

Phytochemical and Pharmacological Review of Allium Species from Georgia

Giorgi Jgerenaia^{1,2, ID}, Michel Frederich^{2, ID}, Lasha Mskhiladze^{1, ID}

DOI: 10.52340/GBMN.2024.01.01.76

ABSTRACT

This research aims to provide a comprehensive phytochemical and pharmacological review of *Allium* species growing in Georgia. Utilizing scientific sources and articles detailing these plants' phytochemical constituents and pharmacological activities. The study identifies 36 *Allium* species in Georgia, including five endemics to the country and 2 to the Caucasus region. Numerous biologically active compounds have been isolated from these species, such as over 20 saponins, 11 types of sapogenins, flavonoids, and sulfur compounds. Extensive research highlights pharmacological activities, including cytotoxic, antioxidant, antimicrobial, hepatoprotective, and fibrinolytic effects. Notably, cytotoxic studies on *Allium* species show significant activity against several cancer cell lines. Antioxidant assays highlight the potential of *Allium* extracts in scavenging free radicals and reducing oxidative stress. Despite these findings, 22 of the 36 *Allium* species in Georgia still need to be studied, with unknown chemical structures and biological activities of their compounds. This review underscores the need for further detailed and extensive studies to enhance understanding of these lesser-known species' pharmacological potential and chemical composition

Keywords: Allium; pharmacological activities; phytochemistry.

INTRODUCTION

The *Allium* genus, part of the Alliaceae family, comprises around 800 species, predominantly found in the northern hemisphere.¹

These herbaceous plants have significant historical and traditional uses as food and medicine, with a cultivation history tracing over 4,000 years to ancient Egypt.²

In Georgian traditional medicine, *Allium* species are utilized for their antifungal, antiseptic, and antibacterial properties.^{3,4} Georgia claims a rich and diverse vegetation due to its varied and contrasting physical and geographical conditions, complex geological history, and strategic location at the crossroads of different botanical and geographical provinces.

The flora of the Caucasus, particularly in Georgia, is highly endemic. Approximately 21% of its flora (around 900 species) is unique to the region, including about 600 species endemics to the Caucasus and 300 specifically endemic to Georgia. The region has a notable endemic genus, with 17 endemic and subendemic genera.⁵

This review aims to provide detailed information on the phytochemistry and pharmacological activities of *Allium* species growing in Georgia.

REVIEW

Allium species in Georgia

According to R. Gagnidze's book "Vascular Plants of Georgia," 36 species of the genus *Allium* are described in Georgia. Among these, five species are endemic to Georgia, and two species are endemic to the Caucasus region (Tab.1).¹

TABLE 1. List of plants genus *Allium* described in Georgia

| | |
|----|--|
| 1 | <i>A. affine</i> Ledeb. |
| 2 | <i>A. albidum</i> Fisch. Ex Bieb |
| 3 | <i>A. albovianum</i> Vved |
| 4 | <i>A. atroviolaceum</i> Boiss |
| 5 | <i>A. aucheri</i> Boiss |
| 6 | <i>A. candolleianum</i> Albov |
| 7 | <i>A. cardiostemon</i> Fisch. Et C.A. Mey |
| 8 | <i>A. chevsuricum</i> Tscholok |
| 9 | <i>A. erubescens</i> K. Koch |
| 10 | <i>A. fuscoviolaceum</i> Fomin |
| 11 | <i>A. globosum</i> Bieb. Ex DC |
| 12 | <i>A. gracilescens</i> Somm. Et Levier |
| 13 | <i>A. gramineum</i> K. Koch |
| 14 | <i>A. karsianum</i> Fomin |
| 15 | <i>A. kunthianum</i> Vved. |
| 16 | <i>A. ledschanense</i> Conrath et Freyn |
| 17 | <i>A. leucanthum</i> K. Koch |
| 18 | <i>A. moschatum</i> L. |
| 19 | <i>A. oreophilum</i> C.A. Mey |
| 20 | <i>A. otschiauriae</i> Tscholok. |
| 21 | <i>A. paczoskianum</i> Tuzs. |
| 22 | <i>A. paradoxum</i> (Bieb.) G. Don. F (Scillaparadoxa Bieb.) |
| 23 | <i>A. ponticum</i> Miscz. Ex Grossh. |
| 24 | <i>A. pseudoflavum</i> Vved |
| 25 | <i>A. pseudostrictum</i> Albov |
| 26 | <i>A. rotundum</i> L. |
| 27 | <i>A. rubellum</i> Bieb |
| 28 | <i>A. rupestre</i> Stev |
| 29 | <i>A. ruprechtii</i> Boiss |
| 30 | <i>A. saxatile</i> Bieb |
| 31 | <i>A. scorodoprasum</i> L. (A. waldsteinii G. Don f. A. jajlae Vved) |
| 32 | <i>A. szovitsii</i> Regel |
| 33 | <i>A. tauricola</i> Bieb(A. subquinqueflorum Boiss) |
| 34 | <i>A. ursinum</i> L. |
| 35 | <i>A. victoralis</i> L. |
| 36 | <i>A. vineale</i> L. |



A. albivianum, *A. candolleianum*, *A. chevsuricum*, *A. gracilescens*, and *A. otschiauriae* are endemic species of Georgia; *A. ledschanense* and *A. leucanthum* are endemic species of the Caucasus region.

