin), are described as both a tyrosinase inhibitor and a substrate.

Actually, the main objective of this review is to provide a useful source of effective tyrosinase inhibitors. However, despite the existence of a wide range of tyrosinase inhibitors from natural and synthetic sources, only a few of them, in addition to being effective, are known as safe compounds. Therefore, it is recommended to examine the efficacy and safety of inhibitors by *in vivo* models, along with *in vitro* and docking experiments, especially for the application of such materials in food and medicinal products. Finally, we hope that the information provided in this study, which is the result of numerous researchers' efforts, could serve as leads in the search for effective anti-tyrosinase agents from natural and synthetic sources with increased efficiency and safety in the food and cosmetics industries.

Disclosure statement

The authors report that they have no conflicts of interest.

Funding

This work was financially supported by Research Council of both University of Tehran and IAU Jahrom Branch.

References

1. Dembitsky VM, Kilimnik A. Anti-melanoma agents derived from fungal species. *M J Pharma* 2016;1:1–16.
2. Maghsoudi S, Adibi H, Hamzeh M, et al. Kinetic of mushroom tyrosinase inhibition by benzaldehyde derivatives. *J Rep Pharma Sci* 2013;2:156–64.
3. Halaoui S, Asther M, Kruus K, et al. Characterization of a new tyrosinase from *Pycnoporus* species with high potential for food technological applications. *J Appl Microbiol* 2005;98:332–43.
4. Sahu RK, Roy A, Dwivedi J, Jha AK. Promotion and computation of inhibitory effect on tyrosinase activity of herbal cream by incorporating indigenous medicinal plants. *Pak J Biol Sci* 2014;17:146–50.
5. Jeon SH, Jong-Uk HK, Kwang-Hoon K. Inhibitory effects on L-dopa oxidation of tyrosinase by skin-whitening agents. *Bull Korean Chem Soc* 2005;26:1135–7.
6. Garcia-Molina F, Munoz JL, Varon R, et al. A review on spectrophotometric methods for measuring the monophenolase and diphenolase activities of tyrosinase. *J Agric Food Chem* 2007;55:9739–49.
7. Ravani Ananda R, Nagaraja P. Quantification of 4-amino-5-hydroxynaphthalene-2,7-disulfonic acid mono sodium salt by oxidation with tyrosinase in the presence of 3-methyl-2-benzothiazolinone hydrazine. *Chem Sci Rev Lett* 2015;4: 342–8.
8. Winder AJ. A stopped spectrophotometric assay for the dopa oxidase activity of tyrosinase. *J Biochem Biophys Methods* 1994;28:173–83.
9. Chai WM, Lin MZ, Song FJ, et al. Rifampicin as a novel tyrosinase inhibitor: inhibitory activity and mechanism. *Int J Biol Macromol* 2017;102:425–30.
10. Lee SY, Baek N, Nam TG. Natural, semisynthetic and synthetic tyrosinase inhibitors. *J Enzyme Inhib Med Chem* 2016; 31:1–13.
11. Zhou J, Tang Q, Wu T, Cheng Z. Improved TLC bioautographic assay for qualitative and quantitative

- estimation of tyrosinase inhibitors in natural products. *Phytochem Anal* 2017;28:115–24.
12. García P, Ramallo IA, Furlan RLE. Reverse phase compatible TLC-bioautography for detection of tyrosinase inhibitors. *Phytochem Anal* 2017;28:101–5.
 13. García P, Furlan RL. Multiresponse optimisation applied to the development of a TLC autography for the detection of tyrosinase inhibitors. *Phytochem Anal* 2015;26:287–92.
 14. Wangthong S, Tonsiripakdee I, Monhaphol T, et al. Post TLC developing technique for tyrosinase inhibitor detection. *Biomed Chromatogr* 2007;21:94–100.
 15. Taibon JAA, Schwaiger S, Magnenat C, et al. Prevention of false-positive results: development of an HPTLC autographic assay for the detection of natural tyrosinase inhibitors. *Planta Med* 2015;81:1198–204.
 16. Misra BB, Dey S. TLC-bioautographic evaluation of in vitro anti-tyrosinase and anti-cholinesterase potentials of sandalwood oil. *Nat Prod Commun* 2013;8:253–6.
 17. Kamagaju L, Morandini R, Bizuru E, et al. Tyrosinase modulation by five Rwandese herbal medicines traditionally used for skin treatment. *J Ethnopharmacol* 2013;146:824–34.
 18. Liu DM, Yang JL, Ha W, et al. Kinetics and inhibition study of tyrosinase by pressure mediated microanalysis. *Anal Biochem* 2017;525:54–9.
 19. Tang L, Zhang W, Zhao H, Chen Z. Tyrosinase inhibitor screening in traditional Chinese medicines by electrophoretically mediated microanalysis. *J Sep Sci* 2015;38:2887–92.
 20. Jiang TF, Liang TT, Wang YH, et al. Immobilized capillary tyrosinase microreactor for inhibitor screening in natural extracts by capillary electrophoresis. *J Pharm Biomed Anal* 2013;84:36–40.
 21. Cheng M, Chen Z. Screening of tyrosinase inhibitors by capillary electrophoresis with immobilized enzyme microreactor and molecular docking. *Electrophoresis* 2017;38:486–93.
 22. Sun BB, Qi L, Mu XY, et al. A Chiral ligand exchange CE system for monitoring inhibitory effect of kojic acid on tyrosinase. *Talanta* 2013;116:1121–5.
 23. Winder AJ, Harris H. New assays for the tyrosine hydroxylase and dopa oxidase activities of tyrosinase. *Eur J Biochem* 1991;198:317–26.
 24. Chen YM, Chavin W. Radiometric assay of tyrosinase and theoretical considerations of melanin formation. *Anal Biochem* 1965;13:234–58.
 25. Vandeput M, Patris S, Silva H, et al. Application of a tyrosinase microreactor – detector in a flow injection configuration for the determination of affinity and dynamics of inhibitor binding. *Sens Actuators B Chem* 2017;248:385–94.
 26. Ruzza P, Serra PA, Davide Fabbri D, et al. Hydroxylated biphenyls as tyrosinase inhibitor: a spectrophotometric and electrochemical study. *Eur J Med Chem* 2017;126:1034–8.
 27. Tang H, Cui F, Li H, et al. Understanding the inhibitory mechanism of tea polyphenols against tyrosinase using fluorescence spectroscopy, cyclic voltammetry, oximetry, and molecular simulations. *RSC Adv* 2018;8:8310–8.
 28. Liu Z, Liu S. A novel fluorescent biosensor for adrenaline detection and tyrosinase inhibitor screening. *Anal Bioanal Chem* 2018;410:4145–52.
 29. Hsu KD, Chan YH, Chen HJ, et al. Tyrosinase-based TLC autography for anti-melanogenic drug screening. *Sci Rep* 2018;8:401.
 30. Bagherzadeh K, Shirgahi Talari F, Sharifi A, et al. A new insight into mushroom tyrosinase inhibitors: docking, pharmacophore-based virtual screening, and molecular modeling studies. *J Biomol Struct Dyn* 2015;33:487–501.
 31. Tang H, Cui F, Liu L, Li Y. Predictive models for tyrosinase inhibitors: challenges from heterogeneous activity data determined by different experimental protocols. *Comput Biol Chem* 2018;73:79–84.
 32. Li Q, Yang H, Mo J, et al. Identification by shape-based virtual screening and evaluation of new tyrosinase inhibitors. *Peer J* 2018;6:e4206.
 33. Suthar SK, Bansal S, Narkhede N, et al. Design, synthesis and biological evaluation of oxindole-based chalcones as small-molecule inhibitors of melanogenic tyrosinase. *Chem Pharm Bull* 2017;65:833–9.
 34. Soares MA, Almeida MA, Marins-Goulart C, et al. Thiosemicarbazones as inhibitors of tyrosinase enzyme. *Bioorg Med Chem Lett* 2017;27:3546–50.
 35. Gou L, Lee J, Hao H, et al. The effect of oxaloacetic acid on tyrosinase activity and structure: Integration of inhibition kinetics with docking simulation. *Int J Biol Macromol* 2017;101:59–66.
 36. Mutahir S, Khan MA, Khan IU, et al. Organocatalyzed and mechanochemical solvent-free synthesis of novel and functionalized bis-biphenyl substituted thiazolidinones as potent tyrosinase inhibitors: SAR and molecular modeling studies. *Eur J Med Chem* 2017;134:406–14.
 37. Fan M, Zhang G, Pan J, Gong D. An inhibition mechanism of dihydromyricetin on tyrosinase and the joint effects of vitamins B6, D3 or E. *Food Funct* 2017;8:2601–10.
 38. Liu J, Li M, Yu Y, Cao S. Novel inhibitors of tyrosinase produced by the 4-substitution of TCT (pi). *Int J Biol Macromol* 2017;103:1096–106.
 39. Chai WM, Lin MZ, Feng HL, et al. Proanthocyanidins purified from fruit pericarp of *Clausena lansium* (Lour.) Skeels as efficient tyrosinase inhibitors: structure evaluation, inhibitory activity and molecular mechanism. *Food Funct* 2017;8:1043–51.
 40. Kwong HC, Chidan Kumar CS, Mah SH, et al. Novel biphenyl ester derivatives as tyrosinase inhibitors: synthesis, crystallographic, spectral analysis and molecular docking studies. *PLoS one* 2017;12:e0170117.
 41. Garcia-Jimenez A, Teruel-Puche JA, Ortiz-Ruiz CV, et al. Study of the inhibition of 3-/4-aminoacetophenones on tyrosinase. *Reac Kinet Mech Cat* 2017;120:1–13.
 42. Cui Y, Hu YH, Yu F, et al. Inhibition kinetics and molecular simulation of *p*-substituted cinnamic acid derivatives on tyrosinase. *Int J Biol Macromol* 2017;95:1289–97.
 43. Ferro S, De Luca L, Germano MP, et al. Chemical exploration of 4-(4-fluorobenzyl)piperidine fragment for the development of new tyrosinase inhibitors. *Eur J Med Chem* 2017;125:992–1001.
 44. Tang J, Liu J, Wu F. Molecular docking studies and biological evaluation of 1,3,4-thiadiazole derivatives bearing schiff base moieties as tyrosinase inhibitors. *Bioorg Chem* 2016;69:29–36.
 45. Lall N, Mogapi E, de Canha MN, et al. Insights into tyrosinase inhibition by compounds isolated from *Greyia radlkoferi* Szyszyl using biological activity, molecular docking and gene expression analysis. *Bioorg Med Chem* 2016;24:5953–9.
 46. Hassani S, Haghbeen K, Fazli M. Non-specific binding sites help to explain mixed inhibition in mushroom tyrosinase activities. *Eur J Med Chem* 2016;122:138–48.

