

Solanum Nigrum Linn: A Detailed Exploration of Its Phytochemistry and Pharmacological Properties- A Review

Ruchika Khatri 

IIS (Deemed to be University) Jaipur, Rajasthan 302020, India

Citation: Ruchika Khatri (2025). *Solanum Nigrum* Linn: A Detailed Exploration of Its Phytochemistry and Pharmacological Properties- A Review. *Acta Biology Forum*. DOI: <https://doi.org/10.51470/ABF.2025.4.2.19>

Corresponding Author: **Ruchika Khatri** | E-Mail: (ruchikakhtr92@gmail.com)

Received 30 April 2025 | Revised 22 May 2025 | Accepted 23 June 2025 | Available Online 27 July 2025

ABSTRACT

Solanum Nigrum Linn, commonly named as makoi or makoy, is a versatile and widely distributed plant species with significant ecological, economic, and medicinal potential. This review explores the botanical, ecological and medicinal characteristics of *Solanum nigrum*, providing a comprehensive analysis of its morphology, taxonomy, distribution, and reproductive strategies. The plant's adaptability to diverse habitats, ranging from tropical to subtropical regions, highlights its invasive potential. In terms of morphology, *Solanum nigrum* is a perennial herb or small tree with heart-shaped leaves and flowers. Its anatomical structure, including its vascular tissue and seed morphology, reflects typical characteristics of dicotyledonous plants. Beyond its botanical significance, *Solanum nigrum* holds substantial economic value. Additionally, *Solanum nigrum* has high medicinal potency like antibacterial, antifungal, anti-inflammatory, anticonvulsant, antioxidant etc. properties along with traditional uses in various cultures. In this review, tends to highlight the potency of plant to cure several diseases because of the existence of many phytometabolites. Hence, it is useful in many pharmaceutical applications.

Keywords: *Solanum Nigrum*, antifungal, secondary metabolites, dicotyledonous, pharmaceuticals.

Introduction

Since the beginning of human life, the hypothesis of "drug" is people have utilised many plant species for curative causes. According to the WHO survey 80% of the peoples be located in developing countries depend on entirely on traditional remedies for their health care requirements. Now a day, herbal medications are famed back into resources because potency of traditional medicines i.e. antibiotics, which have proved globally efficient against severe infections. As the practice of conventional herbal remedies has increased the affair concerning the protection, value and ability in commercial and developing countries is being addressed up. Proved study of the biochemical compounds from medicinal herbs and pharmacological investigation reveals us information for emerging herbal medications from novel agents. However, estimated 400,000 plant species, only 5 to 7 % has reported for biotic pharmacological activity and phytochemically (Potawale, S. E., et al, 2008). More than 2,000 different kinds of tropical plants in the genus *Solanum* can be grown anywhere in the world (Chen, X., et al., 2022). *S. nigrum*, sometimes referred to as Makoi or black nightshade, thrives as a weed in humid areas in a range of soil types, including shallow, deep, stony, and dry soils. By reproducing the seeds in well-kept nursery beds between April and June, it can even be produced in tropical areas. Traditional medicine also frequently uses *S. nigrum* to treat a variety of ailments, such as fever, inflammation, and discomfort. As a diuretic, antipyretic, hepatoprotective, anti-inflammatory, antioxidant, and antitumorigenic, this herb is also utilized as medicine for a variety of medical conditions. Many biocompounds have been reported and isolated those are responsible for innumerable medicinal activities. (Jain, R. et al. (2011))

Objective: The purpose of this study is to discover medicinal and phytochemical potential of the plant *Solanum Nigrum* L.