Phytochemistry of plants genus *Allium*, growing in Georgia

From *Allium* species growing in Georgia, the following compounds have been isolated: Kaempferol-(acetylhexoside)-hexoside, Malondialdehyde (MDA), (E)-Propenyl propyl disulfide, 3,4-Dihydro-3-vinyl-1,2-dithiine, Methyl-2-propenyl trisulfide, (Z)-Di-2-propenyl trisulfide, Cyclopent-1-enecarboxylate, Allicin, Yayoisaponin C, β -Amyrin acetate, Inulin, Diallyl disulfide, 22-Cyclohexyl-1-docosanol, Dideglucoeruboside B, Astragalinal, Allyl (methylthio)methyl disulfide, Nystose, 2-Vinyl-4H-1,3-dithiine, Atroviolaceoside, (E)-Di-2-propenyl trisulfide, Aginoside, Allivictoside A-H, Dimethyl disulfide, 2,4-Dimethylthiophene, Allyl propyl trisulfide, β -Sitosterol acetate, p-Vinylguaicol, Alliumonate, Allyl methyl disulfide, quercetin 3-O- β -glucopyranoside, Dimethyl tetrasulfide, Eruboside B, 2-Furaldehyde, Benzaldehyde, 2-Vinyl-1,3-dithiane, Allyl (Z)-1-propenyl disulfide, diosgenin-3-O- α -rhamnopyranosyl-(1 \rightarrow 2)- β -D-glucopyranoside (Prosapogenin A of dioscin), (Z)-Methyl-2-propenyl disulfide, β -Sitosterol 3-O- β -glucopyranoside, 4-Methyl-1,2,3-trithiolane, Leucospiroside A, Kaempferol-3,7-di-O- β -D-glucopyranoside, isorhamnetin 3,7-di-O- β -glucopyranoside, Deltonine, Dimethyl thiophene-(E)-methyl-2-propyl disulfide, β -Sitosterol, Acetyl-kaempferol-deoxyhexose propylene sulfide, Aginoside, Allumine A and B, 1-Propenyl propyl disulfide, 5-Methyl-1,2,3,4-tetrathiane, 4-Methyl-1,2,3,5,6-pentathiepane, Methyl propyl trisulfide, Allyl (E)-1-propenyl disulfide, Methyl (methylthio)methyl disulfide, Allyl (methylthio)methyl trisulfide, isorhamnetin-3-O- β -D-glucopyranoside, 2-Methoxy tyrosol, (Z)-Propenyl propyl disulfide, Ophiopogonin C, Allyl methyl trisulfide, Diallyl trisulfide, Methyl (Z)-1-propenyl disulfide, Dimethyl trisulfide, Trillin, Kaempferol-3,7-di-O- β -D-glucopyranoside, Diallyl

trisulfide, 2,4-Dimethylthiophene, Methyl (methylthio)methyl trisulfide, Methyl (E)-1-propenyl trisulfide, (Z)-Di-2-propenyl trisulfide, Methyl (Z)-1-propenyl disulfide, 4-Methyl-1,2,3-trithiolane, (E)-Propenyl propyl disulfide, quercetin 3-O- β -glucopyranoside, Cyclopent-1-enecarboxylate, Deltonine, Eruboside B, Ophiopogonin C, Allyl methyl disulfide, Acetyl-kaempferol-deoxyhexose propylene sulfide, Nystose, Allyl propyl trisulfide, Kaempferol-(acetylhexoside)-hexoside, β -Sitosterol acetate, (3Z)-Hexenol, Dideglucoeruboside B, β -Sitosterol 3-O- β -glucopyranoside, Trillin, 3,4-Dihydro-3-vinyl-1,2-dithiine, Alliumonate, Atroviolaceoside, Allyl (Z)-1-propenyl disulfide, 2-Vinyl-1,3-dithiane, Dimethyl trisulfide, Carotenoids, 22-Cyclohexyl-1-docosanol, Diallyl disulfide, Allivictoside A-H, Aginoside, diosgenin-3-O- α -rhamnopyranosyl-(1 \rightarrow 2)- β -D-glucopyranoside (Prosapogenin A of dioscin), Benzaldehyde, Methyl-2-propenyl trisulfide, Allumine A and B, Aginoside, Dimethyl disulfide, 2-Vinyl-4H-1,3-dithiine, (Z)-Methyl-2-propenyl disulfide, Allyl (methylthio)methyl disulfide, Allyl (E)-1-propenyl disulfide, β -Sitosterol, Astragalinal, Methyl propyl trisulfide, 1-Propenyl propyl disulfide, 2-Furaldehyde, (2E)-Hexenal, (Z)-Propenyl propyl disulfide, 5-Methyl-1,2,3,4-tetrathiane, (E)-Di-2-propenyl trisulfide, Yayoisaponin C, β -Amyrin acetate, (Z)-Methyl-2-propenyl disulfide, β -Amyrin, Methyl (E)-1-propenyl trisulfide, Allyl methyl tetrasulfide, 2-Methoxy tyrosol, Diallyl trisulfide, (2E)-Hexenal, Allicin, 4-Methyl-1,2,3,5,6-pentathiepane, Malondialdehyde (MDA), 4-Methyl-1,2,3-trithiolane, Leucospiroside A, (Z)-Propenyl propyl disulfide, Allyl methyl trisulfide, Dimethyl tetrasulfide, isorhamnetin 3,4'-di-O- β -glucopyranoside, Inulin, Dimethyl thiophene-(E)-methyl-2-propyl disulfide.

Different types of sapogenins have also been isolated from these species: β -chlorogenin, Diosgenin, Yucagenin, Tigogenin, Nuatigenin, Hecogenine, Agigenin, Ruscogenin, Atroviolacegenin, Gitogenin, and Isonuatigenin. Information about the biological activities and constituents of these plants is given in Table 2.