47. Wang R, Chai WM, Yang Q, et al. (4-Fluorophenyl)-quinazolin-4(3H)-one as a novel tyrosinase inhibitor: Synthesis, inhibitory activity, and mechanism. *Bioorg Med Chem* 2016;24:4620–5.
48. Yue LM, Lee J, Lü ZR, et al. Effect of Cd²⁺ on tyrosinase: integration of inhibition kinetics with computational simulation. *Int J Biol Macromol* 2017; 94:836–44.
49. Gao H. Predicting tyrosinase inhibition by 3D QSAR pharmacophore models and designing potential tyrosinase inhibitors from traditional Chinese medicine database. *Phytomedicine* 2018;38:145–57.
50. Khan MT. Novel tyrosinase inhibitors from natural resources – their computational studies. *Curr Med Chem* 2012;19: 2262–72.
51. Chan CF, Huang CC, Lee MY, Lin YS. Fermented broth in tyrosinase- and melanogenesis inhibition. *Molecules* 2014; 19:13122–35.
52. Chang TS. An updated review of tyrosinase inhibitors. *Int J Mol Sci* 2009;10:2440–75.
53. Chang TS. Natural melanogenesis inhibitors acting through the down-regulation of tyrosinase activity. *Materials* 2012;5: 1661–85.
54. Chen CY, Lin LC, Yang WF, et al. An updated organic classification of tyrosinase inhibitors on melanin biosynthesis. *Curr Org Chem* 2015;19:4–18.
55. Hassan Khan MT. Molecular design of tyrosinase inhibitors: a critical review of promising novel inhibitors from synthetic origins. *Pure Appl Chem* 2007;79:2277–95.
56. Likhitwitayawuid K. Stilbenes with tyrosinase inhibitory activity. *Curr Sci* 2008;94:44–52.
57. Lin JW, Chiang HM, Lin YC, Wen KC. Natural products with skin – whitening effects. *J Food Drug Anal* 2008;16:1–10.
58. Loizzo MR, Tundis R, Menichini F. Natural and synthetic tyrosinase inhibitors as antibrowning agents: an update. *Compr Rev Food Sci Food Saf* 2012;11:378–98.
59. Parvez S, Kang M, Chung HS, Bae H. Naturally occurring tyrosinase inhibitors: mechanism and applications in skin health, cosmetics and agriculture industries. *Phytother Res* 2007;21:805–16.
60. Fernandes MS, Kerkar S. Microorganisms as a source of tyrosinase inhibitors: a review. *Ann Microbiol* 2017;67: 343–58.
61. Pillaiyar T, Manickam M, Namasivayam V. Skin whitening agents: medicinal chemistry perspective of tyrosinase inhibitors. *J Enzyme Inhib Med Chem* 2017;32:403–25.
62. Wu B. Tyrosinase inhibitors from terrestrial and marine resources. *Curr Top Med Chem* 2014;14:1425–49.
63. Brenner M, Hearing VJ. The protective role of melanin against UV damage in human skin. *Photochem Photobiol* 2008;84:539–49.
64. Cestari TF, Dantas LP, Boza JC. Acquired hyperpigmentations. *An Bras Dermatol* 2014;89:11–25.
65. Nouveau S, Agrawal D, Kohli M, et al. Skin hyperpigmentation in Indian population: insights and best practice. *Indian J Dermatol* 2016;61:487–95.
66. Dorga S, Sarangal R. Pigmentary disorders: an insight. *Pigment Int* 2014;1:5–7.
67. Garcia-Jimenez A, Teruel-Puche JA, Garcia-Ruiz PA, et al. Action of 2,2',4,4'-tetrahydroxybenzophenone in the biosynthesis pathway of melanin. *Int J Biol Macromol* 2017; 98:622–9.
68. Bull AT, Carter BLA. The isolation of tyrosinase from *Aspergillus nidulans*, its kinetic and molecular properties and some consideration of its activity in vivo. *J Gen Microbiol* 1973;75:61–73.
69. Silva S. C d, Wisniewski C, Luccas PO, Magalhães CSD. Enzyme from banana (*Musa* sp.) extraction procedures for sensitive adrenaline biosensor construction. *Am J Analyt Chem* 2013;04: 293–300.
70. Dolashki A, Voelter W, Gushterova A, et al. Isolation and characterization of novel tyrosinase from *Laceyella sacchari*. *Protein Pept Lett* 2012;19:538–43.
71. Haghbeen K, Rastegar j. F, Karkhaneh AA, Shareefi Borojerdi SH. Purification of tyrosinase from edible mushroom. *Iran J Biotechnol* 2004;2:189–94.
72. Sambasiva Rao KRS, Tripathy NK, Srinivasa Rao D, Prakasham RS. Production, characterization, catalytic and inhibitory activities of tyrosinase. *Res J Biotech* 2013;8: 187–99.
73. Vieira NCS, Ferreira RA, Valquiria da CR, et al. Self-assembled films containing crude extract of avocado as a source of tyrosinase for monophenol detection. *Mater Sci Eng C* 2013;33:3899–902.
74. Yamauchi K, Mitsunaga T, Batubara I. Isolation, identification and tyrosinase inhibitory activities of the extractives from *Allamanda cathartica*. *Nat Res* 2011;2:167–72.
75. Yuan H, Ke-wu L, Dong Y, et al. Some properties of potato tyrosinase, chemical research and application. *J Chem Res App* 2005;1:22–7.
76. Zh Y, Wu F. Catalytic properties of tyrosinase from potato and edible fungi. *Biotechnology* 2006;5:344–8.
77. Harir M, Bellahcene M, Baratto MC, et al. Isolation and characterization of a novel tyrosinase produced by Sahara soil actinobacteria and immobilization on nylon nanofiber membranes. *J Biotechnol* 2018;265:54–64.
78. Vanitha M, Soundhari C. Isolation and characterisation of mushroom tyrosinase and screening of herbal extracts for anti-tyrosinase activity. *Int J ChemTech Research* 2017;10: 1156–67.
79. Zekiri F, Molitor C, G.Mauracher SG, et al. Purification and characterization of tyrosinase from walnut leaves (*Juglans regia*). *Phytochemistry* 2014;101:5–15.
80. Gasparetti C. Biochemical and structural characterisation of the copper containing oxidoreductases catechol oxidase, tyrosinase, and laccase from ascomycete fungi. Espoo: VTT Technical Research Centre of Finland; 2012.
81. Boekelheide K, Graham DG, Mize PD, Jeffs PW. The metabolic pathway catalyzed by the tyrosinase of *Agaricus bisporus*. *J Biol Chem* 1980;255:4766–71.
82. Ioniță E, Stănciuc N, Aprodu I, et al. pH-induced structural changes of tyrosinase from *Agaricus bisporus* using fluorescence and in silico methods. *J Sci Food Agric* 2014;94: 2338–44.
83. Ioniță E, Aprodu I, Stănciuc N, et al. Advances in structure-function relationships of tyrosinase from *Agaricus bisporus* – investigation on heat-induced conformational changes. *Food Chem* 2014;156:129–36.
84. Khan IA, Ali R. Antigenicity, catalytic activity and conformation of *Agaricus bisporus* tyrosinase: interaction of conformation-directed antibodies with the native and irradiated enzyme. *J Biochem* 1986;99:445–52.
85. Zhou L, Liu W, Zou L, et al. Aggregation and conformational change of mushroom (*Agaricus bisporus*) polyphenoloxidase subjected to thermal treatment. *Food Chem* 2017;214:423–31.

86. Gheibi N, Saboury AA, Haghbeen K, Moosavi Movahedi AA. The effect of some osmolytes on the activity and stability of mushroom tyrosinase. *J Biosci* 2006;31:355–62.
87. Narin R, Cresswell W, J Narin J. Mushroom tyrosinase: a model system to combine experimental investigation of enzyme-catalyzed reactions, data handling using R, and enzyme-inhibitor structural studies. *Biochem Mol Biol Educ* 2015;43:370–6.
88. Della Longa S, Ascone I, Bianconi A, et al. The dinuclear copper site structure of *Agaricus bisporus* tyrosinase in solution probed by X-ray absorption spectroscopy. *J Biol Chem* 1996;271:21025–30.
89. Ismaya WT, Tandrasasmita OM, Sundari S, et al. The light subunit of mushroom *Agaricus bisporus* tyrosinase: its biological characteristics and implications. *Int J Biol Macromol* 2017;102:308–14.
90. Ismaya WT, Rozeboom HJ, Weijn A, et al. Crystal structure of *Agaricus bisporus* mushroom tyrosinase: identity of the tetramer subunits and interaction with tropolone. *Biochemistry* 2011;50:5477–86.
91. Strothkamp KG, Jolley RL, Mason HS. Quaternary structure of mushroom tyrosinase. *Biochem Biophys Res Commun* 1976;70:519–24.
92. Bourquelot E, Bertrand A. Le bluissement et le noircissement des champignons. *Comp Rend Soc Biol* 1895;2:582–4.
93. van Gelder CW, Flurkey WH, Wichers HJ. Sequence and structural features of plant and fungal tyrosinases. *Phytochemistry* 1997;45:1309–23.
94. Lopez-Tejedor D, Palomo JM. Efficient purification of a highly active H-subunit of tyrosinase from *Agaricus bisporus*. *Protein Expr Purif* 2018;145:64–70.
95. Sanchez-Ferrer A, Rodriguez-Lopez JN, Garcia-Canovas F, Garcia-Carmona F. Tyrosinase: a comprehensive review of its mechanism. *Biochim Biophys Acta* 1995;1247:1–11.
96. Ortiz-Ruiz CV, Maria-Solano MA, Garcia-Molina Mdel M, et al. Kinetic characterization of substrate-analogous inhibitors of tyrosinase. *IUBMB Life* 2015;67:757–67.
97. Dec J, Bollag JM. Effect of various factors on dehalogenation of chlorinated phenols and anilines during oxidative coupling. *Environ Sci Technol* 1995;29:657–63.
98. Munoz-Munoz JL, Garcia-Molina F, Varon R, et al. Suicide inactivation of the diphenolase and monophenolase activities of tyrosinase. *IUBMB Life* 2010;62:539–47.
99. Land EJ, Ramsden CA, Riley PA. The mechanism of suicide-inactivation of tyrosinase: a substrate structure investigation. *Tohoku J Exp Med* 2007;212:341–8.
100. Haghbeen K, Saboury AA, Karbassi F. Substrate share in the suicide inactivation of mushroom tyrosinase. *Biochim Biophys Acta* 2004;1675:139–46.
101. Saboury AA, Karbassi F, Haghbeen K, et al. Stability, structural and suicide inactivation changes of mushroom tyrosinase after acetylation by *n*-acetylimidazole. *Int J Biol Macromol* 2004;34:257–62.
102. Munoz-Munoz JL, Garcia-Molina F, Garcia-Ruiz PA, et al. Phenolic substrates and suicide inactivation of tyrosinase: kinetics and mechanism. *Biochem J* 2008;416:431–40.
103. Munoz-Munoz JL, Garcia-Molina F, Garcia-Ruiz PA, et al. Stereospecific inactivation of tyrosinase by L- and D-ascorbic acid. *Biochim Biophys Acta* 2009;1794:244–53.
104. Munoz-Munoz JL, Acosta-Motos JR, Garcia-Molina F, et al. Tyrosinase inactivation in its action on dopa. *Biochim Biophys Acta* 2010;1804:1467–75.
105. Munoz-Munoz JL, Garcia-Molina F, Berna J, et al. Kinetic characterisation of *o*-aminophenols and aromatic *o*-diamines as suicide substrates of tyrosinase. *Biochim Biophys Acta* 2012;1824:647–55.
106. Muñoz-Muñoz JL, Berna J, Garcia-Molina F, et al. Unravelling the suicide inactivation of tyrosinase: a discrimination between mechanisms. *J Mol Catal B Enzym* 2012;75:11–9.
107. Munoz-Munoz JL, Garcia-Molina Mdel M, Garcia-Molina F, et al. Indirect inactivation of tyrosinase in its action on 4-*tert*-butylphenol. *J Enzyme Inhib Med Chem* 2014;29:344–52.
108. Munoz-Munoz JL, Garcia-Molina F, Acosta-Motos JR, et al. Indirect inactivation of tyrosinase in its action on tyrosine. *Acta Biochim Pol* 2011;58:477–88.
109. del Mar Garcia-Molina M, Munoz-Munoz JL, Berna J, et al. Catalysis and inactivation of tyrosinase in its action on hydroxyhydroquinone. *IUBMB Life* 2014;66:122–7.
110. Munoz-Munoz JL, Garcia-Molina F, Arribas E, et al. Suicide inactivation of tyrosinase in its action on tetrahydropterines. *J Enzyme Inhib Med Chem* 2011;26:728–33.
111. Garcia-Molina F, Munoz-Munoz JL, Martinez-Ortiz F, et al. Tetrahydrofolic acid is a potent suicide substrate of mushroom tyrosinase. *J Agric Food Chem* 2011;59:1383–91.
112. Garcia-Molina F, Munoz-Munoz JL, Garcia-Molina M, et al. Melanogenesis inhibition due to NADH. *Biosci Biotechnol Biochem* 2010;74:1777–87.
113. Park J, Jung H, Kim K, et al. D-tyrosine negatively regulates melanin synthesis by competitively inhibiting tyrosinase activity. *Pigment Cell Melanoma Res* 2018;31:374–83.
114. Hassani S, Gharechaei B, Nikfard S, et al. New insight into the allosteric effect of L-tyrosine on mushroom tyrosinase during L-dopa production. *Int J Biol Macromol* 2018;114:821–9.
115. Zhao DY, Zhang MX, Dong XW, et al. Design and synthesis of novel hydroxypyridinone derivatives as potential tyrosinase inhibitors. *Bioorg Med Chem Lett* 2016;26:3103–8.
116. Yin SJ, Si YX, Qian GY. Inhibitory effect of phthalic acid on tyrosinase: the mixed-type inhibition and docking simulations. *Enzyme Res* 2011;2011:1. doi: [10.4061/2011/294724](https://doi.org/10.4061/2011/294724).
117. Yin SJ, Si YX, Chen YF, et al. Mixed-type inhibition of tyrosinase from *Agaricus bisporus* by terephthalic acid: computational simulations and kinetics. *Protein J* 2011;30:273–80.
118. Liu HJ, Ji S, Fan YQ, et al. The effect of D-(–)-arabinose on tyrosinase: an integrated study using computational simulation and inhibition kinetics. *Enzyme Res* 2012;2012:731427. doi: [10.1155/2012/731427](https://doi.org/10.1155/2012/731427).
119. Hridya H, Amrita A, Sankari M, et al. Inhibitory effect of brazilein on tyrosinase and melanin synthesis: kinetics and in silico approach. *Int J Biol Macromol* 2015;81:228–34.
120. Ashraf Z, Rafiq M, Seo SY, et al. Kinetic and in silico studies of novel hydroxy-based thymol analogues as inhibitors of mushroom tyrosinase. *Eur J Med Chem* 2015;98:203–11.
121. Karbassi F, Saboury AA, Khan MT, et al. Mushroom tyrosinase inhibition by two potent uncompetitive inhibitors. *J Enzyme Inhib Med Chem* 2004;19:349–53.
122. Seo B, Yun J, Lee S, et al. Barbarin as a new tyrosinase inhibitor from *Barbarea orthoceras*. *Planta Med* 1999;65:683–6.
123. Hu YH, Liu X, Jia YL, et al. Inhibitory kinetics of chlorocinnamic acids on mushroom tyrosinase. *J Biosci Bioeng* 2014;117:142–6.