Methods and Methodology:

The primary resource of data used for this review work comprises of research papers and review articles published by reputable publishers such as Springer, Elsevier, Wiley, etc. Literature research was conducted by using Google Scholar and PubMed with the following keywords: makoy, *Solanum nigrum*, Makoi, Indian medicinal weed *Solanum nigrum*, pharmaceutical properties of *Solanum nigrum*, biochemical components of *Solanum nigrum*, medicinal properties of makoy, etc. Several research papers published in the trustworthy have contributed meaningfully to the foundational consideration of this review paper. For instance, prior researches have detailed the taxonomical and geographical description, traditional usage, phytochemical composition and pharmacological activities including antioxidant, antimicrobial, antihelminthic, cytoprotective, Anti-convulsant activity were screened of crude extracts of various parts of this medicinal weed plants were used to explore the information about *Solanum Nigrum* L.

Traditional benefits:

Instead of using other plant parts for medicinal purposes, *Solanum nigrum* fruits and leaves are frequently used. The leaves were used as an anti-tuberculosis treatment and as a dressing for gouty and rheumatic joints as well as skin diseases. Additionally, leaves can be used to treat nausea, dropsy, and neurological diseases. The berry and floral extracts are useful for treating colds and coughs. The plant part is useful in bronchitis and pulmonary tuberculosis medications, diuretics, etc.

Copyright: This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY 4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

The fruit's juice is discarded as a remedy for heart problems, hydrophobia, ophthalmopathy, and diarrhea. Berries are helpful in treating skin infections and inflammations because of their tonic, diuretic, and cathartic qualities. Hepatitis, osteopathology, and ophthalmopathy can all be treated with the roots. Leprosy, hemorrhoids, nephropathy, dropsy, digestive, diuretic, laxative, diaphoretic, sedative, edema, cardiovascular disorders, and general debility are all treated with whole plant components.



Leaves



Fruit



Flower



Seeds



Root

THERAPEUTIC USES

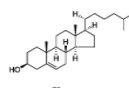
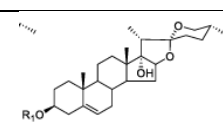
Plants have been utilized with varying success to cure and prevent diseases due to their therapeutic efficacy and pharmacological potential. *Solanum nigrum* as a medicinal plants has been described in ancient literature of Indian system of medicine (ayurveda). *Solanum Nigrum* possesses various therapeutic usages due to the presence of alkaloid, flavonoids and other phytochemicals. (Gabrani, R., et al, 2012)

- "To treating TB, nausea, and neurological system diseases.
- Works in the medication of rheumatoid arthritis, gout, skin conditions".
- It works against cancer.
- "Functions in the management of bacterial infections that might result in colds and coughing".
- Shows potential towards the growth and the oxidation of biomolecules.
- Effective in dropping inflammation and curing liver from damage.
- Act as anticonvulsant, which prevents seizures.
- Fruit shows their effectiveness in the ailment of diarrhoea and gastrointestinal illnesses

Taxonomical classification

- "Division: Magnoliophyta"
- "Class: Magnoliopsida"
- "Subclass: Asteridae"
- "Order: Solanales"
- "Genus: Solanum"
- "Species: *Solanum Nigrum* L."
- "Authority: Linn."

Phytoconstituents of *Solanum Nigrum* L

S.no	Chemical component	Extract	Formula	References
1	Cholesterol	EtoH		He, J., et al (2015)
2	SolanigrosideH	EtOH		He, J., et al (2015)

GEOGRAPHICAL DISTRIBUTION

People from Europe and Asia first familiarized it in Americas, Australia and Africa. Other nations including "India, Afghanistan, Bangladesh, Bhutan, Indonesia, Iraq, Iran, Japan, Pakistan, Europe, North America, South America, Brazil, Peru, Colombia, etc. can also access it".