TABLE 2. Chemical compounds isolated from *Allium* species widespread in Georgia

| Species | Compounds | Aglycone | Biological activity |
|------------------------------|--|---|--|
| <i>Allium affine</i> | | Diosgenin, Tigogenin, Ruscogenin ⁶ | Antioxidant activity ⁷ Fibrinolytic activity ⁷ |
| <i>Allium albidum</i> | | Diosgenin, Ruscogenin ⁸ Hecogenine ⁹ | |
| <i>Allium atroviolaceum</i> | Atroviolaceoside ¹⁰ | Atroviolacegenin ¹⁰ | Antiplatelet activity ¹¹ Antibacterial activity ¹² Cytotoxic and antiproliferative effect ¹³⁻¹⁵ |
| <i>Allium erubescens</i> | Eruboside B ¹⁶ | β -chlorogenin ¹⁶ | |
| <i>Allium fuscoviolaceum</i> | | Diosgenin ¹⁷ | |
| <i>Allium gramineum</i> | isorhamnetin-3-O- β -D-glucopyranoside, diosgenin-3-O- α -rhamnopyranosyl-(1 \rightarrow 2)- β -D-glucopyranoside (Prosapogenin A of dioscin), Deltonine ¹⁸ Isorhamnetin 3,7-di-O- β -glucopyranoside, Quercetin 3-O- β -glucopyranoside, β -sitosterol 3-O- β -glucopyranoside, Isorhamnetin 3,4'-di-O- β -glucopyranoside, ¹⁹ Eruboside B ⁹ | | Cytotoxicity and Antioxidant activity ¹⁸ |

TABLE 2. Chemical compounds isolated from Allium species widespread in Georgia (continued)

| Species | Compounds | Aglycone | Biological activity |
|----------------------|---|--|---|
| Allium leucanthum | Yayoisaponin C, Eruboside B, Aginoside, Leucospiroside A, Three unknown saponins, ²⁰ (25R)-5 α -spirostane-3 β ,6 β -diol-O- β -D-glucopyranosyl-(1-2)-O- β -D-glucopyranosyl-(1-4)- β -D-galactopyranoside, Leucofuranoside A ²¹⁻²⁴ | Diosgenin β -chlorogenin Yucagenin, Agigenin ²⁴ | Cytotoxic activity ²⁰ Antifungal activity Antileishmanial activity ²⁵ Antibacterial activity ²⁶ |
| Allium paradoxum | Astragalgin; 2-Methoxy tyrosol ²⁷ | Dioscin related saponins ²⁸ | Leishmanicidal activity, ²⁸ Antioxidant activity, ²⁹ Hepatoprotective effect, ³⁰ Antiplasmodial activity, ³¹ Analgesic activity ³² Nephroprotective effect ³³ |
| Allium rotundum | Trillin; Dideglucoeruboside B, Aginoside, Eruboside B, Yayoisaponin C, ³⁴ Quercetin, Luteolin, Apigenin, Hyperin, Cinaroside, Apigenin-7-O- β -D-glucopyranoside, Scopoletin and Umbelliferone, ³⁵ β -Carotene, Violaxanthin, Flavoxanthin, Lutein, Rubixanthin, and Zeaxanthin ³⁶ | Tigogenin, Diosgenin, Gitogenin, β - Chlorogenin, Yucagenin, Agigenin ³⁷ | Antioxidant and Antihemolytic effects ³⁸ |
| Allium rubellum | | Tigogenin ³⁹ | Antioxidant activity ⁴⁰ |
| Allium scorodoprasum | | | Antioxidant activity ⁴¹⁻⁴³ Cytotoxicity, Antioxidant, Antimicrobial and anti-inflammatory activity ⁴⁴ |
| Allium ursinum | Dimethyl disulfide, Di-2-propenyl disulfide, (Z)-Di-2-propenyl trisulfide, Di-2-propyl trisulfide, Di-2-propenyl tetrasulfide, (E)-Di-2-propenyl trisulfide, Methyl-2-propenyl trisulfide, Dimethyl trisulfide, 2-Vinyl-1,3-dithiane, (Z)-Methyl-2-propenyl disulfide, (E)-Propenyl propyl disulfide, (Z)-Propenyl propyl disulfide, 3,4-Dihydro-3-vinyl-1,2-dithiine, 2-Vinyl-4H-1,3-dithiine, Dimethyl tetrasulfide, Carotenoids, Allicin, Malondialdehyde (MDA), Kaempferol-3,7-di-O- β -D-glucopyranoside, Kaempferol-(acetylhexoside)-hexoside, Acetyl-kaempferol-deoxyhexose propylene sulfide, I-Ketose, Nystose and Inulin ⁴⁵ | | Antioxidant and Anti-tyrosinase activity ^{46,47} Cytotoxicity ⁴⁸ Antifungal activity ⁴⁹ Antiplatelet activity ⁵⁰ |
| Allium victorialis | Allivictoside A-H ⁵¹ Allumine A and B, Cyclopent-1-enecarboxylate, ⁵² 22-Cyclohexyl-1-docosanol, Alliumonoate, β -Sitosterol acetate, β -Amyrin, β -Amyrin acetate, β -Sitosterol, β -Sitosterol 3-O-b-D-glucopyranoside ⁵³ | | Anti-neuroinflammatory effects ⁵¹ Chemopreventive and anticancer activities ⁵⁴ Anti-diabetic activity ⁵⁵ |
| Allium vineale | Deltonin, Ophiopogonin C and 7 other saponins ⁵⁶ Allyl methyl disulfide, 1-Propenyl propyl disulfide, Dimethyl trisulfide, 2-Furaldehyde, 2,4-Dimethylthiophene, Benzaldehyde, (2E)-Hexenal, (3Z)-Hexenol, Methyl (Z)-1-propenyl disulfide, Methyl (E)-1-propenyl disulfide, Diallyl disulfide, Allyl (Z)-1-propenyl disulfide, Allyl (E)-1-propenyl disulfide, Methyl (methylthio)methyl disulfide, Allyl methyl trisulfide, 4-Methyl-1,2,3-trithiolane, Methyl propyl trisulfide, Methyl (Z)-1-propenyl trisulfide, Methyl (E)-1-propenyl trisulfide, Dimethyl tetrasulfide, Allyl (methylthio)methyl disulfide, Diallyl trisulfide, Allyl (Z)-1-propenyl trisulfide, p-Vinylguaiacol, Allyl propyl trisulfide, 5-Methyl-1,2,3,4-tetrathiane, Methyl (methylthio)methyl trisulfide, 4-Methyl-1,2,3,5,6-pentathiepane, Allyl methyl tetrasulfide ⁵⁷ | Diosgenin, Nuatigenin and Isonuatigenin ⁵⁶ | Antioxidant activity ⁵⁷ |

Biological activity

Numerous pharmacological studies have revealed that Allium species exhibit a variety of exciting activities, including cytotoxic, antioxidant, thrombolytic, antibacterial, and other beneficial effects.