124. Gheibi N, Saboury AA, Haghbeen K, et al. Dual effects of aliphatic carboxylic acids on cresolase and catecholase reactions of mushroom tyrosinase. *J Enzyme Inhib Med Chem* 2009;24:1076–81.
125. Shiino M, Watanabe Y, Umezawa K. Synthesis and tyrosinase inhibitory activity of novel N-hydroxybenzyl-N-nitrosodihydroxylamines. *Bioorg Chem* 2003;31:129–35.
126. Chen QX, Song KK, Wang Q, Huang H. Inhibitory effects on mushroom tyrosinase by some alkylbenzaldehydes. *J Enzyme Inhib Med Chem* 2003;18:491–6.
127. Saeed A, Mahesar PA, Channar PA, et al. Synthesis, molecular docking studies of coumarinyl-pyrazolinyl substituted thiazoles as non-competitive inhibitors of mushroom tyrosinase. *Bioorg Chem* 2017;74:187–96.
128. Mann T, Gerwat W, Batzer J, et al. Inhibition of human tyrosinase requires molecular motifs distinctively different from mushroom tyrosinase. *J Invest Dermatol* 2018; 138:1601–8.
129. Hamed SH, Sriwiranont P, deLong MA, et al. Comparative efficacy and safety of deoxyarbutin, a new tyrosinase-inhibiting agent. *J Cosmet Sci* 2006;57:291–308.
130. Sugimoto K, Nishimura T, Nomura K, et al. Syntheses of arbutin- α -glycosides and a comparison of their inhibitory effects with those of α -arbutin and arbutin on human tyrosinase. *Chem Pharm Bull (Tokyo)* 2003;51:798–801.
131. Di Petrillo A, Gonzalez-Paramas AM, Era B, et al. Tyrosinase inhibition and antioxidant properties of *Asphodelus microcarpus* extracts. *BMC Complement Altern Med* 2016;16:453.
132. Koyu H, Kazan A, Demir S, et al. Optimization of microwave assisted extraction of *Morus nigra* L. Fruits maximizing tyrosinase inhibitory activity with isolation of bioactive constituents. *Food Chem* 2018;248:183–91.
133. Lee SG, Karadeniz F, Seo Y, Kong CS. Anti-melanogenic effects of flavonoid glycosides from *Limonium tetragonum* (thunb.) bullock via inhibition of tyrosinase and tyrosinase-related proteins. *Molecules* 2017;22:1480–90.
134. Matsuo K, Kobayashi M, Takuno Y, et al. Anti-tyrosinase activity constituents of *Arctostaphylos uva-ursi*. *Yakugaku Zasshi* 1997;117:1028–32.
135. Alam N, Yoon KN, Lee JS, et al. Consequence of the antioxidant activities and tyrosinase inhibitory effects of various extracts from the fruiting bodies of *Pleurotus ferulae*. *Saudi J Biol Sci* 2012;19:111–18.
136. Kim NY, Kwon HS, Lee HY. Effect of inhibition on tyrosinase and melanogenesis of *Agastache rugosa* Kuntze by lactic acid bacteria fermentation. *J Cosmet Dermatol* 2017;16:407–15.
137. Taherkhani M. Chemical constituents, total phenolic content, antimicrobial, antioxidant and radical scavenging properties, chelating ability, tyrosinase inhibition and in vitro cytotoxic effects of *Artemisia aucheri* herbs. *Pharm Chem J* 2017;50:736–45.
138. Lee GY, Cho BO, Shin JY, et al. Tyrosinase inhibitory components from the seeds of *Cassia tora*. *Arch Pharm Res* 2018;41:490–6.
139. Senol FS, Orhan I, Yilmaz G, et al. Acetylcholinesterase, butyrylcholinesterase, and tyrosinase inhibition studies and antioxidant activities of 33 *Scutellaria* L. taxa from Turkey. *Food Chem Toxicol* 2010;48:781–8.
140. Ya W, Chun-Meng Z, Tao G, et al. Preliminary screening of 44 plant extracts for anti-tyrosinase and antioxidant activities. *Pak J Pharm Sci* 2015;28:1737–44.
141. Nithitanakool S, Pithayanukul P, Bavovada R, Sapparpakorn P. Molecular docking studies and anti-tyrosinase activity of Thai mango seed kernel extract. *Molecules* 2009;14:257–65.
142. Abdillahi HS, Finnie JF, Van Staden J. Anti-inflammatory, antioxidant, anti-tyrosinase and phenolic contents of four *Podocarpus* species used in traditional medicine in South Africa. *J Ethnopharmacol* 2011;136:496–503.
143. Saeio K, Yotsawimonwat S, Anuchapreeda S, Okonogi S. Development of microemulsion of a potent anti-tyrosinase essential oil of an edible plant. *Drug Discov Ther* 2011;5:246–52.
144. Mapunya MB, Hussein AA, Rodriguez B, Lall N. Tyrosinase activity of *Greyia flanaganii* (Bolus) constituents. *Phytomedicine* 2011;18:1006–12.
145. Lin YS, Chen HJ, Huang JP, et al. Kinetics of tyrosinase inhibitory activity using *Vitis vinifera* leaf extracts. *Biomed Res Int* 2017;2017:5232680. doi: 10.1155/2017/5232680.
146. Huang MH, Tai HM, Wang BS, Chang LW. Inhibitory effects of water extract of Flos Inulae on mutation and tyrosinase. *Food Chem* 2013;139:1015–20.
147. Chiari ME, Joray MB, Ruiz G, et al. Tyrosinase inhibitory activity of native plants from central argentina: isolation of an active principle from *Lithrea molleoides*. *Food Chem* 2010;120:10–4.
148. Burlando B, Clericuzio M, Cornara L. Moraceae plants with tyrosinase inhibitory activity: A review. *Mini Rev Med Chem* 2017;17:108–21.
149. Chatatikun M, Chiabchalard A. Thai plants with high antioxidant levels, free radical scavenging activity, anti-tyrosinase and anti-collagenase activity. *BMC Complement Altern Med* 2017;17:487.
150. Issa RA, Afifi FU, Amro BI. Studying the anti-tyrosinase effect of *Arbutus andrachne* L. extracts. *Int J Cosmet Sci* 2008;30:271–6.
151. Hameed A, Akhtar N. Comparative chemical investigation and evaluation of antioxidant and tyrosinase inhibitory effects of *Withania somnifera* (L.) Dunal and *Solanum nigrum* (L.) berries. *Acta Pharm* 2018;68:47–60.
152. Neagu E, Radu GL, Albu C, Paun G. Antioxidant activity, acetylcholinesterase and tyrosinase inhibitory potential of *Pulmonaria officinalis* and *Centarium umbellatum* extracts. *Saudi J Biol Sci* 2018;25:578–85.
153. Panda P, Dash P, Ghosh G. Chemometric profile, antioxidant and tyrosinase inhibitory activity of Camel's foot creeper leaves (*Bauhinia vahlii*). *Nat Prod Res* 2018;32:596–9.
154. Quispe YN, Hwang SH, Wang Z, Lim SS. Screening of peruvian medicinal plants for tyrosinase inhibitory properties: identification of tyrosinase inhibitors in *Hypericum laricifolium* Juss. *Molecules* 2017;22:doi: 10.3390/molecules 22030402.
155. Suh SS, Hwang J, Park M, et al. Phenol content, antioxidant and tyrosinase inhibitory activity of mangrove plants in Micronesia. *Asian Pac J Trop Med* 2014;7:531–5.
156. Hun Son K, Young Heo M. Inhibitory effects of Korean indigenous plants on tyrosinase and melanogenesis. *J Cosmet Sci* 2013;64:145–58.
157. Souza PM, Elias ST, Simeoni LA, et al. Plants from Brazilian cerrado with potent tyrosinase inhibitory activity. *PLoS One* 2012;7:e48589.
158. Khazaeli P, Goldoosian R, Shariffar F. An evaluation of extracts of five traditional medicinal plants from Iran