Biophysical Limit & botanical description

Altitude: up to Sea level at 3048m,

Temperature: temperatures at 20 and 30°C,

Photoperiod: Approximately 16 to 18 hours a day

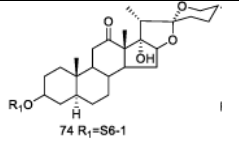
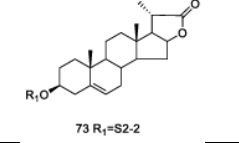
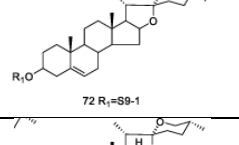
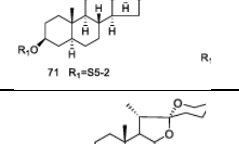
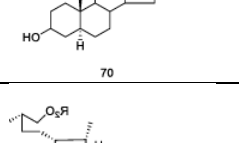
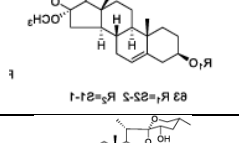
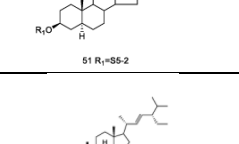
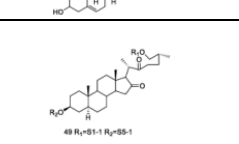
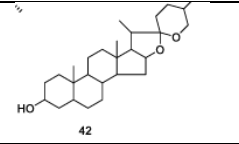
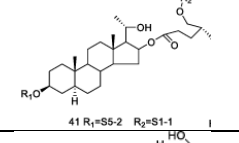
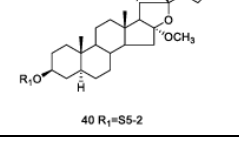
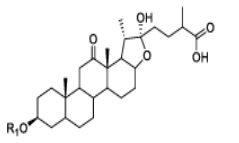

Moisture: 40-45% of seed moisture necessary for optimal germination

Rainfall: 500-1200 mm annually

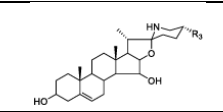
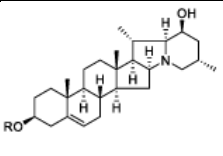
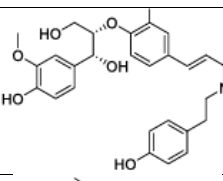
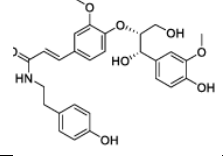
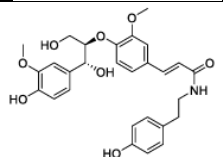
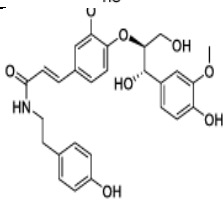
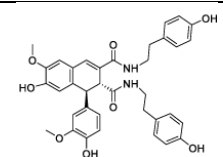
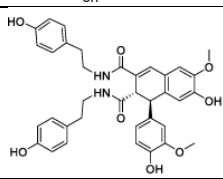
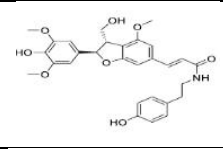
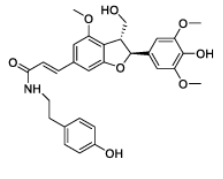
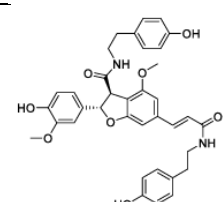
Soil type: sandy, loamy, clay rich soil (alkaline soil)

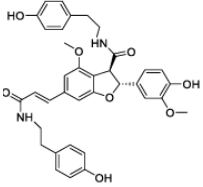
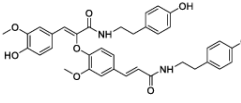
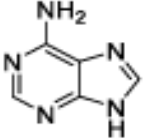
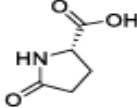
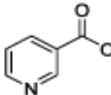
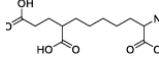
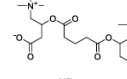
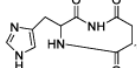
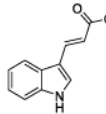
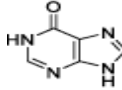
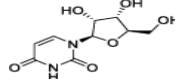
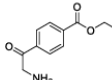
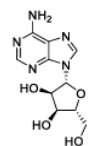
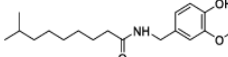