Cytotoxic activity

Numerous experimental articles have discussed the cytotoxic activity of fractions and compounds isolated from Allium species. Khazaei et al. evaluated the cytotoxic activity of bulbs of Allium atroviolaceum on various cancer cell lines, including MCF7, MDA-MB-231, HeLa, and HepG2. Using the MTT cytotoxicity assay, they observed different growth responses across these cell lines. The IC50 values were as follows after 24, 48, and 72 hours of treatment:

- MCF-7 cells: 91.5, 88, and 75.7 μ g/ml

- MDA-MB-231 cells: 149, 114, and 101 μ g/ml
- HeLa cells: 154, 89.7, and 74.7 μ g/ml
- HepG2 cells: 97, 70, and 58.7 μ g/ml.¹⁴

Additionally, the anticancer effect of 4',5,7-Trihydroxy-3',5'-dimethoxyflavone (Tricin), initially isolated from Allium atroviolaceum, was investigated in combination with Docetaxel on the PC3 cell line. The IC50 values for Tricin and Docetaxel were 117.5 \pm 4.4 μ M and 0.1 \pm 0.02 nM, respectively, as assessed by the MTT assay.¹⁵

The 80% ethanol extract of Allium gramineum flowers significantly inhibits the growth of breast adenocarcinoma cell lines, with IC50 values of 4.5 \pm 0.7 μ g/mL for MDAMB-231 and 4.8 \pm 0.9 μ g/mL for MCF-7 cells. This potent cytotoxicity is linked to saponins, which have an IC50 of about 3 μ M.¹⁸

Seven glycosides extracted from *Allium leucanthum* were tested for cytotoxicity against A549 and DLD-1 cell lines. Compounds 1-3 and 5 exhibited comparable cytotoxic effects on both cell lines, with IC50 values ranging from 5.6 to 8.2 μM for DLD-1 and 3.7 to 5.8 μM for A549.²⁰

Demir et al. (2022) found that the extract of *Allium scorodoprasum* exhibits cytotoxic activity, with IC50 values of 82.78 $\mu\text{g}/\text{mL}$ for MCF-7 cells and 76.53 $\mu\text{g}/\text{mL}$ for MG-63 cells.⁴⁴

Korga et al. (2019) evaluated the cytotoxicity of *Allium ursinum* extract on MKN28 and MKN74 cell lines, finding it toxic to both, with cell viability at 66.77 \pm 3.00% and 31.55 \pm 2.04%, respectively.⁴⁸

The apoptosis-inducing potential of these extracts was evaluated using the MTT assay, DAPI staining, and DNA fragmentation assay in human colon cancer HT-29 cells.⁵⁴

Antioxidant activity

Antioxidants play a crucial role in health by reducing disease risk and protecting the body against oxidative damage, which can lead to conditions such as diabetes, cancer, and neurodegenerative disorders. They help control oxidative processes that can deteriorate food quality due to reactive oxygen species (ROS) and free radical reactions.⁴⁵

Antioxidants can naturally occur in plants, animals, and microorganisms, or they can be chemically synthesized. Higher plants are particularly rich in natural antioxidants like tocopherols and polyphenols, which are found in abundance in spices, herbs, fruits, vegetables, cereals, grains, seeds, teas, and oils.⁵⁸

Numerous researches have been done on *Allium* species growing in Georgia to evaluate their antioxidant activity.

The DPPH scavenging test was used to evaluate the free radical scavenging activity of *Allium affine* hydroalcoholic extract, with an RC50 of 201 $\mu\text{g}/\text{mL}$, compared to 43 $\mu\text{g}/\text{mL}$ for vitamin C.⁷

Isolated compounds from *Allium gramineum*, particularly isorhamnetin-3-O- β -D-glucopyranoside, showed strong scavenging effects with an IC50 of 20.1 \pm 0.8 μM , while Prosopogenin A of dioscin and Deltonine had weak activity with IC50 values over 100 μM .¹⁸

Ebrahimzadeh et al. investigated the antioxidant activity of the aerial parts and bulbs of *Allium paradoxum* using eight in vitro assays. The total phenolic content was 62.7 \pm 3.5 mg/g in the aerial parts and 7.4 \pm 0.2 mg/g in the bulbs, while the total flavonoid content was 47.9 \pm 2.6 mg/g in the aerial parts and 23.61 \pm 1.1 mg/g in the bulbs. Both parts exhibited significant antioxidant activity and contained high levels of iron and manganese.²⁹

Assadpour et al. investigated the antioxidant and antihemolytic effects of *Allium rotundum*'s essential oil and methanolic extract. The methanolic extract displayed superior activity compared to the essential oil in reducing effects, nitric oxide-scavenging, DPPH scavenging (IC50 values: 284 \pm 11.64 $\mu\text{g}/\text{mL}$ for extract vs. 1264 \pm 45.60 $\mu\text{g}/\text{mL}$ for oil), nitric oxide-

scavenging (IC50 values: 464 \pm 19.68 $\mu\text{g}/\text{mL}$ for extract vs. 1093 \pm 38.25 $\mu\text{g}/\text{mL}$ for oil), and Fe2+ chelating (IC50 values: 100 \pm 3.75 $\mu\text{g}/\text{mL}$ for extract vs. 1223 \pm 36.25 $\mu\text{g}/\text{mL}$ for oil). The extract and oil demonstrated significant dose-dependent H2O2 scavenging effects during H2O2-induced hemolysis (IC50: 786 \pm 29.08 mg/mL for oil).³⁸

Research by Motamed et al. revealed that *Allium rubellum* had the highest deoxyribose degradation inhibitory activity among the ten plants studied, with a 56.45 \pm 1.56% inhibition.⁴⁰