- on the inhibition of mushroom tyrosinase activity and scavenging of free radicals. *Int J Cosmet Sci* 2009;31:375–81.
159. Masuda T, Fujita N, Odaka Y, et al. Tyrosinase inhibitory activity of ethanol extracts from medicinal and edible plants cultivated in Okinawa and identification of a water-soluble inhibitor from the leaves of *Nandina domestica*. *Biosci Biotechnol Biochem* 2007;71:2316–20.
 160. Masuda T, Yamashita D, Takeda Y, Yonemori S. Screening for tyrosinase inhibitors among extracts of seashore plants and identification of potent inhibitors from *Garcinia subelliptica*. *Biosci Biotechnol Biochem* 2005;69:197–201.
 161. Baurin N, Arnoult E, Scior T, et al. Preliminary screening of some tropical plants for anti-tyrosinase activity. *J Ethnopharmacol* 2002;82:155–8.
 162. Kubo I, Yokokawa Y, Kinst-Hori I. Tyrosinase inhibitors from Bolivian medicinal plants. *J Nat Prod* 1995;58:739–43.
 163. Bonesi M, Xiao J, Tundis R, et al. Advances in the tyrosinase inhibitors from plant source. *Curr Med Chem* 2018;25:1. doi: [10.2174/0929867325666180522091311](https://doi.org/10.2174/0929867325666180522091311).
 164. Vasantha KY, Muruges CS, Sattur AP. A tyrosinase inhibitor from *Aspergillus niger*. *J Food Sci Technol* 2014;51:2877–80.
 165. Tsuchiya T, Yamada K, Minoura K, et al. Purification and determination of the chemical structure of the tyrosinase inhibitor produced by *Trichoderma viride* strain H1-7 from a marine environment. *Biol Pharm Bull* 2008;31:1618–20.
 166. Lu R, Liu X, Gao S, et al. New tyrosinase inhibitors from *Paecilomyces gunnii*. *J Agric Food Chem* 2014;62:11917–23.
 167. Kang HS, Choi JH, Cho WK, et al. A sphingolipid and tyrosinase inhibitors from the fruiting body of *Phellinus linteus*. *Arch Pharm Res* 2004;27:742–50.
 168. Morimura K, Yamazaki C, Hattori Y, et al. A tyrosinase inhibitor, daedalin a, from mycelial culture of *Daedalea dickinsii*. *Biosci Biotechnol Biochem* 2007;71:2837–40.
 169. Sharma VK, Choi J, Sharma N, et al. In vitro anti-tyrosinase activity of 5-(hydroxymethyl)-2-furfural isolated from *Dictyophora indusiata*. *Phytother Res* 2004;18:841–4.
 170. Ishihara A, Ide Y, Bito T, et al. Novel tyrosinase inhibitors from liquid culture of *Neolentinus lepideus*. *Biosci Biotechnol Biochem* 2018;82:22–30.
 171. Li X, Kim MK, Lee U, et al. Myrothenones A and B, cyclopentenone derivatives with tyrosinase inhibitory activity from the marine-derived fungus *Myrothecium* sp. *Chem Pharm Bull (Tokyo)* 2005;53:453–5.
 172. Wu B, Wu X, Sun M, Li M. Two novel tyrosinase inhibitory sesquiterpenes induced by cucl2 from a marine-derived fungus *Pestalotiopsis* sp. Z233. *Mar Drugs* 2013;11:2713–21.
 173. Chang TS, Tseng M, Ding HY, Shou-Ku Tai S. Isolation and characterization of *Streptomyces hiroshimensis* strain TI-C3 with anti-tyrosinase activity. *J Cosmet Sci* 2008;59:33–40.
 174. le Roes-Hill M, Prins A, Meyers PR. *Streptomyces swartbergensis* sp. Nov., a novel tyrosinase and antibiotic producing actinobacterium. *Antonie Van Leeuwenhoek* 2018;111:589–600.
 175. Nakashima T, Anzai K, Kuwahara N, et al. Physicochemical characters of a tyrosinase inhibitor produced by *Streptomyces roseolilacinus* NBRC 12815. *Biol Pharm Bull* 2009;32:832–6.
 176. Deering RW, Chen J, Sun J, et al. N-acyl dehydrotyrosines, tyrosinase inhibitors from the marine bacterium *Thalassotalea* sp. PP2-459. *J Nat Prod* 2016;79:447–50.
 177. Sano T, Kaya K. Oscillapeptin G, a tyrosinase inhibitor from toxic *Oscillatoria agardhii*. *J Nat Prod* 1996;59:90–2.
 178. Ji K, Cho YS, Kim YT. Tyrosinase inhibitory and anti-oxidative effects of lactic acid bacteria isolated from dairy cow feces. *Probiotics Antimicrob Proteins* 2018;10:43–55.
 179. Wang GH, Chen CY, Tsai TH, et al. Evaluation of tyrosinase inhibitory and antioxidant activities of *Angelica dahurica* root extracts for four different probiotic bacteria fermentations. *J Biosci Bioeng* 2017;123:679–84.
 180. Crozier A, Jaganath IB, Clifford MN. Phenols, polyphenols and tannins: an overview. *Plant Secondary Metabolites* 2007;1–24. [10.1002/9780470988558.ch1](https://doi.org/10.1002/9780470988558.ch1).
 181. Sakuma K, Ogawa M, Sugibayashi K, et al. Relationship between tyrosinase inhibitory action and oxidation-reduction potential of cosmetic whitening ingredients and phenol derivatives. *Arch Pharm Res* 1999;22:335–9.
 182. Hashimoto A, Ichihashi M, Mishima Y. The mechanism of depigmentation by hydroquinone: a study on suppression and recovery processes of tyrosinase activity in the pigment cells in vivo and in vitro. *Nihon Hifuka Gakkai Zasshi* 1984;94:797–804.
 183. Chen YR, Y-Y R, Lin TY, et al. Identification of an alkylhydroquinone from *Rhus succedanea* as an inhibitor of tyrosinase and melanogenesis. *J Agric Food Chem* 2009;57:2200–5.
 184. Sasaki A, Yamano Y, Sugimoto S, et al. Phenolic compounds from the leaves of *Breynia officinalis* and their tyrosinase and melanogenesis inhibitory activities. *J Nat Med* 2018;72:381–9.
 185. Chawla S, deLong MA, Visscher MO, et al. Mechanism of tyrosinase inhibition by deoxyarbutin and its second-generation derivatives. *Br J Dermatol* 2008;159:1267–74.
 186. Boissy RE, Visscher M, DeLong MA. Deoxyarbutin: a novel reversible tyrosinase inhibitor with effective in vivo skin lightening potency. *Exp Dermatol* 2005;14:601–8.
 187. Chawla S, Kvalnes K, deLong MA, et al. Deoxyarbutin and its derivatives inhibit tyrosinase activity and melanin synthesis without inducing reactive oxygen species or apoptosis. *J Drugs Dermatol* 2012;11:e28–34.
 188. Tasaka K, Kamei C, Nakano S, et al. Effects of certain resorcinol derivatives on the tyrosinase activity and the growth of melanoma cells. *Methods Find Exp Clin Pharmacol* 1998;20:99–109.
 189. Kolbe L, Mann T, Gerwat W, et al. 4-*n*-Butylresorcinol, a highly effective tyrosinase inhibitor for the topical treatment of hyperpigmentation. *J Eur Acad Dermatol Venereol* 2013;27:19–23.
 190. Nguyen MH, Nguyen HX, Nguyen MT, Nguyen NT. Phenolic constituents from the heartwood of *Artocapus altalis* and their tyrosinase inhibitory activity. *Nat Prod Commun* 2012;7:185–6.
 191. Ashraf Z, Rafiq M, Seo SY, et al. Synthesis, kinetic mechanism and docking studies of vanillin derivatives as inhibitors of mushroom tyrosinase. *Bioorg Med Chem* 2015;23:5870–80.
 192. Shirota S, Miyazaki K, Aiyama R, et al. Tyrosinase inhibitors from crude drugs. *Biol Pharm Bull* 1994;17:266–9.
 193. Matsumoto T, Nakajima T, Iwadate T, Nihei KI. Chemical synthesis and tyrosinase-inhibitory activity of isotachioside and its related glycosides. *Carbohydr Res* 2018;465:22–8.
 194. Deri B, Kanteev M, Goldfeder M, et al. The unravelling of the complex pattern of tyrosinase inhibition. *Sci Rep* 2016;6:34993.
 195. Garcia-Jimenez A, Teruel-Puche JA, Berna J, et al. Action of tyrosinase on alpha and beta-arbutin: a kinetic study. *PLoS One* 2017;12:e0177330.

196. Garcia-Molina MO, Munoz-Munoz JL, Garcia-Molina F, et al. Study of umbelliferone hydroxylation to esculetin catalyzed by polyphenol oxidase. *Biol Pharm Bull* 2013;36:1140–5.
197. Maria Del Mar Garcia-Molina JB, Muñoz-Muñoz JL, García-Ruiz PA, Moreno MG, Martínez JR, García-Canovas F. Action of tyrosinase on hydroquinone in the presence of catalytic amounts of *o*-diphenol. A kinetic study. *React Kinet Mech Cat* 2014;112:305–20.
198. Garcia-Molina Mdel M, Munoz Munoz JL, Martinez-Ortiz F, et al. Tyrosinase-catalyzed hydroxylation of hydroquinone, a depigmenting agent, to hydroxyhydroquinone: a kinetic study. *Bioorg Med Chem* 2014;22:3360–9.
199. Ortiz-Ruiz CV, Berna J, Garcia-Molina Mdel M, et al. Identification of *p*-hydroxybenzyl alcohol, tyrosol, phloretin and its derivate phloridzin as tyrosinase substrates. *Bioorg Med Chem* 2015;23:3738–46.
200. Ortiz-Ruiz CV, Berna J, Rodriguez-Lopez JN, et al. Tyrosinase-catalyzed hydroxylation of 4-hexylresorcinol, an antibrowning and depigmenting agent: A kinetic study. *J Agric Food Chem* 2015;63:7032–40.
201. Ortiz-Ruiz CV, Ballesta de Los Santos M, Berna J, et al. Kinetic characterization of oxyresveratrol as a tyrosinase substrate. *IUBMB Life* 2015;67:828–36.
202. Garcia-Jimenez A, Teruel-Puche JA, Ortiz-Ruiz CV, et al. 4-*n*-Butylresorcinol, a depigmenting agent used in cosmetics, reacts with tyrosinase. *IUBMB Life* 2016;68:663–72.
203. Garcia-Jimenez A, Teruel-Puche JA, Berna J, et al. Characterization of the action of tyrosinase on resorcinols. *Bioorg Med Chem* 2016;24:4434–43.
204. Garcia-Jimenez A, Munoz-Munoz JL, Garcia-Molina F, et al. Spectrophotometric characterization of the action of tyrosinase on *p*-coumaric and caffeic acids: characteristics of *o*-caffeoquinone. *J Agric Food Chem* 2017;65:3378–86.
205. Garcia-Jimenez A, Garcia-Molina F, Teruel-Puche JA, et al. Catalysis and inhibition of tyrosinase in the presence of cinnamic acid and some of its derivatives. *Int J Biol Macromol* 2018;119:548–54.
206. Garcia-Jimenez A, Teruel-Puche JA, Garcia-Ruiz PA, et al. Structural and kinetic considerations on the catalysis of deoxyarbutin by tyrosinase. *PLoS One* 2017;12:e0187845.
207. Ortiz-Ruiz CV, Garcia-Molina Mdel M, Serrano JT, et al. Discrimination between alternative substrates and inhibitors of tyrosinase. *J Agric Food Chem* 2015;63:2162–71.
208. Lin D, Xiao M, Zhao J, et al. An overview of plant phenolic compounds and their importance in human nutrition and management of type 2 diabetes. *Molecules* 2016;21:1374. <https://doi.org/10.3390/molecules21101374>
209. Huang WY, Cai YZ, Zhang Y. Natural phenolic compounds from medicinal herbs and dietary plants: potential use for cancer prevention. *Nutr Cancer* 2010;62:1–20.
210. Wang Y, Curtis-Long MJ, Lee BW, et al. Inhibition of tyrosinase activity by polyphenol compounds from *Flemingia philippinensis* roots. *Bioorg Med Chem* 2014;22:1115–20.
211. Xue YL, Miyakawa T, Hayashi Y, et al. Isolation and tyrosinase inhibitory effects of polyphenols from the leaves of persimmon, *Diospyros kaki*. *J Agric Food Chem* 2011;59:6011–7.
212. Sirat HM, Rezali MF, Ujang Z. Isolation and identification of radical scavenging and tyrosinase inhibition of polyphenols from *Tibouchina semidecandra* L. *J Agric Food Chem* 2010;58:10404–9.
213. Yoon NY, Eom TK, Kim MM, Kim SK. Inhibitory effect of phlorotannins isolated from *Ecklonia cava* on mushroom tyrosinase activity and melanin formation in mouse B16F10 melanoma cells. *J Agric Food Chem* 2009;57:4124–9.
214. Fujimoto A, Shingai Y, Nakamura M, et al. A novel ring-expanded product with enhanced tyrosinase inhibitory activity from classical Fe-catalyzed oxidation of rosmarinic acid, a potent antioxidative lamiaceae polyphenol. *Bioorg Med Chem Lett* 2010;20:7393–6.
215. Solimine J, Garo E, Wedler J, et al. Tyrosinase inhibitory constituents from a polyphenol-enriched fraction of rose oil distillation wastewater. *Fitoterapia* 2016;108:13–9.
216. Du ZY, Jiang YF, Tang ZK, et al. Antioxidation and tyrosinase inhibition of polyphenolic curcumin analogs. *Biosci Biotechnol Biochem* 2011;75:2351–8.
217. Jiang Y, Du Z, Xue G, et al. Synthesis and biological evaluation of unsymmetrical curcumin analogues as tyrosinase inhibitors. *Molecules* 2013;18:3948–61.
218. Ng LT, Ko HH, Lu TM. Potential antioxidants and tyrosinase inhibitors from synthetic polyphenolic deoxybenzoins. *Bioorg Med Chem* 2009;17:4360–6.
219. Zheng ZP, Zhang YN, Zhang S, Chen J. One-pot green synthesis of 1,3,5-triarylpentane-1,5-dione and triarylmethane derivatives as a new class of tyrosinase inhibitors. *Bioorg Med Chem Lett* 2016;26:795–8.
220. Orhan IE, Khan MT. Flavonoid derivatives as potent tyrosinase inhibitors – a survey of recent findings between 2008–2013. *Curr Top Med Chem* 2014;14:1486–93.
221. Jegal J, Park SA, Chung K, et al. Tyrosinase inhibitory flavonoid from *Juniperus communis* fruits. *Biosci Biotechnol Biochem* 2016;80:2311–7.
222. Muhammad D, Hubert J, Lalun N, et al. Isolation of flavonoids and triterpenoids from the fruits of *Alphitonia neocaledonica* and evaluation of their anti-oxidant, anti-tyrosinase and cytotoxic activities. *Phytochem Anal* 2015;26:137–44.
223. Erdogan Orhan I, Senol FS, Aslan Erdem S, et al. Tyrosinase and cholinesterase inhibitory potential and flavonoid characterization of *Viola odorata* L. (Sweet Violet). *Phytother Res* 2015;29:1304–10.
224. Liang CP, Chang CH, Liang CC, et al. In vitro antioxidant activities, free radical scavenging capacity, and tyrosinase inhibitory of flavonoid compounds and ferulic acid from *Spiranthes sinensis* (Pers.). *AMES Mol* 2014;19:4681–94.
225. Badria FA, elGayyar MA. A new type of tyrosinase inhibitors from natural products as potential treatments for hyperpigmentation. *Boll Chim Farm* 2001;140:267–71.
226. Promden W, Viriyabancha W, Monthakantirat O, et al. Correlation between the potency of flavonoids on mushroom tyrosinase inhibitory activity and melanin synthesis in melanocytes. *Molecules* 2018;23:1403. doi: [10.3390/molecules23061403](https://doi.org/10.3390/molecules23061403).
227. Demirkiran O, Sabudak T, Ozturk M, Topcu G. Antioxidant and tyrosinase inhibitory activities of flavonoids from *Trifolium nigrescens* Subsp. petrisavi. *J Agric Food Chem* 2013;61:12598–603.
228. Yao Y, Cheng X, Wang L, et al. Mushroom tyrosinase inhibitors from mung bean (*Vigna radiatae* L.) extracts. *Int J Food Sci Nutr* 2012;63:358–61.
229. Lou SN, Yu MW, Ho CT. Tyrosinase inhibitory components of immature calamondin peel. *Food Chem* 2012;135:1091–6.
230. Hu X, Wu JW, Wang M, et al. 2-Arylbenzofuran, flavonoid, and tyrosinase inhibitory constituents of *Morus yunnanensis*. *J Nat Prod* 2012;75:82–7.