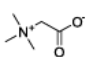
Plant habit

Simple multicellular glandular hairs provide protection for subglabrous plants, which can grow up to 80 cm in height. The branching root system is tapped. The stem of the plant is erect, glabrous, green, unbranched, and slightly woody. The leaves have an oval shape, sinuate dentate margins, and a maximum width of 6.0 cm. Inflorescences are simple cymes with up to ten blooms and tiny pedicels. The calyces are 2 mm long and accrescent, sticking to the base of the ripe fruit, while the sepals are oval in shape. Corolla are stellate and white. Anthers are between 1.6 and 2.9 mm long and have a golden hue. Pollen has a diameter of between 26.6 and 35.8 μm . Styles are 3.5 mm long. The fruit has axile placentation, a thin, papery epicarp, pulpy mesocarp, and seeds that stay in the berry pulp. The berry is broadly round, obtuse, dark purple to blackish green, and 12 mm wide. Each fruit contains 15 to 60 seeds, each up to 2.3 mm long. The bark is having pale yellow wood inside. (Potawale, S. E., et al (2008)) (Rani, Y. S., Reddy, et al (2017))

3	NigrumninII	EtoH		He, J., et al (2015)
4	Dumoside	EtoH		Gao, S, H., et al (2021)
5	UttroninB	EtoH		He, J., et al (2015)
6	UttroninA	EtoH		Wu, X. W. (2011).
7	Tigogenin	EtoH		He, J., et al (2015)
8	"(25R)-26-O-β-D-glucopyranosylfurost-5(6)-ene-16α-methoxy-3β,26-diol-3-O-α-L-rhamnopyranosyl C39H60O16 MeOHWangetal. (2017)-(1→2)-[α-L-rhamno"	MeoH		Wang, Y., (2017).
9	InunigrosideA	MeOH		Ohno, M.,(2012).
10	"Stigmast-5,22-dien-3β-ol"	EtOH MeOH		Sharma, B. K., (2012).
11	"(25R)-26-O-β-D-glucopyranosyl-cholest-5α-3β,26-diol-16,22-dione-3-O-β-D-glucopyranosyl-(1→2)-[β-D-glucopyranosyl-(1→3)]-β-D-glucopyranosyl-(1→4) β-D-galactopyranoside(25R)-26-O-β-D-glucopyranosyl-cholest-5α-3β,26-diol-16,22-dione-3-O-β-D-glucopyranosyl-(1→2)-[β-D-glucopyranosyl-(1→3)]-β-D-glucopyranosyl-(1→4) β-D-galactopyranoside"	MeOH		Xiang, L., et al (2018).
12	"Tigogenin/(25R)-5α-spirostan-3β-ol"	EtOH		Wu, X. W. (2011).
13	NigrosideA	EtOH		Wu, X. W. (2011)
14	SolanigrosideO	EtOH		Wu, X. W. (2011).
15	"(25R)-5α-furost-3β,22α-diol-12-one-26-carboxylicacid-3-O-β-D-glucopyranosyl-(1→4)-[O-β-D-glucopyranosyl-(1→2)]-O-β-D-glucopyranosyl (1→4)-O-β-D-galactopyranoside"	EtOH		Yang, Y. (2014).