The analyses revealed that the bulb and leaf parts of *A. scorodoprasum* contained significant antioxidants. Total phenolic content ranged from 254.51 to 927.81 mg/kg in bulbs and 1929.05 to 19645.24 mg/kg in leaves. FRAP values ranged from 0.80 to 5.20 mM TE/g in bulbs and 14.31 to 47.83 mM TE/g in leaves. DPPH free radical scavenger effect varied from 0.99 to 9.02 $\mu\text{mol TE/g}$ in bulbs and 36.61 to 241.06 $\mu\text{mol TE/g}$ in leaves. Ascorbic acid content ranged from 29.14 to 314.01 mg/kg in bulbs and 200.64 to 1383.16 mg/kg in leaves. These findings highlight the abundant antioxidant richness of *A. scorodoprasum* subsp. *rotundum* leaves.⁴¹ Antioxidant enzyme activity analysis of *A. scorodoprasum* revealed notable results: compared to *A. sativum*. *A. scorodoprasum* exhibited an increase in catalase activity. Moreover, GPx and GSH-Px activities were detected in bulbs across all investigated *Allium* species. Among them, wild *A. scorodoprasum* displayed the highest GPx activity, showing a remarkable increase compared to *A. sativum*.⁴³

The antioxidant and anti-tyrosinase activity of different *A. ursinum* extracts and their metal complexes were evaluated using DPPH radical scavenging and mushroom tyrosinase assay.⁴⁶ The outcomes indicated that the polarity of the extracting solvents and the solubility of the phenolic compounds in these solvents significantly impacted the yield and phenolic content as the antioxidant and anti-tyrosinase activities.⁴⁶

Four flavonoids were isolated (1-4) From the water extract of *A. vineale*.⁵⁷ The assessment of total antioxidant activity was conducted using the ferric thiocyanate method in the linoleic acid system for both the crude extract and isolated compounds 1, 2, and 3, as well as Trolox and α -tocopherol. During the 20-hour incubation period at a concentration of 80 $\mu\text{g}/\text{mL}$, the activities were observed to be 64.8%, 79%, 75.6%, 82.2%, 75.7%, and 31.4%, respectively, for the crude extract, flavonoids 1, 2, and 3, Trolox, and α -tocopherol.⁵⁷

Other biological activities

In addition to the activities previously mentioned, *Allium* species possess several other significant biological properties.

For instance, steroidal saponins and sapogenins with thrombolytic activity, such as diosgenin, tigogenin, and ruscogenin, have been isolated from *A. affine*.⁷ Research indicates that *A. atroviolaceum* extract exhibits excellent antiplatelet activity, effectively inhibiting platelet aggregation

in vitro induced by ARA and ADP, with IC50 values of 0.4881 (0.4826–0.4937) mg/mL and 0.4945 (0.4137–0.5911) mg/mL, respectively.¹¹ Similarly, the leaf extract of *A. ursinum* demonstrates antiplatelet activity. In vitro, testing of antiplatelet activity was conducted using light transmission aggregometry, with induction by ADP, collagen, A23187, epinephrine, and arachidonic acid (ARA).⁵⁰

The antibacterial properties exhibited by aqueous and alcoholic extracts of *A. atroviolaceum* have also been investigated. Results show that after 48 hours of incubation, the minimum inhibitory concentration (MIC) of the aqueous extract against *S. aureus* was 3.125 mg/mL, while the alcoholic extract had an MIC of 6.25 mg/mL. For *Escherichia coli*, the MIC of the aqueous extract was 3.125 mg/mL, and the alcoholic extract's MIC was 12.50 mg/mL.¹² Additionally, the concentrated extract from *A. scorodoprasum* was tested against various microorganisms relevant to food technology, showing notable antimicrobial activity. The extract demonstrated the highest antibacterial activity against *S. aureus* (20.00 mm inhibition zone) and significant antifungal activity against *A. niger* (18.50 mm inhibition zone). However, *E. coli* exhibited a high resistance (MIC of 7.5 mg/mL) to the *A. scorodoprasum* extract. Among the tested bacteria and mold, *A. scorodoprasum* extract showed the most potent antibacterial effect against *S. aureus* and the most substantial antifungal effect against *A. niger*.⁴⁴

Research by Mskhiladze et al. demonstrated that although the total steroid saponins from *A. leucanthum* show moderate antibacterial activity, the spirostanol fraction exhibits vigorous activity. At the same time, bacterial strains were resistant to the furostanol fraction. Glycosides of β -chlorogenine displayed more potent activity than agigenin glycosides.²⁶

Phytochemical studies of *A. paradoxum*, particularly its saponin constituents, led to the isolation and identification of a dioscin-related steroidal saponin from the plant's bulbs by Rezaee et al. This compound exhibited significant leishmanicidal effects on promastigotes of *L. major* in both 10 and 50 μ g/mL concentrations.²⁸

Further investigation into *A. leucanthum* by Mskhiladze et al. found that the spirostanol fraction was ten times more active against *Leishmania amastigotes* than human cells.²⁵

Extracts from the aerial parts and bulbs of *A. paradoxum* at doses of 500 and 750 mg/kg intraperitoneally demonstrated a significant hepatoprotective effect by reducing serum aspartate aminotransferase (AST) and alkaline phosphatase (ALP) levels.³⁰

In vitro anti-plasmodial activity of *A. paradoxum* extract was most effective at an 80 μ g/mL dosage, resulting in 60.43% parasite growth inhibition compared to control groups. At 40 μ g/mL, growth inhibition was 52.48%, also statistically significant compared to controls.³¹

The analgesic activity of methanolic extracts of *A. paradoxum* was evaluated in male Balb/C mice using acetic acid-induced writhing and hot plate tests. The extracts

demonstrated notable analgesic activity in both experimental models. In the writhing test, the extract exhibited significant analgesic effects across all tested doses compared to the control group, reducing writhing behaviors ($p < 0.001$). In the hot plate test, the extract notably elevated pain thresholds by the 30th minute ($p < 0.001$) compared to the control group.³²

The nephroprotective effects of extracts derived from the aerial parts and bulbs of *A. paradoxum* were evident against gentamicin-induced nephrotoxicity in mice. Both extracts administered at 200 mg/kg/day exhibited protective effects by modulating blood urea nitrogen and creatinine levels.³³