231. Fawole OA, Makunga NP, Opara UL. Antibacterial, antioxidant and tyrosinase-inhibition activities of pomegranate fruit peel methanolic extract. *BMC Complement Altern Med* 2012;12:200.
232. Kim JM, Ko RK, Jung DS, et al. Tyrosinase inhibitory constituents from the stems of *Maackia fauriei*. *Phytother Res* 2010;24:70–5.
233. Alam N, Yoon KN, Lee KR, et al. Antioxidant activities and tyrosinase inhibitory effects of different extracts from *Pleurotus ostreatus* fruiting bodies. *Mycobiology* 2010;38:295–301.
234. Piao X, Tian Y, Mi X, Cui J. tyrosinase inhibition of *Potentilla bifurca*. *Zhongguo Zhong Yao Za Zhi* 2009;34:1952–4.
235. Lu YH, Lin T, Wang ZT, et al. Mechanism and inhibitory effect of galangin and its flavonoid mixture from *Alpinia officinarum* on mushroom tyrosinase and B16 murine melanoma cells. *J Enzyme Inhib Med Chem* 2007;22:433–8.
236. Jeong SH, Ryu YB, Curtis-Long MJ, et al. Tyrosinase inhibitory polyphenols from roots of *Morus lhou*. *J Agric Food Chem* 2009;57:1195–203.
237. Kishore N, Twilley D, Blom van Staden A, et al. Isolation of flavonoids and flavonoid glycosides from *Myrsine africana* and their inhibitory activities against mushroom tyrosinase. *J Nat Prod* 2018;81:49–56.
238. Lee HS. Tyrosinase inhibitors of *Pulsatilla cernua* root-derived materials. *J Agric Food Chem* 2002;50:1400–3.
239. Wang YL, Hu G, Zhang Q, et al. Screening and characterizing tyrosinase inhibitors from *Salvia miltiorrhiza* and *Carthamus tinctorius* by spectrum-effect relationship analysis and molecular docking. *J Anal Methods Chem* 2018;2018:1.
240. Azizuddin, Khan AM, Choudhary MI. Tyrosinase inhibitory potential of natural products isolated from various medicinal plants. *Nat Prod Res* 2011;25:750–3.
241. Falcone Ferreyra ML, Rius SP, Casati P. Flavonoids: biosynthesis, biological functions, and biotechnological applications. *Front Plant Sci* 2012;3:222.
242. Lee NK, Son KH, Chang HW, et al. Prenylated flavonoids as tyrosinase inhibitors. *Arch Pharm Res* 2004;27:1132–5.
243. Rho HS, Ahn SM, Lee BC, et al. Changes in flavonoid content and tyrosinase inhibitory activity in kenaf leaf extract after far-infrared treatment. *Bioorg Med Chem Lett* 2010;20:7534–6.
244. Manthey JA, Cesar TB, Jackson E, Mertens-Talcott S. Pharmacokinetic study of nobiletin and tangeretin in rat serum by high-performance liquid chromatography-electrospray ionization-mass spectrometry. *J Agric Food Chem* 2011;59:145–51.
245. Shang C, Zhang Y, You X, et al. The effect of 7,8,4-trihydroxyflavone on tyrosinase activity and conformation: spectroscopy and docking studies. *Luminescence* 2018;33:681–91.
246. Gao H, Nishida J, Saito S, Kawabata J. Inhibitory effects of 5,6,7-trihydroxyflavones on tyrosinase. *Molecules* 2007;12:86–97.
247. Mu Y, Li L, Hu SQ. Molecular inhibitory mechanism of tricetin on tyrosinase. *Spectrochim Acta A Mol Biomol Spectrosc* 2013;107:235–40.
248. Guo N, Wang C, Shang C, et al. Integrated study of the mechanism of tyrosinase inhibition by baicalein using kinetic, multispectroscopic and computational simulation analyses. *Int J Biol Macromol* 2018;118:57–68.
249. Zhang L, Tao G, Chen J, Zheng ZP. Characterization of a new flavone and tyrosinase inhibition constituents from the Twigs of *Morus alba* L. *Molecules* 2016;21(9):1130. doi: 10.3390/molecules21091130
250. Ryu YB, Ha TJ, Curtis-Long MJ, et al. Inhibitory effects on mushroom tyrosinase by flavones from the stem barks of *Morus lhou* (S.) Koidz. *J Enzyme Inhib Med Chem* 2008;23:922–30.
251. Kubo I, Kinoshita H. Flavonols from saffron flower: tyrosinase inhibitory activity and inhibition mechanism. *J Agric Food Chem* 1999;47:4121–5.
252. Omar SH, Scott CJ, Hamlin AS, Obied HK. Biophenols: enzymes (β -secretase, cholinesterases, histone deacetylase and tyrosinase) inhibitors from olive (*Olea europaea* L.). *Fitoterapia* 2018;128:118–29.
253. Yang Z, Zhang Y, Sun L, et al. An ultrafiltration high-performance liquid chromatography coupled with diode array detector and mass spectrometry approach for screening and characterising tyrosinase inhibitors from mulberry leaves. *Anal Chim Acta* 2012;719:87–95.
254. Wang Y, Zhang G, Yan J, Gong D. Inhibitory effect of morin on tyrosinase: insights from spectroscopic and molecular docking studies. *Food Chem* 2014;163:226–33.
255. Zheng ZP, Zhu Q, Fan CL, et al. Phenolic tyrosinase inhibitors from the stems of *Cudrania cochinchinensis*. *Food Funct* 2011;2:259–64.
256. Kim JH, Cho IS, So YK, et al. Kushenol a and 8-prenylkaempferol, tyrosinase inhibitors, derived from *Sophora flavescens*. *J Enzyme Inhib Med Chem* 2018;33:1048–54.
257. Park JS, Kim DH, Lee JK, et al. Natural ortho-dihydroxyisoflavone derivatives from aged Korean fermented soybean paste as potent tyrosinase and melanin formation inhibitors. *Bioorg Med Chem Lett* 2010;20:1162–4.
258. Chang TS, Ding HY, Lin HC. Identifying 6,7,4'-trihydroxyisoflavone as a potent tyrosinase inhibitor. *Biosci Biotechnol Biochem* 2005;69:1999–2001.
259. Chang TS. Two potent suicide substrates of mushroom tyrosinase: 7,8,4'-trihydroxyisoflavone and 5,7,8,4'-tetrahydroxyisoflavone. *J Agric Food Chem* 2007;55:2010–5.
260. Chen J, Yu X, Huang Y. Inhibitory mechanisms of glabridin on tyrosinase. *Spectrochim Acta A Mol Biomol Spectrosc* 2016;168:111–7.
261. Deshmukh K, Poddar SS. Tyrosinase inhibitor-loaded micro-sponge drug delivery system: new approach for hyperpigmentation disorders. *J Microencapsul* 2012;29:559–68.
262. Jirawattanapong W, Saifah E, Patarapanich C. Synthesis of glabridin derivatives as tyrosinase inhibitors. *Arch Pharm Res* 2009;32:647–54.
263. Nerya O, Vaya J, Musa R, et al. Glabrene and isoliquiritigenin as tyrosinase inhibitors from licorice roots. *J Agric Food Chem* 2003;51:1201–7.
264. Heo do Y, Kim YM, Lee J, et al. Desmodianone H and uncianone B, potential tyrosinase inhibitors obtained from *Lespedeza maximowiczii* by using bioactivity-guided isolation. *Biosci Biotechnol Biochem* 2014;78:943–5.
265. Kim HJ, Seo SH, Lee BG, Lee YS. Identification of tyrosinase inhibitors from *Glycyrrhiza uralensis*. *Planta Med* 2005;71:785–7.
266. Kim JH, Kim HY, Kang SY, et al. Chemical constituents from *Apios americana* and their inhibitory activity on tyrosinase. *Molecules* 2018;23:232. doi: 10.3390/molecules23010232.

267. Si YX, Wang ZJ, Park D, et al. Effect of hesperetin on tyrosinase: inhibition kinetics integrated computational simulation study. *Int J Biol Macromol* 2012;50:257–62.
268. Chiari ME, Vera DM, Palacios SM, Carpinella MC. Tyrosinase inhibitory activity of a 6-isoprenoid-substituted flavanone isolated from *Dalea elegans*. *Bioorg Med Chem* 2011;19:3474–82.
269. Hu X, Yu MH, Yan GR, et al. Isoprenylated phenolic compounds with tyrosinase inhibition from *Morus nigra*. *J Asian Nat Prod Res* 2018;20:488–93.
270. Kim SS, Hyun CG, Choi YH, Lee NH. Tyrosinase inhibitory activities of the compounds isolated from *Neolitsea aciculata* (Blume) Koidz. *J Enzyme Inhib Med Chem* 2013;28:685–9.
271. Ko RK, Kim GO, Hyun CG, et al. Compounds with tyrosinase inhibition, elastase inhibition and DPPH radical scavenging activities from the branches of *Distylium racemosum* Sieb. et Zucc. *Phytother Res* 2011;25:1451–6.
272. Chai WM, Lin MZ, Wang YX, et al. Inhibition of tyrosinase by cherimoya pericarp proanthocyanidins: structural characterization, inhibitory activity and mechanism. *Food Res Int* 2017;100:731–9.
273. Kim YJ, Chung JE, Kurisawa M, et al. New tyrosinase inhibitors, (+)-catechin-aldehyde polycondensates. *Biomacromolecules* 2004;5:474–9.
274. Chai WM, Huang Q, Lin MZ, et al. Condensed tannins from longan bark as inhibitor of tyrosinase: structure, activity, and mechanism. *J Agric Food Chem* 2018;66:908–17.
275. Jhan JK, Chung YC, Chen GH, et al. Anthocyanin contents in the seed coat of black soya bean and their anti-human tyrosinase activity and antioxidative activity. *Int J Cosmet Sci* 2016;38:319–24.
276. Bukhari SN, Jantan I, Unsal Tan O, et al. Biological activity and molecular docking studies of curcumin-related α,β -unsaturated carbonyl-based synthetic compounds as anticancer agents and mushroom tyrosinase inhibitors. *J Agric Food Chem* 2014;62:5538–47.
277. Liu J, Wu F, Chen L, et al. Biological evaluation of coumarin derivatives as mushroom tyrosinase inhibitors. *Food Chem* 2012;135:2872–8.
278. Le-Thi-Thu H, Casanola-Martin GM, Marrero-Ponce Y, et al. Novel coumarin-based tyrosinase inhibitors discovered by OECD principles-validated QSAR approach from an enlarged, balanced database. *Mol Divers* 2011;15:507–20.
279. Hassan M, Ashraf Z, Abbas Q, et al. Exploration of novel human tyrosinase inhibitors by molecular modeling, docking and simulation studies. *Interdiscip Sci* 2018;10:68–80.
280. Pintus F, Matos MJ, Vilar S, et al. New insights into highly potent tyrosinase inhibitors based on 3-heteroarylcoumarins: anti-melanogenesis and antioxidant activities, and computational molecular modeling studies. *Bioorg Med Chem* 2017;25:1687–95.
281. Masamoto Y, Ando H, Murata Y, et al. Mushroom tyrosinase inhibitory activity of esuletin isolated from seeds of *Euphorbia lathyris* L. *Biosci Biotechnol Biochem* 2003;67:631–4.
282. Ahmad VU, Ullah F, Hussain J, et al. Tyrosinase inhibitors from *Rhododendron collettianum* and their structure–activity relationship (SAR) studies. *Chem Pharm Bull (Tokyo)* 2004;52:1458–61.
283. Ashraf Z, Rafiq M, Seo SY, et al. Design, synthesis and bioevaluation of novel umbelliferone analogues as potential mushroom tyrosinase inhibitors. *J Enzyme Inhib Med Chem* 2015;30:874–83.
284. Matos MJ, Santana L, Uriarte E, et al. New halogenated phenylcoumarins as tyrosinase inhibitors. *Bioorg Med Chem Lett* 2011;21:3342–5.
285. Ueno T, Fukami H, Ohkishi H, et al. Synthesis of 3,4-dihydro-3-amino-7-hydroxycoumarin from *s*-methyl-l-cysteine and resorcinol by crystalline- β -tyrosinase. *Biochim Biophys Acta* 1970;206:476–9.
286. Asthana S, Zucca P, Vargiu AV, et al. Structure–activity relationship study of hydroxycoumarins and mushroom tyrosinase. *J Agric Food Chem* 2015;63:7236–44.
287. Gardelly M, Trimech B, Belkacem MA, et al. Synthesis of novel diazaphosphinanes coumarin derivatives with promoted cytotoxic and anti-tyrosinase activities. *Bioorg Med Chem Lett* 2016;26:2450–4.
288. Tocco G, Fais A, Meli G, et al. PEG-immobilization of cardol and soluble polymer-supported synthesis of some cardol-coumarin derivatives: preliminary evaluation of their inhibitory activity on mushroom tyrosinase. *Bioorg Med Chem Lett* 2009;19:36–9.
289. Fais A, Corda M, Era B, et al. Tyrosinase inhibitor activity of coumarin-resveratrol hybrids. *Molecules* 2009;14:2514–20.
290. Yang HH, Oh KE, Jo YH, et al. Characterization of tyrosinase inhibitory constituents from the aerial parts of *Humulus japonicus* using LC-MS/MS coupled online assay. *Bioorg Med Chem* 2018;26:509–15.
291. Takahashi M, Takara K, Toyozato T, Wada K. A novel bioactive chalcone of *Morus australis* inhibits tyrosinase activity and melanin biosynthesis in B16 melanoma cells. *J Oleo Sci* 2012;61:585–92.
292. Zhang X, Hu X, Hou A, Wang H. Inhibitory effect of 2,4,2',4'-tetrahydroxy-3-(3-methyl-2-butenyl)-chalcone on tyrosinase activity and melanin biosynthesis. *Biol Pharm Bull* 2009;32:86–90.
293. Niesen DB, Ma H, Yuan T, et al. Phenolic constituents of *Carex vulpinoidea* seeds and their tyrosinase inhibitory activities. *Nat Prod Commun* 2015;10:491–3.
294. Suthar SK, Bansal S, Narkhede N, et al. Design, synthesis and biological evaluation of oxindole-based chalcones as small-molecule inhibitors of melanogenic tyrosinase. *Chem Pharm Bull (Tokyo)* 2017;65:833–9.
295. Kim BH, Park KC, Park JH, et al. Inhibition of tyrosinase activity and melanin production by the chalcone derivative 1-(2-cyclohexylmethoxy-6-hydroxy-phenyl)-3-(4-hydroxymethyl-phenyl)-propenone. *Biochem Biophys Res Commun* 2016;480:648–54.
296. Niu C, Yin L, Nie LF, et al. Synthesis and bioactivity of novel isoxazole chalcone derivatives on tyrosinase and melanin synthesis in murine B16 cells for the treatment of vitiligo. *Bioorg Med Chem* 2016;24:5440–8.
297. Radhakrishnan SK, Shimmon RG, Conn C, Baker AT. Evaluation of novel chalcone oximes as inhibitors of tyrosinase and melanin formation in B16 cells. *Arch Pharm (Weinheim)* 2016;349:20–9.
298. Radhakrishnan SK, Shimmon RG, Conn C, Baker AT. Inhibitory kinetics of azachalcones and their oximes on mushroom tyrosinase: a facile solid-state synthesis. *Chem Biodivers* 2016;13:531–8.
299. Dong X, Zhang Y, He JL, et al. Preparation of tyrosinase inhibitors and antibrowning agents using green technology. *Food Chem* 2016;197:589–96.