16	β -daucosterol	EtOH		Yang, Y. (2014).
17	28-O- β -D-glucopyranosylbetulinic acid 3 β -O- β -D-glucopyranoside	EtOH		Yang, Y. (2014)
18	12-keto-porrigenin	EtOH		Zhao, Y. (2010).
19	Pterosterone	EtOH		Zhao, Y. (2010).
20	Stigmasterol	EtOH		Zhao, Y. (2010).
21	Degalactotigonin	EtOH		Wang, L. Y.,(2007).
22	Diosgenin	MeOH		Gao, S, H.,et al (2021).
23	Tomatidenol	EtOH		Gao, S, H., et al 2021
24	Solanocapsine	EtOH		Gao, S, H.,et al (2021).
25	SolanigrosideQ	EtOH		Gao, S, H.,et al (2021). Zhou, X. L. (2006).
26	"(25R)-22 α N-4-nor-spirosol-5(6)-en-3 β -ol-6-al-3-O-L-rhamnopyranosyl-(1 \rightarrow 2) [α -L-rhamnopyranosyl-(1 \rightarrow 4)]- β -D-glucopyranoside"	MeOH		Gao, S, H.,et al (2021). Zhou, X. L. (2006).
27	N-methylsolasodine	EtOH		Zhou, X. L. (2006). He, J.,(2015) Zhou, et al (2015).
28	23-O-acetyl-12 β -hydroxysolasodine	EtOH		He, J., (2015).

29	15 α -hydroxysolasodine	EtOH		Wu, X. W. (2011).
30	Leptininel	EtOH		Gao, S, H., et al. (2021).
31	"(7R,8S)-1-(4-hydroxy-3-methoxyphenyl)-2-{4-[2-[N-2-(4-hydroxyphenyl) ethyl] Carbamoylphenyl]C28H32NO8 Lietal. (2019)-2-methoxyphen"	EtOH		Li, J. H., et al (2021).
32	"(7S,8R)-1-(4-hydroxy-3-methoxyphenyl)-2-{4-[2-[N-2-(4-hydroxyphenyl) ethyl] Carbamoylphenyl]Liatal. (2019)-2-methoxyphenoxy}}-1"	EtOH		Li, J. H., et al (2021).
33	"(7R,8R)-1-(4-hydroxy-3-methoxyphenyl)-2-{4-[2-[N-2-(4-hydroxyphenyl)ethyl] Carbamoylphenyl 2-methoxyphenoxy}}-1,3-propanodiolnamed"	EtOH		Li, J. H., et al (2021).
34	"(7S,8S)-1-(4-hydroxy-3-methoxyphenyl)-2-{4-[2-[N-2-(4-hydroxyphenyl)ethyl] Carbamoylphenyl 2-methoxyphenoxy}}-1,3-propanodiolnamed"	EtOH		Li, J. H., et al (2021).
35	"7'S, 8'R-7-hydroxy-1-(4-hydroxy-3-methoxyphenyl)-N2, N3-bis (4 hydroxyphenethyl)-6-methoxy Lietal. (2019)-1,2-dihydronaphthalene-2,3-d"	EtOH		Li, J. H., et al (2021).
36	"7'R,8'S-7-hydroxy-1-(4-hydroxy-3-methoxyphenyl)-N2,N3-bis(4 hydroxyphenethyl)-6-methoxy 1,2-dihydronaphthalene-2,3-dicarboxamide"	EtOH		Li, J. H., et al (2021).
37	"7'R,8'S-7-(4-hydroxy-3,5-dimethoxyphenyl)-3'-hydroxymethyl-1'-[N-7''-(4'' hydroxyphenyl)ethyl] carbamoylphenyl-3'-methoxybenzodihydrofuran"	EtOH		Li, J. H., et al (2021).
38	"7'S,8'R-7-(4-hydroxy-3,5-dimethoxyphenyl)-3'-hydroxymethyl-1'-[N-7''-(4'' hydroxyphenyl)ethyl] carbamoylphenyl-3'-methoxybenzodihydrofuran"	EtOH		Li, J. H., et al (2021).
39	"(7'R,8'R)-2-(4-Hydroxy-3-methoxyphenyl)-3-[N-2-(4-hydroxyphenyl)ethyl] carbamoyl-5-[N-2-(4-hydroxyphenyl)ethyl] carbamoylphenyl-7-methoxybenzodihydrofuran"	EtOH		Li, J. H., et al (2021).