Woo et al. investigated the inhibitory activities of compounds isolated from *A. victoralis* (allivictoside A–H) on neuroinflammation by measuring NO levels in LPS-activated BV-2 cells. Compounds 2 and 6 significantly inhibited NO production, indicating that flavonoid derivatives from *A. victoralis* possess anti-neuroinflammatory properties.⁵¹

The anti-inflammatory activity of *A. scorodoprasum* extract was evaluated, showing that the extract concentrations required to inhibit 50% of LOX and XO activities were 9.75 and 9.71 mg extract/mL, respectively. The IC50 values of quercetin and allopurinol for LOX and XO were 1.22 and 2.69 mg/mL, respectively.⁴⁴

CONCLUSIONS

In conclusion, this review compiles available literature on plants from the genus *Allium* growing in Georgia, focusing on the compounds isolated from these species and their biological activity. The gathered texts show that the major constituents of *Allium* plants in this region are saponins, phenolic, and sulfur compounds, which likely contribute significantly to their diverse pharmacological potentials.

No phytochemical studies have been conducted on several species of the genus *Allium* growing in Georgia, including *A. ledchanense*, *A. otchiauriae*, *A. saxatile*, *A. moschatum*, *A. aucheri*, *A. gracilescens*, *A. rupetchii*, *A. tauricola*, *A. pseudoflavum*, *A. candolleianum*, *A. albobianum*, *A. ponticum*, *A. rupestre*, *A. karsianum*, *A. chevsuricum*, *A. globosum*, *A. oreophilum*, *A. pseudostrictum*, *A. kunthianum*, *A. paczoskianum*, *A. szovistii*, *A. cardiostemon*. Therefore, future comprehensive studies are necessary to enhance understanding of these plants' pharmacological activities, chemical constituents, and effectiveness.

AUTHOR AFFILIATION

- 1 Direction of Pharmacognosy and Pharmaceutical Botany, Faculty of Pharmacy, Tbilisi State Medical University, Tbilisi, Georgia;
- 2 Laboratory of Pharmacognosy, Department of Pharmacy, Centre Interdisciplinaire de Recherche du Médicament-CIRM, University of Liège, CHU-B36, B-4000 Liège, Belgium.