300. Jun N, Hong G, Jun K. Synthesis and evaluation of 2',4',6'-trihydroxychalcones as a new class of tyrosinase inhibitors. *Bioorg Med Chem* 2007;15:2396–402.
301. Radhakrishnan S, Shimmon R, Conn C, Baker A. Integrated kinetic studies and computational analysis on naphthyl chalcones as mushroom tyrosinase inhibitors. *Bioorg Med Chem Lett* 2015;25:4085–91.
302. Liu J, Chen C, Wu F, Zhao L. Microwave-assisted synthesis and tyrosinase inhibitory activity of chalcone derivatives. *Chem Biol Drug Des* 2013;82:39–47.
303. Nerya O, Musa R, Khatib S, et al. Chalcones as potent tyrosinase inhibitors: the effect of hydroxyl positions and numbers. *Phytochemistry* 2004;65:1389–95.
304. Okombi S, Rival D, Bonnet S, et al. Discovery of benzylidenebenzofuran-3(2H)-one (aurones) as inhibitors of tyrosinase derived from human melanocytes. *J Med Chem* 2006;49:329–33.
305. Zhu JJ, Yan GR, Xu ZJ, et al. Inhibitory effects of (2'R)-2',3'-dihydro-2'-(1-hydroxy-1-methylethyl)-2,6'-bibenzofuran-6,4'-diol on mushroom tyrosinase and melanogenesis in B16-F10 melanoma cells. *Phytother Res* 2015;29:1040–5.
306. Hu X, Wang M, Yan GR, et al. 2-Arylbzofuran and tyrosinase inhibitory constituents of *Morus notabilis*. *J Asian Nat Prod Res* 2012;14:1103–8.
307. Koirala P, Seong SH, Zhou Y, et al. Structure(–)activity relationship of the tyrosinase inhibitors kuwanon G, mulberrofuran G, and albanol B from *Morus* species: a kinetics and molecular docking study. *Molecules* 2018;23:1413. doi: [10.3390/molecules23061413](https://doi.org/10.3390/molecules23061413).
308. Wang Y, Xu L, Gao W, et al. Isoprenylated phenolic compounds from *Morus macroura* as potent tyrosinase inhibitors. *Planta Med* 2018;84:336–43.
309. Lin YF, Hu YH, Lin HT, et al. Inhibitory effects of propyl gallate on tyrosinase and its application in controlling pericarp browning of harvested longan fruits. *J Agric Food Chem* 2013;61:2889–95.
310. Lopes TIB, Coelho RG, Honda NK. Inhibition of mushroom tyrosinase activity by orsellinates. *Chem Pharm Bull (Tokyo)* 2018;66:61–4.
311. Lim JY, Ishiguro K, Kubo I. Tyrosinase inhibitory *p*-coumaric acid from ginseng leaves. *Phytother Res* 1999;13:371–5.
312. Cabanes J, Garcia-Carmona F, Garcia-Canovas F, et al. Kinetic study on the slow inhibition of epidermis tyrosinase by *m*-coumaric acid. *Biochim Biophys Acta* 1984;790:101–7.
313. An SM, Koh JS, Boo YC. *p*-Coumaric acid not only inhibits human tyrosinase activity in vitro but also melanogenesis in cells exposed to UVB. *Phytother Res* 2010;24:1175–80.
314. Garcia-Jimenez A, Teruel-Puche JA, Garcia-Ruiz PA, et al. Action of tyrosinase on caffeic acid and its *n*-nonyl ester. Catalysis and suicide inactivation. *Int J Biol Macromol* 2018;107:2650–9.
315. Hu YH, Chen QX, Cui Y, et al. 4-Hydroxy cinnamic acid as mushroom preservation: anti-tyrosinase activity kinetics and application. *Int J Biol Macromol* 2016;86:489–95.
316. Kwak SY, Yang JK, Choi HR, et al. Synthesis and dual biological effects of hydroxycinnamoyl phenylalanyl/prolyl hydroxamic acid derivatives as tyrosinase inhibitor and antioxidant. *Bioorg Med Chem Lett* 2013;23:1136–42.
317. Iwai K, Kishimoto N, Kakino Y, et al. In vitro antioxidative effects and tyrosinase inhibitory activities of seven hydroxycinnamoyl derivatives in green coffee beans. *J Agric Food Chem* 2004;52:4893–8.
318. Oyama T, Takahashi S, Yoshimori A, et al. Discovery of a new type of scaffold for the creation of novel tyrosinase inhibitors. *Bioorg Med Chem* 2016;24:4509–15.
319. Bernard P, Berthon JY. Resveratrol: an original mechanism on tyrosinase inhibition. *Int J Cosmet Sci* 2000;22:219–26.
320. Park J, Boo YC. Isolation of resveratrol from *Vitis Viniferae caulis* and its potent inhibition of human tyrosinase. *Evid Based Complement Alternat Med* 2013;2013:645257.
321. Gilly R, Mara D, Oded S, Zohar K. Resveratrol and a novel tyrosinase in carignan grape juice. *J Agric Food Chem* 2001;49:1479–85.
322. Likhitwitayawuid K, Sritularak B, De-Eknamkul W. Tyrosinase inhibitors from *Artocarpus gomezianus*. *Planta Med* 2000;66:275–7.
323. Lee N, Kim EJ, Kim BG. Regioselective hydroxylation of trans-resveratrol via inhibition of tyrosinase from *Streptomyces avermitilis* MA4680. *ACS Chem Biol* 2012;7:1687–92.
324. Ohguchi K, Tanaka T, Ito T, et al. Inhibitory effects of resveratrol derivatives from dipterocarpaceae plants on tyrosinase activity. *Biosci Biotechnol Biochem* 2003;67:1587–9.
325. Franco DC, de Carvalho GS, Rocha PR, et al. Inhibitory effects of resveratrol analogs on mushroom tyrosinase activity. *Molecules* 2012;17:11816–25.
326. Shin NH, Ryu SY, Choi EJ, et al. Oxyresveratrol as the potent inhibitor on DOPA oxidase activity of mushroom tyrosinase. *Biochem Biophys Res Commun* 1998;243:801–3.
327. Zheng ZP, Tan HY, Wang M. Tyrosinase inhibition constituents from the roots of *Morus australis*. *Fitoterapia* 2012;83:1008–13.
328. Bae SJ, Ha YM, Kim JA, et al. A novel synthesized tyrosinase inhibitor: (E)-2-((2,4-dihydroxyphenyl) diazenyl) phenyl 4-methylbenzenesulfonate as an azo-resveratrol analog. *Biosci Biotechnol Biochem* 2013;77:65–72.
329. Song YM, Ha YM, Kim JA, et al. Synthesis of novel azo-resveratrol, azo-oxyresveratrol and their derivatives as potent tyrosinase inhibitors. *Bioorg Med Chem Lett* 2012;22:7451–5.
330. Yanagihara M, Yoshimatsu M, Inoue A, et al. Inhibitory effect of gnetin c, a resveratrol dimer from melinjo (*Gnetum gnemon*), on tyrosinase activity and melanin biosynthesis. *Biol Pharm Bull* 2012;35:993–6.
331. Kim YM, Yun J, Lee CK, et al. Oxyresveratrol and hydroxystilbene compounds. Inhibitory effect on tyrosinase and mechanism of action. *J Biol Chem* 2002;277:16340–4.
332. Ohguchi K, Tanaka T, Kido T, et al. Effects of hydroxystilbene derivatives on tyrosinase activity. *Biochem Biophys Res Commun* 2003;307:861–3.
333. Kim DH, Kim JH, Baek SH, et al. Enhancement of tyrosinase inhibition of the extract of *Veratrum patulum* using cellulase. *Biotechnol Bioeng* 2004;87:849–54.
334. Uesugi D, Hamada H, Shimoda K, et al. Synthesis, oxygen radical absorbance capacity, and tyrosinase inhibitory activity of glycosides of resveratrol, pterostilbene, and pinostilbene. *Biosci Biotechnol Biochem* 2017;81:226–30.
335. Ismail T, Shafi S, Srinivas J, et al. Synthesis and tyrosinase inhibition activity of trans-stilbene derivatives. *Bioorg Chem* 2016;64:97–102.
336. Lima LL, Lima RM, da Silva AF, et al. Azastilbene analogs as tyrosinase inhibitors: new molecules with depigmenting potential. *Scientific World Journal* 2013;2013:274643. doi: [10.1155/2013/274643](https://doi.org/10.1155/2013/274643).