40	"(7'S,8'S)-2-(4-Hydroxy-3-methoxyphenyl)-3-[N-2-(4-hydroxyphenyl)ethyl] carbamoyl-5-[N-2-(4-hydroxyphenyl)ethyl]carbamoyl-7-methoxybenzodihydrofuran"	EtOH		Li, J. H., et al (2021).
41	Cannabisin F	EtOH		Li, J. H., et al (2021).
42	Adenine	MeOH		Gao, S, H.,et al. (2021).
43	Pyroglutamicacid	MeOH		Gao, S, H.,et al. (2021).
44	Nicotinicacid	MeOH		Gao, S, H.,et al. (2021).
45	9-aminononane-1,3,9-tricarboxylicacid	MeOH		Gao, S, H.,et al. (2021).
46	Glutarylcarntine	MeOH		Gao, S, H.,et al. (2021).
47	"6S)-3-((1H-imidazol-4-yl)methyl)-6-amino-1,4-diazocane-2,5,8-trione"	MeOH		Gao, S, H.,et al. (2021).
48	3-Indoleacrylicacid	MeOH		Gao, S, H.,et al. (2021).
49	6-Hydroxypurine	MeOH		Gao, S, H.,et al. (2021).
50	Uridine	MeOH		Gao, S, H.,et al. (2021).
51	Ethyl4-glycylbenzoate	MeOH		Gao, S, H.,et al. (2021).
52	Adenosine	MeOH		Gao, S, H.,et al. (2021).
53	Dihydrocapsaicin	MeOH		Gao, S, H.,et al. (2021).
54	Choline	MeOH		Gao, S, H.,et al. (2021).
55	Betaine	MeOH		Gao, S, H.,et al. (2021).

Pharmacological activities of *Solanum Nigrum*

1. Hepatoprotective activity

Aqueous extracts of *S. nigrum* shows hepatoprotective activity against CCl₄ intoxication. As observed in the CCl₄-infused groups, increases in AST, ALT, ALP, and serum bilirubin in groups 1 and 2 suggested severe pathological states of the infected rats' livers. These values dropped in Groups 3 and 4 in a dose-dependent way, particularly in Group 4 (500 mg/kg), which led to less liver damage in the rats. According to the research, liver dysfunction is indicated by a drop in Hb, RBC, and MCHC values in the CCl₄-treated groups (Groups 2 and 4). [R. A. M. Elhag et al., 2011].

2. Antiulcerogenic and antidiarrhoeal activity

Methanolic extract of *Solanum Nigrum* resulted anti ulcerogenic effect on headache drug-induced ulceration in rodents. *Solanum nigrum* berries shows gastroprotective effect due to the presence for free radicals. A crude extract of SN dried fruit has also been shown to have antidiarrheal qualities, and SN berries may be helpful in treating gastrointestinal issues. At two dosages of 250 and 500 mg/kg body weight, the fruit extracts show notable antidiarrheal activity against castor oil-induced diarrhea in rats, reducing the frequency of stools in rodents. (Mani, R. K. et al., 2022).

3. Neuropharmacological activity:

The neuropharmacological effects of ethanolic extracts of *Solanum nigrum* L. (Solanaceae) berries were tested on experimental animals. The extract significantly lengthened pentobarbital-induced naps, changed general behavior patterns, reduced exploratory behavior patterns, suppressed aggressive behavior, changed locomotor activity, and reduced spontaneous motility when given intraperitoneally. The results suggest that the fruit of *S. nigrum* may have CNS-depressant qualities. (Perez G, R. M., et al., 1998).