REFERENCES

1. Gagnidze R. Vascular Plants of Georgia a Nomenclatural Checklist. Institute of Botany.; 2005.

2. Rahman K. Historical Perspective on Garlic and Cardiovascular Disease. *J Nutr.* 2001;131(3):977S-979S. doi:10.1093/jn/131.3.977S.
3. Bagrationi D. Iadigar Daudi. Edition of Tbilisi University.; 1993.
4. Panaskerteli-Tsitsishvili Z. Karabadini. Sabchota Sakartvelo; 1978.
5. Davlianidze M, Ghviniashvili T, Mukbaniani M, Jinjolia-Imnadze L, Jugheli T. Nomenclatural Checklist of Flora of Georgia. The Botanical Institute of Ilia State University; 2018.
6. Sobolewska D, Michalska K, Podolak I, Grabowska K. Steroidal saponins from the genus *Allium*. *Phytochem Rev.* 2016;15(1):1-35. doi:10.1007/s11101-014-9381-1.
7. Sadeghi M, Safaeian L, Ghazvini MA, Ramezani M. Evaluation of fibrinolytic and antioxidant effects of *Allium affine* hydroalcoholic extract. *Res Pharm Sci.* 2017;12(4):299-306. doi:10.4103/1735-5362.212047
8. Pkheidze TA, Kereselidze EV, Kemertelidze ÉP. Diosgenin, neoruscogenin, and ruscogenin from *Ruscus ponticus*, *R. hypophyllum*, and *Allium albidum*. *Chem Nat Compd.* 1971;7(6):823-823. doi:10.1007/BF00567962
9. Kravets SD, Vollerner YuS, Gorovits MB, Abubakirov NK. Steroids of the spirostan and furostan series from plants of the genus *Allium*. *Chem Nat Compd.* 1991;26(4):359-373. doi:10.1007/BF00598985
10. Zolfaghari B, Barile E, Capasso R, Izzo AA, Sajjadi SE, Lanzotti V. The Sapogenin Atroviolacegenin and Its Diglycoside Atroviolaceoside from *Allium atroviolaceum*. *J Nat Prod.* 2006;69(2):191-195. doi:10.1021/np0503350
11. Lorigooini Z, Ayatollahi SA, Amidi S, Kobarfard F. Evaluation of Anti-Platelet Aggregation Effect of Some *Allium* Species. *Iran J Pharm Res IJPR.* 2015;14(4):1225-1231.
12. Hafeznia B, Anvar S a. A, Kakoolaki S, Choobkar N. Antimicrobial efficiency of *Allium atroviolaceum* extract on Rainbow trout in different temperature and storage time. *Iran J Aquat Anim Health.* 2018;4(2):86-94. doi:10.29252/ijaah.4.2.86.
13. Khazaei S, Abdul Hamid R, Ramachandran V, et al. Cytotoxicity and Proapoptotic Effects of *Allium atroviolaceum* Flower Extract by Modulating Cell Cycle Arrest and Caspase-Dependent and p53-Independent Pathway in Breast Cancer Cell Lines. *Evid-Based Complement Altern Med ECAM.* 2017;2017:1468957. doi:10.1155/2017/1468957.
14. Khazaei S, Esa NM, Ramachandran V, et al. In vitro Antiproliferative and Apoptosis Inducing Effect of *Allium atroviolaceum* Bulb Extract on Breast, Cervical, and Liver Cancer Cells. *Front Pharmacol.* 2017;8:5. doi:10.3389/fphar.2017.00005.
15. Ghasemi S, Lorigooini Z, Wibowo J, Amini-khoei H. Tricin isolated from *Allium atroviolaceum* potentiated the effect of docetaxel on PC3 cell proliferation: role of miR-21. *Nat Prod Res.* 2019;33(12):1828-1831. doi:10.1080/14786419.2018.1437439.
16. Chincharadze DG, Kel'ginbaev AN, Gorovits MB, Éristavi LI, Gorovits TT, Abubakirov NK. Steroid saponins and sapogenins of *Allium*. XV. Eruboside B from *Allium erubescens*. *Chem Nat Compd.* 1979;15(4):442-446. doi:10.1007/BF00565042.
17. Eristavi L. The onion, *Allium fuscoviolaceum* Fom.-As a new stuff of Diosgenin. 1972;3(66). Accessed January 13, 2022. <https://dspace.nplg.gov.ge/handle/1234/219425>.
18. Mskhiladze L, Chincharadze D, Tits M, et al. IDENTIFICATION OF CYTOTOXIC AND ANTIOXIDANT COMPOUNDS FROM ALLIUM GRAMINEUM FLOWERS. *Int J Pharm Sci Drug Res.* 2015;7(1). Accessed October 14, 2021. <https://ijpsdr.com/index.php/ijpsdr/article/view/382>.
19. Mskhiladze L, Chincharadze D, St-Gelais A, et al. Isolation and Identification of Steroid and Flavonoid Glycosides from the Flowers of *Allium gramineum*. *Int J Pharm Sci Drug Res.* 2016;8(04). doi:10.25004/IJPSDR.2016.080409
20. Mskhiladze L, Legault J, Lavoie S, et al. Cytotoxic Steroidal Saponins from the Flowers of *Allium leucanthum*. *Molecules.* 2008;13(12):2925-2934. doi:10.3390/molecules13122925.
21. Mskhiladze L, Jokhadze M, Chincharadze D, Kuchukhidze J. Quantitative analysis of steroidal saponins from the *Allium leucanthum* C. Koch. *Collect Scientific Works.* 2008;44:150-151.
22. Mskhiladze L, Chincharadze D, Mshvildadze V, et al. Steroidal Glycosides from the Flowers of *Allium leucanthum*. *Chem Nat Compd.* 2015;51(5):900-904. doi:10.1007/s10600-015-1444-z.
23. Mskhiladze L, Chincharadze D, Elias R, Faure R, Balansard G, Kuchukhidze J. Steroidal saponins of *Allium leucanthum* C.Koch. *Ga Chem J.* 2007;7(1):70-72.
24. Mskhiladze L, Chincharadze D, Eristavi L, Kuchukhidze J. Steroidal sapogenins of *Allium leucanthum* C.Koch. *Ga Chem J.* 2007;7(1):73-74.
25. Mskhiladze L, Kuchukhidze J, Chincharadze D, Delmas F, Elias R, Favel A. In vitro antifungal and antileishmanial activities of steroidal saponins from *Allium leucanthum* C. Koch a Caucasian endemic species. *Georgian Med News.* 2008;(154):39-43.
26. Mskhiladze L, Galdava G, Mikaiia M, Chincharadze D, Kuchukhidze J. Antibacterial activity of Steroid saponins of the *Allium leucanthum* C. Koch. *Collect Scientific Works.* 2008;44:147-149.
27. Ghavam-Haghi F, Sadeghi Dinani M. Isolation and identification of *Allium astragalum* and 2-methoxy tyrosol from the bulbs of *Allium paradoxum*. *J HerbMed Pharmacol.* 2017;6:114-118.
28. Rezaee F, Zolfaghari B, Dinani M. Isolation of dioscin-related steroidal saponin from the bulbs of *Allium paradoxum* L. with leishmanicidal activity. *Res Pharm Sci.* 2018;13(5):469. doi:10.4103/1735-5362.236875.
29. Ebrahimzadeh MA, Nabavi SF, Nabavi SM, Eslami B. Antihemolytic and antioxidant activities of *Allium paradoxum*. *Cent Eur J Biol.* 2010;5(3):338-345. doi:10.2478/s11535-010-0013-5.
30. Nabavi SM, Hajizadeh Moghaddam A, Fazli M, et al. Hepatoprotective activity of *Allium paradoxum*. *Eur Rev Med Pharmacol Sci.* 2012;16 Suppl 3:43-46.
31. Elmi T, Hajjalilani F, Asadi MR, et al. Antimalarial effects of the hydroalcoholic extract of *Allium paradoxum* in vitro and in vivo. *J Parasit Dis Off Organ Indian Soc Parasitol.* 2021;45(4):1055-1064. doi:10.1007/s12639-021-01359-0.
32. Maghsoodi R, Afsha V, Rabani M, Mahmoodi M, Ebrahimzadeh MA. Antinociceptive effects of methanolic extract of *Allium paradoxum* in mice in Hot-plate and writhing tests. *Int Pharm Acta.* 2018;1(1):71-72. doi:10.22037/ipa.v1i1.20121.
33. Nabavi SF, Nabavi SM, Moghaddam AH, Naqinezhad A, Bigdelou R, Mohammadzadeh S. Protective effects of *Allium paradoxum* against gentamicin-induced nephrotoxicity in mice. *Food Funct.* 2012;3(1):28-29. doi:10.1039/c1fo10173k.
34. Maisashvili MR, Kuchukhidze DK, Gvazava LN, Eristavi LI. Steroidal Glycosides from *Allium rotundum*. *Chem Nat Compd.* 2008;44(4):545-547. doi:10.1007/s10600-008-9116-x.
35. Maisashvili MR, Gvazava LN, Kuchukhidze JK. Flavonoids and coumarins from *Allium rotundum*. *Chem Nat Compd.* 2009;45(1):87-88. doi:10.1007/s10600-009-9246-9.
36. Maisashvili MR, Kuchukhidze DzhK, Gvazava LN. Carotenoids and amino acids from *Allium rotundum*. *Chem Nat Compd.* 2009;45(5):769. doi:10.1007/s10600-009-9452-5.
37. Maisashvili MR, Eristavi LI, Gvazava LN, Gugunishvili DM. Steroidal sapogenins from *Allium rotundum*. *Chem Nat Compd.* 2007;43(6):756-757. doi:10.1007/s10600-007-0259-y.
38. Assadpour S, Nabavi SM, Nabavi SF, Dehpour AA, Ebrahimzadeh MA. In vitro antioxidant and antihemolytic effects of the essential oil and methanolic extract of *Allium rotundum* L. *Eur Rev Med Pharmacol Sci.* 2016;20(24):5210-5215.
39. Ismailov AI, Tagiev SA, Rasulov ÉM. Steroid saponins and sapogenins from *Allium rubellum* and *A. albanum*. *Chem Nat Compd.* 1976;12(4):495-495. doi:10.1007/BF00564837.
40. Motamed SM, Naghibi F. Antioxidant activity of some edible plants of the Turkmen Sahara region in northern Iran. *Food Chem.* 2010;119(4):1637-1642. doi:10.1016/j.foodchem.2009.09.057.
41. Taşçı B, Kütük H, Koca İ. Antioxidant Activity of *Allium scorodoprasum* L. subsp. *rotundum* (L.) STEARN Plant Grown in