337. Choi J, Bae SJ, Ha YM, et al. A newly synthesized, potent tyrosinase inhibitor: 5-(6-hydroxy-2-naphthyl)-1,2,3-benzenetriol. *Bioorg Med Chem Lett* 2010;20:4882–4.
338. Song S, Lee H, Jin Y, et al. Syntheses of hydroxy substituted 2-phenyl-naphthalenes as inhibitors of tyrosinase. *Bioorg Med Chem Lett* 2007;17:461–4.
339. Ha YM, Chung SW, Song S, et al. 4-(6-Hydroxy-2-naphthyl)-1,3-benzenediol: a potent, new tyrosinase inhibitor. *Biol Pharm Bull* 2007;30:1711–5.
340. Satooka H, Kubo I. Resveratrol as a kcat type inhibitor for tyrosinase: potentiated melanogenesis inhibitor. *Bioorg Med Chem* 2012;20:1090–9.
341. Fachinetti N, Rigon RB, Eloy JO, et al. Comparative study of glyceryl behenate or polyoxyethylene 40 stearate-based lipid carriers for trans-resveratrol delivery: development, characterization and evaluation of the in vitro tyrosinase inhibition. *AAPS PharmSciTech* 2018;19:1401–9.
342. Wu B, Zhang X, Wu X. New lignan glucosides with tyrosinase inhibitory activities from exocarp of *Castanea henryi*. *Carbohydr Res* 2012;355:45–9.
343. Karioti A, Protopappa A, Megoulas N, Skaltsa H. Identification of tyrosinase inhibitors from *Marrubium velutinum* and *Marrubium cylleneum*. *Bioorg Med Chem* 2007;15:2708–14.
344. Wu YY, Huang XX, Wu J, et al. A new cyclolignan glycoside from the tubers of *Pinellia ternata*. *J Asian Nat Prod Res* 2015;17:1097–103.
345. Huang XX, Liu QB, Wu J, et al. Antioxidant and tyrosinase inhibitory effects of neolignan glycosides from *Crataegus pinnatifida* seeds. *Planta Med* 2014;80:1732–8.
346. Ashraf Z, Rafiq M, Nadeem H, et al. Carvacrol derivatives as mushroom tyrosinase inhibitors; synthesis, kinetics mechanism and molecular docking studies. *PLoS One* 2017; 12:e0178069.
347. Mutschlechner B, Rainer B, Schwaiger S, Stuppner H. Tyrosinase inhibitors from the aerial parts of *Wulfenia carinthiaca* Jacq. *Chem Biodivers* 2018;15:e1800014
348. Lehbili M, Alabdul Magid A, Hubert J, et al. Two new bis-iridoids isolated from *Scabiosa stellata* and their antibacterial, antioxidant, anti-tyrosinase and cytotoxic activities. *Fitoterapia* 2018;125:41–8.
349. Tan C, Zhu W, Lu Y. Aloin, cinnamic acid and sophorcarpine are potent inhibitors of tyrosinase. *Chin Med J (Engl)* 2002;115:1859–62.
350. Leu YL, Hwang TL, Hu JW, Fang JY. Anthraquinones from *Polygonum cuspidatum* as tyrosinase inhibitors for dermal use. *Phytother Res* 2008;22:552–6.
351. Bao K, Dai Y, Zhu ZB, et al. Design and synthesis of biphenyl derivatives as mushroom tyrosinase inhibitors. *Bioorg Med Chem* 2010;18:6708–14.
352. Kim YJ, No JK, Lee JH, Chung HY. 4,4'-Dihydroxybiphenyl as a new potent tyrosinase inhibitor. *Biol Pharm Bull* 2005; 28:323–7.
353. van Rensburg WJ, Ferreira D, Malan E, Steenkamp JA. Tyrosinase catalysed biphenyl construction from flavan-3-ol substrates. *Phytochemistry* 2000;53:285–92.
354. Oyama T, Yoshimori A, Takahashi S, et al. Structural insight into the active site of mushroom tyrosinase using phenylbenzoic acid derivatives. *Bioorg Med Chem Lett* 2017;27: 2868–72.
355. Le Mellay-Hamon V, Criton M. Phenylethylamide and phenylmethylamide derivatives as new tyrosinase inhibitors. *Biol Pharm Bull* 2009;32:301–3.
356. Chiku K, Dohi H, Saito A, et al. Enzymatic synthesis of 4-hydroxyphenyl beta-D-oligoxylosides and their notable tyrosinase inhibitory activity. *Biosci Biotechnol Biochem* 2009;73:1123–8.
357. Saboury AA, Zolghadri S, Haghbeen K, Moosavi-Movahedi AA. The inhibitory effect of benzenethiol on the cresolase and catecholase activities of mushroom tyrosinase. *J Enzyme Inhib Med Chem* 2006;21:711–7.
358. Alijanianzadeh M, Saboury AA, Ganjali MR, et al. Inhibition of mushroom tyrosinase by a newly synthesized ligand: inhibition kinetics and computational simulations. *J Biomol Struct Dyn* 2012;30:448–59.
359. Shareefi Borojerdi S, Haghbeen K, Asghar Karkhane A, et al. Successful resonance Raman study of cresolase activity of mushroom tyrosinase. *Biochem Biophys Res Commun* 2004;314:925–30.
360. Shao LL, Wang XL, Chen K, et al. Novel hydroxypyridinone derivatives containing an oxime ether moiety: synthesis, inhibition on mushroom tyrosinase and application in anti-browning of fresh-cut apples. *Food Chem* 2018;242:174–81.
361. Saghaie L, Pourfarzam M, Fassihi A, Sartippour B. Synthesis and tyrosinase inhibitory properties of some novel derivatives of kojic acid. *Res Pharm Sci* 2013;8:233–42.
362. Li DF, Hu PP, Liu MS, et al. Design and synthesis of hydroxypyridinone-l-phenylalanine conjugates as potential tyrosinase inhibitors. *J Agric Food Chem* 2013;61:6597–603.
363. Hider RC, Lerch K. The inhibition of tyrosinase by pyridinones. *Biochem J* 1989;257:289–90.
364. Dong H, Liu J, Liu X, et al. Molecular docking and QSAR analyses of aromatic heterocycle thiosemicarbazone analogues for finding novel tyrosinase inhibitors. *Bioorg Chem* 2017;75:106–17.
365. Song S, You A, Chen Z, et al. Study on the design, synthesis and structure-activity relationships of new thiosemicarbazone compounds as tyrosinase inhibitors. *Eur J Med Chem* 2017;139:815–25.
366. Xie J, Dong H, Yu Y, Cao S. Inhibitory effect of synthetic aromatic heterocycle thiosemicarbazone derivatives on mushroom tyrosinase: insights from fluorescence, (1)H-NMR titration and molecular docking studies. *Food Chem* 2016; 190:709–16.
367. Zhu TH, Cao SW, Yu YY. Synthesis, characterization and biological evaluation of paeonol thiosemicarbazone analogues as mushroom tyrosinase inhibitors. *Int J Biol Macromol* 2013;62:589–95.
368. Xu J, Liu J, Zhu X, et al. Novel inhibitors of tyrosinase produced by the 4-substitution of TCT. *Food Chem* 2017; 221:1530–8.
369. Choi J, Choi KE, Park SJ, et al. Ensemble-based virtual screening led to the discovery of new classes of potent tyrosinase inhibitors. *J Chem Inf Model* 2016;56:354–67.
370. Yi W, Cao R, Chen Z, et al. Rational design and synthesis of 4-o-substituted phenylmethylenethiosemicarbazones as novel tyrosinase inhibitors. *Chem Pharm Bull (Tokyo)* 2010; 58:752–4.
371. Yi W, Cao RH, Chen ZY, et al. Design, synthesis and biological evaluation of hydroxy- or methoxy-substituted phenylmethylenethiosemicarbazones as tyrosinase inhibitors. *Chem Pharm Bull (Tokyo)* 2009;57:1273–7.
372. Chen LH, Hu YH, Song W, et al. Synthesis and antityrosinase mechanism of benzaldehyde thiosemicarbazones: novel tyrosinase inhibitors. *J Agric Food Chem* 2012;60: 1542–7.

373. Li ZC, Chen LH, Yu XJ, et al. Inhibition kinetics of chlorobenzaldehyde thiosemicarbazones on mushroom tyrosinase. *J Agric Food Chem* 2010;58:12537–40.
374. El-Sadek MM, Hassan SY, Abdelwahab HE, Yacout GA. Synthesis of new 1,3,4-thiadiazole and 1,2,3,4-oxathiadiazole derivatives from carbohydrate precursors and study of their effect on tyrosinase enzyme. *Molecules* 2012;17:8378–96.
375. Yi W, Dubois C, Yahiaoui S, et al. Refinement of arylthiosemicarbazone pharmacophore in inhibition of mushroom tyrosinase. *Eur J Med Chem* 2011;46:4330–5.
376. Buitrago E, Vuillamy A, Boumendjel A, et al. Exploring the interaction of n/s compounds with a dicopper center: tyrosinase inhibition and model studies. *Inorg Chem* 2014;53:12848–58.
377. Yang MH, Chen CM, Hu YH, et al. Inhibitory kinetics of DABT and DABPT as novel tyrosinase inhibitors. *J Biosci Bioeng* 2013;115:514–7.
378. Liu J, Wu F, Chen C. Design and synthesis of aloe-emodin derivatives as potent anti-tyrosinase, antibacterial and anti-inflammatory agents. *Bioorg Med Chem Lett* 2015;25:5142–6.
379. You A, Zhou J, Song S, et al. Structure-based modification of 3-/4-aminoacetophenones giving a profound change of activity on tyrosinase: from potent activators to highly efficient inhibitors. *Eur J Med Chem* 2015;93:255–62.
380. Yi W, Cao R, Wen H, et al. Discovery of 4-functionalized phenyl-*o*-beta-d-glycosides as a new class of mushroom tyrosinase inhibitors. *Bioorg Med Chem Lett* 2009;19:6157–60.
381. Liu J, Yi W, Wan Y, et al. 1-(1-Arylethylidene) thiosemicarbazide derivatives: a new class of tyrosinase inhibitors. *Bioorg Med Chem* 2008;16:1096–102.
382. Yan Q, Cao R, Yi W, et al. Synthesis and evaluation of 5-benzylidene(thio)barbiturate-beta-d-glycosides as mushroom tyrosinase inhibitors. *Bioorg Med Chem Lett* 2009;19:4055–8.
383. Gheibi N, Saboury AA, Mansuri-Torshizi H, et al. The inhibition effect of some n-alkyl dithiocarbamates on mushroom tyrosinase. *J Enzyme Inhib Med Chem* 2005;20:393–9.
384. Amin E, Saboury AA, Mansouri-Torshizi H, et al. Evaluation of p-phenylene-bis and phenyl dithiocarbamate sodium salts as inhibitors of mushroom tyrosinase. *Acta Biochim Pol* 2010;57:277–83.
385. Amin E, Saboury AA, Mansuri-Torshizi H, Moosavi-Movahedi AA. Potent inhibitory effects of benzyl and p-xylylidine-bis dithiocarbamate sodium salts on activities of mushroom tyrosinase. *J Enzyme Inhib Med Chem* 2010;25:272–81.
386. Zarivi O, Bonfigli A, Cesare P, et al. Truffle thio-flavours reversibly inhibit truffle tyrosinase. *FEMS Microbiol Lett* 2003;220:81–8.
387. Choi J, Jee JG. Repositioning of thiourea-containing drugs as tyrosinase inhibitors. *Int J Mol Sci* 2015;16:28534–48.
388. De B, Adhikari I, Nandy A, et al. In silico modelling ofazole derivatives with tyrosinase inhibition ability: application of the models for activity prediction of new compounds. *Comput Biol Chem* 2018;74:105–14.
389. Loriga M, Paglietti G, Sparatore F, et al. Synthesis of substituted DL-3(5-benzazolyl)alanines as dopa and alpha-methyl-dopa analogs and their effects on dopamine beta-hydroxylase, tyrosinase and diphenoloxidase. *Farmacol* 1992;47:439–48.
390. Channar PA, Saeed A, Larik FA, et al. Synthesis of aryl pyrazole via suzuki coupling reaction, in vitro mushroom tyrosinase enzyme inhibition assay and in silico comparative molecular docking analysis with kojic acid. *Bioorg Chem* 2018;79:293–300.
391. Gawande SS, Warangkar SC, Bandgar BP, Khobragade CN. Synthesis of new heterocyclic hybrids based on pyrazole and thiazolidinone scaffolds as potent inhibitors of tyrosinase. *Bioorg Med Chem* 2013;21:2772–7.
392. Zhou Z, Zhuo J, Yan S, Ma L. Design and synthesis of 3,5-diaryl-4,5-dihydro-1H-pyrazoles as new tyrosinase inhibitors. *Bioorg Med Chem* 2013;21:2156–62.
393. Bandgar BP, Totre JV, Gawande SS, et al. Synthesis of novel 3,5-diaryl pyrazole derivatives using combinatorial chemistry as inhibitors of tyrosinase as well as potent anticancer, anti-inflammatory agents. *Bioorg Med Chem* 2010;18:6149–55.
394. Khan KM, Maharvi GM, Khan MT, et al. A facile and improved synthesis of sildenafil (viagra) analogs through solid support microwave irradiation possessing tyrosinase inhibitory potential, their conformational analysis and molecular dynamics simulation studies. *Mol Divers* 2005;9:15–26.
395. Mojzych M, Tarasiuk P, Kotwica-Mojzych K, et al. Synthesis of chiral pyrazolo[4,3-*e*][1,2,4]triazine sulfonamides with tyrosinase and urease inhibitory activity. *J Enzyme Inhib Med Chem* 2017;32:99–105.
396. Mojzych M, Dolashki A, Voelter W. Synthesis of pyrazolo[4,3-*e*][1,2,4]triazine sulfonamides, novel sildenafil analogs with tyrosinase inhibitory activity. *Bioorg Med Chem* 2014;22:6616–24.
397. Qamar R, Saeed A, Larik FA, et al. Novel 1,3-oxazine-tetrazole hybrids as mushroom tyrosinase inhibitors and free radical scavengers: synthesis, kinetic mechanism and molecular docking studies. *Chem Biol Drug Des* 2018; doi: [10.1111/cbdd.13352](https://doi.org/10.1111/cbdd.13352).
398. Nikalje APG, Gawhane P, Tiwari S, et al. Ultrasound promoted green synthesis, docking study of indole spliced thiadiazole, alpha-amino phosphonates as anticancer agents and anti-tyrosinase agents. *Anticancer Agents Med Chem* 2018;18:1. doi: [10.2174/1871520618666180417163226](https://doi.org/10.2174/1871520618666180417163226).
399. Mahdavi M, Ashtari A, Khoshneviszadeh M, et al. Synthesis of new benzimidazole-1,2,3-triazole hybrids as tyrosinase inhibitors. *Chem Biodivers* 2018; 15:e1800120.
400. Chekir S, Debbabi M, Regazzetti A, et al. Design, synthesis and biological evaluation of novel 1,2,3-triazole linked coumarinopyrazole conjugates as potent anticholinesterase, anti-5-lipoxygenase, anti-tyrosinase and anti-cancer agents. *Bioorg Chem* 2018;80:189–94.
401. Kim SJ, Yang J, Lee S, et al. The tyrosinase inhibitory effects of isoxazolone derivatives with a (z)-beta-phenyl-alpha, beta-unsaturated carbonyl scaffold. *Bioorg Med Chem* 2018; doi: [10.1016/j.bmc.2018.05.047](https://doi.org/10.1016/j.bmc.2018.05.047).
402. Hamidian H, Tagizadeh R, Fozooni S, et al. Synthesis of novel azo compounds containing 5(4h)-oxazolone ring as potent tyrosinase inhibitors. *Bioorg Med Chem* 2013;21:2088–92.
403. Heitz MP, Rupp JW. Determining mushroom tyrosinase inhibition by imidazolium ionic liquids: a spectroscopic and molecular docking study. *Int J Biol Macromol* 2018;107:1971–81.
404. Mann T, Scherner C, Rohm KH, Kolbe L. Structure–activity relationships of thiazolyl resorcinols, potent and selective