4. Antitumor activity

Using a range of chromatographic procedures, six steroidal glycoalkaloids were extracted, separated, and purified from "*Solanum nigrum* L. (SNL)" by acid and alkaline precipitation. These chemicals' structures were clarified using spectroscopic data. The anticancer activity and its molecular mechanism were examined using the "Methyl Thiazolyl Tetrazolium (MTT)" method, flow cytometry, colorimetric assay, and an immunocytochemical method. MGC-803 cells were found to be cytotoxically affected by "solasonine, β 1-solasonine, solamargine, and solanigraside P". Three sugar complex units, "a-L-rhamnopyranose at C-2", or "hydroxyl group" effects on the steroidal backbone are possible therapeutic options for gastric cancer. (X. Ding and others, 2013).

5. Antioxidant effect

Effective antioxidant qualities have been reported for *S. nigrum* L. A thorough investigation was conducted to determine *S. nigrum*'s antioxidant capacity. Total phenolic content (TPC) is a potent indicator of plant antioxidant efficacy and shows the mechanism of peroxidation inhibition against the linoleic acid peroxidation system. The TPC readings showed a small change. All species revealed significant antioxidant activity from linoleic acid, which was similar to Trolox as the positive control. [A. Mohyuddin et al., 2022].

6. Molluscicidal and larvicidal activities

It has been shown that *S. Nigrum* root extracts are effective against "*Alternaria Brassicicola* Isolates ABA-31" and ABA104, which induce black leaf spot on "Chinese cabbage (*Brassica pekinensis*)". Methanolic preparations of dried black nightshade root tissue exhibit antifungal properties that combat *A. brassicicola*. When root extracts were evaluated for antibacterial activity using ethyl acetate, n-butanol, and water fractions, n-butanol extracts showed the best results. Saponins were identified as the active components responsible for SN's antibacterial qualities (Muto, H., et al., 2006). The "sub-lethal (LC25)" concentration of SN leaves was observed to affect the AST, ALT, and LDH activities of the Saudi Arabian mollusk *Biomphalaria Arabica*, which could reveal how its molluscicidal actions work (Ajayi, E. I. O. (2011)). According to Amer and Manal (2005) and Ahmed, A. H. (1998), the binary combination of SN and "*Iris pseudacorus*" demonstrates molluscicidal and cercaricidal efficacy against *Biomphalaria alexandrina* and *Schistosoma mansoni* cercariae, respectively. Following a 30-minute pre-treatment of mice with different dosages (2.5–10 mg/ml) of crude water extract of SN, *S. mansoni* cercariae penetration "($p < 0.001$)" and "infectivity ($p < 0.01$)" were considerably reduced (Amer and Manal, 2005). Additionally, a recent study by Raghavendra et al. (2009) and Ahmed et al. (2002) assessed Sn extracts as a larvicidal agent against five laboratory-colonized strains of mosquito species.

7. Anticancer

The investigation showed that total alkaloids may be important in preventing the growth of tumor cells. However, the glycoprotein (150 x 10³) that was extracted from *S. nigrum* may have shown anti-cancer properties by inhibiting NF-kappa B's anti-apoptotic pathway, activating to boost the caspase cascade response, and boosting nitric oxide synthesis. (L. An and others, 2006).

8. Antimicrobial activity

Bacterial and fungal strains were suppressed by *Solanum nigrum* methanolic extract. The findings show that the zone of inhibition against the investigated microbial strains grew as the extract content rose. *S. nigrum* methanolic extract was observed to inhibit Gram positive bacteria at all tested concentrations "(100 mg, 75 mg, 50 mg, and 25 mg)", however Gram negative bacteria exhibited no activity at a concentration of 25 mg/ml. SN extracts demonstrated good action against the investigated microorganisms, with the exception of *P. aeruginosa*, which showed low activity compared to the other tested germs. SN extract was found to have the largest zone of inhibition against *S. aureus*, *B. subtilis* came next, and ethanolic extract proved to be the most effective against Gram-negative bacteria, with *E. coli* exhibiting the largest zone of inhibition, followed by *P. aeruginosa*. The extract's activity against *B. subtilis* and *S. aureus* was similar to that of the common medication ciprofloxacin. According to Kaushik et al. (2009), ethanolic extract had more action against *Candida albicans* with a zone of inhibition than the conventional medication amphotericin B.