- Turkey. *Turk J Agric - Food Sci Technol.* 2019;7(10):1561-1567. doi:10.24925/turjaf.v7i10.1561-1567.2583.
42. Mollica A, Zengin G, Locatelli M, Picot-Allain CMN, Mahomoodally MF. Multidirectional investigations on different parts of *Allium scorodoprasum* L. subsp. *rotundum* (L.) Stearn: Phenolic components, in vitro biological, and in silico propensities. *Food Res Int.* 2018;108:641-649. doi:10.1016/j.foodres.2018.03.064.
 43. Stajner D, Igić R, Popović BM, Malencić D. Comparative study of antioxidant properties of wild growing and cultivated *Allium* species. *Phytother Res PTR.* 2008;22(1):113-117. doi:10.1002/ptr.2278.
 44. Demir T, Akpınar Ö, Kara H, Güngör H. Phenolic profile and investigation of biological activities of *Allium scorodoprasum* L. subsp. *rotundum*. *Food Biosci.* Published online January 10, 2022;101548. doi:10.1016/j.fbio.2022.101548.
 45. Kurnia D, Ajiati D, Heliawati L, Sumiarsa D. Antioxidant Properties and Structure-Antioxidant Activity Relationship of *Allium* Species Leaves. *Mol Basel Switz.* 2021;26(23):7175. doi:10.3390/molecules26237175.
 46. Nikkhahi M, Souri E, Sarkhail P, Baeeri M, Mohammadhosseini N. Evaluation of anti-tyrosinase activity of *Allium ursinum* extracts and their metal complexes. *Acta Sci Pol Technol Aliment.* 2018;17(3):219-226. doi:10.17306/J.AFS.0585.
 47. Barla GF, Poroch-Seritan M, Sandulaec E, Ciornei SE. Antioxidant Activity and Total Phenolic Content in *Allium ursinum* and *Ranunculus ficaria*. *Food Environ Saf J.* 2016;13(4). <http://fens.usv.ro/index.php/FENS/article/view/123/121>.
 48. Korga A, Ostrowska M, Iwan M, et al. Ethanol extracts of *Allium* sp. regulate cyclooxygenase-2 and E-cadherin expression in gastric cancer MKN74 cell line and enhance doxorubicin toxicity. *Food Nutr Res.* 2019;63. doi:10.29219/fnr.v63.3449.
 49. Pârvu M, Alina Elena P, Vlase L, Roșca-Casian O, Parvu O. Antifungal properties of *Allium ursinum* L. ethanol extract. *J Med PLANTS Res.* 2011;5:2041-2046.
 50. Hiyasat B, Sabha D, Grötzinger K, et al. Antiplatelet Activity of *Allium ursinum* and *Allium sativum*. *Pharmacology.* 2009;83(4):197-204. doi:10.1159/000196811.
 51. Woo KW, Moon E, Park SY, Kim SY, Lee KR. Flavonoid glycosides from the leaves of *Allium victorialis* var. *platyphyllum* and their anti-neuroinflammatory effects. *Bioorg Med Chem Lett.* 2012;22(24):7465-7470. doi:10.1016/j.bmcl.2012.10.043.
 52. Khan S, Fatima I, Kazmi M, Malik A. New Secondary Metabolites from *Allium victorialis*. *Helv Chim Acta.* 2013;96. doi:10.1002/hlca.201200581.
 53. Khan S, Mehmood R, Kazmi MH, Malik A. Alliumonoate: a new cyclopentane derivative from *Allium victorialis*. *J Asian Nat Prod Res.* 2011;13(12):1165-1169. doi:10.1080/10286020.2011.619183.
 54. Kim HJ, Park MJ, Park HJ, Chung WY, Kim KR, Park KK. Chemopreventive and Anticancer Activities of *Allium victorialis* var. *platyphyllum* Extracts. *J Cancer Prev.* 2014;19(3):179-186. doi:10.15430/JCP.2014.19.3.179.
 55. Kim YS, Jung DH, Lee IS, et al. Effects of *Allium victorialis* leaf extracts and its single compounds on aldose reductase, advanced glycation end products and TGF-β1 expression in mesangial cells. *BMC Complement Altern Med.* 2013;13:251. doi:10.1186/1472-6882-13-251.
 56. Chen S, Snyder JK. Diosgenin-bearing, molluscicidal saponins from *Allium vineale*: an NMR approach for the structural assignment of oligosaccharide units. *J Org Chem.* 1989;54(15):3679-3689. doi:10.1021/jo00276a033.
 57. Demirtas I, Erenler R, Elmastas M, Goktasoglu A. Studies on the antioxidant potential of flavones of *Allium vineale* isolated from its water-soluble fraction. *Food Chem.* 2013;136(1):34-40. doi:10.1016/j.foodchem.2012.07.086.
 58. Shahidi F, Zhong Y. Measurement of antioxidant activity. *J Funct Foods.* 2015;18:757-781. doi:10.1016/j.jff.2015.01.047.