- inhibitors of human tyrosinase. *Int J Mol Sci* 2018;19:690. doi: [10.3390/ijms19030690](https://doi.org/10.3390/ijms19030690).
405. Rezaei M, Mohammadi HT, Mahdavi A, et al. Evaluation of thiazolidinone derivatives as a new class of mushroom tyrosinase inhibitors. *Int J Biol Macromol* 2018;108:205–13.
 406. Kim SH, Ha YM, Moon KM, et al. Anti-melanogenic effect of (z)-5-(2,4-dihydroxybenzylidene) thiazolidine-2,4-dione, a novel tyrosinase inhibitor. *Arch Pharm Res* 2013;36:1189–97.
 407. Ha YM, Park YJ, Kim JA, et al. Design and synthesis of 5-(substituted benzylidene)thiazolidine-2,4-dione derivatives as novel tyrosinase inhibitors. *Eur J Med Chem* 2012;49:245–52.
 408. Han YK, Park YJ, Ha YM, et al. Characterization of a novel tyrosinase inhibitor, (2RS,4R)-2-(2,4-dihydroxyphenyl)thiazolidine-4-carboxylic acid (MHY384). *Biochim Biophys Acta* 2012;1820:542–9.
 409. Ha YM, Park YJ, Lee JY, et al. Design, synthesis and biological evaluation of 2-(substituted phenyl)thiazolidine-4-carboxylic acid derivatives as novel tyrosinase inhibitors. *Biochimie* 2012;94:533–40.
 410. Jung HJ, Lee MJ, Park YJ, et al. A novel synthetic compound, (z)-5-(3-hydroxy-4-methoxybenzylidene)-2-iminothiazolidin-4-one (mhy773) inhibits mushroom tyrosinase. *Biosci Biotechnol Biochem* 2018;82:759–67. doi: [10.1080/09168451.2018.1445518](https://doi.org/10.1080/09168451.2018.1445518).
 411. Kahn V. Effect of kojic acid on the oxidation of DL-dopa, norepinephrine, and dopamine by mushroom tyrosinase. *Pigment Cell Res* 1995;8:234–40.
 412. Xie W, Zhang H, He J, et al. Synthesis and biological evaluation of novel hydroxybenzaldehyde-based kojic acid analogues as inhibitors of mushroom tyrosinase. *Bioorg Med Chem Lett* 2017;27:530–2.
 413. Chen MJ, Hung CC, Chen YR, et al. Novel synthetic kojic acid-methimazole derivatives inhibit mushroom tyrosinase and melanogenesis. *J Biosci Bioeng* 2016;122:666–72.
 414. Asadzadeh A, Sirous H, Pourfarzam M, et al. In vitro and in silico studies of the inhibitory effects of some novel kojic acid derivatives on tyrosinase enzyme. *Iran J Basic Med Sci* 2016;19:132–44.
 415. Xie W, Zhang J, Ma X, et al. Synthesis and biological evaluation of kojic acid derivatives containing 1,2,4-triazole as potent tyrosinase inhibitors. *Chem Biol Drug Des* 2015;86:1087–92.
 416. Lima CR, Silva JR, de Tassia Carvalho Cardoso E, et al. Combined kinetic studies and computational analysis on kojic acid analogous as tyrosinase inhibitors. *Molecules* 2014;19:9591–605.
 417. Noh JM, Kwak SY, Seo HS, et al. Kojic acid-amino acid conjugates as tyrosinase inhibitors. *Bioorg Med Chem Lett* 2009;19:5586–9.
 418. Lee YS, Park JH, Kim MH, et al. Synthesis of tyrosinase inhibitory kojic acid derivative. *Arch Pharm (Weinheim)* 2006;339:111–4.
 419. Cho JC, Rho HS, Joo YH, et al. Depigmenting activities of kojic acid derivatives without tyrosinase inhibitory activities. *Bioorg Med Chem Lett* 2012;22:4159–62.
 420. Nihei KI, Kubo I. Substituent effect of benzaldehydes on tyrosinase inhibition. *Plant Physiol Biochem* 2017;112:278–82.
 421. Rafiee M, Javaheri M. A theoretical study of benzaldehyde derivatives as tyrosinase inhibitors using ab initio calculated NQCC parameters. *Mol Biol Res Commun* 2015;4:151–9.
 422. Ley JP, Bertram HJ. Hydroxy- or methoxy-substituted benzaldoximes and benzaldehyde-*o*-alkyloximes as tyrosinase inhibitors. *Bioorg Med Chem* 2001;9:1879–85.
 423. Lopes ND, Chaves OA, de Oliveira MCC, et al. Novel piperonal 1,3,4-thiadiazolium-2-phenylamines mesoionic derivatives: synthesis, tyrosinase inhibition evaluation and HSA binding study. *Int J Biol Macromol* 2018;112:1062–72.
 424. Yi W, Cao R, Peng W, et al. Synthesis and biological evaluation of novel 4-hydroxybenzaldehyde derivatives as tyrosinase inhibitors. *Eur J Med Chem* 2010;45:639–46.
 425. Ha TJ, Tamura S, Kubo I. Effects of mushroom tyrosinase on anisaldehyde. *J Agric Food Chem* 2005;53:7024–8.
 426. Yu L. Inhibitory effects of (s)- and (r)-6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acids on tyrosinase activity. *J Agric Food Chem* 2003;51:2344–7.
 427. Alijanianzadeh M, Saboury AA, Mansuri-Torshizi H, et al. The inhibitory effect of some new synthesized xanthates on mushroom tyrosinase activities. *J Enzyme Inhib Med Chem* 2007;22:239–46.
 428. Saboury AA, Alijanianzadeh M, Mansoori-Torshizi H. The role of alkyl chain length in the inhibitory effect *n*-alkyl xanthates on mushroom tyrosinase activities. *Acta Biochim Pol* 2007;54:183–91.
 429. Xing R, Wang F, Zheng A, et al. Biological evaluation of two Keggin-type polyoxometalates containing glycine as mushroom tyrosinase inhibitors. *Biotechnol Appl Biochem* 2016;63:746–50.
 430. Park KH, Lee JR, Hahn HS, et al. Inhibitory effect of ammonium tetrathiotungstate on tyrosinase and its kinetic mechanism. *Chem Pharm Bull (Tokyo)* 2006;54:1266–70.
 431. Chen XX, Zhang J, Chai WM, et al. Reversible and competitive inhibitory kinetics of amoxicillin on mushroom tyrosinase. *Int J Biol Macromol* 2013;62:726–33.
 432. Hemachandran H, Jain F, Mohan S, et al. Glandular hair constituents of *Mallotus philippinensis* muell. Fruit act as tyrosinase inhibitors: insights from enzyme kinetics and simulation study. *Int J Biol Macromol* 2018;107:1675–82.
 433. Lin YF, Hu YH, Jia YL, et al. Inhibitory effects of naphthols on the activity of mushroom tyrosinase. *Int J Biol Macromol* 2012;51:32–6.
 434. Wu LC, Chen YC, Ho JA, Yang CS. Inhibitory effect of red koji extracts on mushroom tyrosinase. *J Agric Food Chem* 2003;51:4240–6.
 435. Fourche J, Jensen H, Neuzil E, Bellegarde B. [Alpha-hydrazinophloretic acid, competitive inhibitor of fungal tyrosinase]. *CR Hebd Seances Acad Sci Ser D* 1977;284:2163–6.
 436. Gheibi N, Saboury AA, Haghbeen K, Moosavi-Movahedi AA. Activity and structural changes of mushroom tyrosinase induced by *n*-alkyl sulfates. *Colloids Surf B Biointerfaces* 2005;45:104–7.
 437. Jena K, Pandey JP, Kumari R, et al. Tasar silk fiber waste sericin: new source for anti-elastase, anti-tyrosinase and anti-oxidant compounds. *Int J Biol Macromol* 2018;114:1102–8.
 438. Hwang SH, Wang Z, Suh HW, Lim SS. Antioxidant activity and inhibitory effects of 2-hydroxy-3-methylcyclopent-2-enone isolated from ribose-histidine maillard reaction products on aldose reductase and tyrosinase. *Food Funct* 2018;9:1790–9.
 439. Huang XX, Yan ZY, Liu S, et al. Investigation of chemical constituents of safflower and their tyrosinase inhibitory activity. *J Asian Nat Prod Res* 2018; 1–9. doi: [10.1080/10286020.2018.1430775](https://doi.org/10.1080/10286020.2018.1430775).

440. Aljanianzadeh M, Saboury AA, Ganjali MR, et al. The inhibitory effect of ethylenediamine on mushroom tyrosinase. *Int J Biol Macromol* 2012;50:573–7.
441. Wang Y, Hao MM, Sun Y, et al. Synergistic promotion on tyrosinase inhibition by antioxidants. *Molecules* 2018;23:106. doi: [10.3390/molecules23010106](https://doi.org/10.3390/molecules23010106).
442. Liang C, Lim JH, Kim SH, Kim DS. Dioscin: a synergistic tyrosinase inhibitor from the roots of *Smilax china*. *Food Chem* 2012;134:1146–8.
443. Jin YH, Lee SJ, Chung MH, et al. Aloesin and arbutin inhibit tyrosinase activity in a synergistic manner via a different action mechanism. *Arch Pharm Res* 1999;22:232–6.
444. Schved F, Kahn V. Synergism exerted by 4-methyl catechol, catechol, and their respective quinones on the rate of DL-dopa oxidation by mushroom tyrosinase. *Pigment Cell Res* 1992;5:41–8.
445. Chen X, Haniu A, Kashiwagi T, et al. The evaluation of the synergistic effect of 3-(2,4-dihydroxyphenyl)propionic acid and L-ascorbic acid on tyrosinase inhibition. *Z Naturforsch C* 2017;72:119–21.
446. Hseu YC, Cheng KC, Lin YC, et al. Synergistic effects of linderanolide B combined with arbutin, PTU or kojic acid on tyrosinase inhibition. *Curr Pharm Biotechnol* 2015;16:1120–6.