9. Anti-convulsant activity

It was discovered that the extracts significantly affected rats' supramaximal electric shock.

When compared to the common medication phenytoin, the extract's percentage inhibitory activity at 100, 200, and 300 mg/kg was 36.01, 25.21, and 38.86%, in that order. "The effects of ethanolic extracts of *S. nigrum* in supra maximal electric shock technique in rats. Ravi, V. (2009)"

10. Antinociceptive and Antipyretic Effects

the chloroform extract from *Solanumnigrum* leaves' antipyretic and antinociceptive qualities in a range of animal models. "To make the extract, the air-dried powdered leaves (20 g) were steeped (1:20; w/v) in chloroform for 72 hours, evaporated (40°C) under low pressure to dryness (1.26 g), and then dissolved (1:50; w/v) in dimethylsulfoxide (DMSO)". Mice and rats received all dosages (s.c.; 10 ml/kg) half an hour prior to testing. DMSO was used to dilute the supernatant, which was considered the stock solution with a dosage of 200 mg/kg, to 20 and 100 mg/kg. The extract showed significant ($p < 0.05$) antinociceptive efficacy in the abdominal constriction, hot plate, and formalin tests. Furthermore, the extract showed significant ($p < 0.05$) antipyretic and anti-inflammatory properties in the carrageenan-induced paw edema and brewer's yeast-induced pyrexia tests. The actions were dose-independent overall. By demonstrating that the lipid-soluble extract of *S. nigrum* leaves possessed antinociceptive, anti-inflammatory, and anti-pyretic properties, the current study confirmed the traditional claims, Zakaria, Z. A. (2009)

11. Cytoprotective Activity

Potawale and colleagues (2008) examined the ability of a 50% ethanolic-produced plant extract from *Solanumnigrum* Linn. To protect Vero cells from gentamycin damage in vitro. To find out if cytotoxicity was present, an investigation used the assays for mitochondrial dehydrogenase activity and Trypan Blue exclusion. The ability of the ethanol extract to function as a hepatoprotective agent was also evaluated by looking at the liver under a microscope to look for any histological abnormalities. It has a strong hepatoprotective effect as a direct consequence of the discovery 1599. When the toxicant (carbon tetrachloride) was given to a rat, the liver tissue showed significant centrilobular necrosis. Treatment side effects include the regeneration of healthy liver cells and the elimination of necrosis and vacuoles. An adult male albino Wistar rats weighing between 150 and 170 grams served as the study's subjects. The study made use of these rats. The animals were kept in polypropylene cages with a 12-hour day and 12-hour night cycle and a constant temperature of 25 degrees Celsius. Six groups of six creatures each were created from the animals. After that, each group was put in its own cage. Each group's stomach was intubated to administer the medicine. This was the technique of administration used. Injections of 0.2 ml of acacia, 20% ethanol, and "SNFET" should be administered to groups 1, 2, and 3, respectively. Groups 2 and 3 should also receive the 20% ethanol injection. All of the rats in the group were allowed to go without food for the whole night before receiving ketamine chloride to induce anesthesia. Following the collection of blood, plasma was obtained for use in a number of biochemical assessments. These calculations relied on blood. A few drops of heparin were added to each sterile, fully dry test tube used to draw the blood. (M. Sankaran, 2012).

12. Anti Seizure Activity

The studies demonstrate that administering an intraperitoneal leaf extract to rats and mice has an antiseizure effect. Amphetamine use may have a significant impact on the antiseizure's effectiveness. It is beneficial to improve the plant SN's antiseizure properties. (N. N. Wannang, 2008)

Conclusion

NigrumSolanum Linn is an exceptional remedy for a variety of illnesses, including hepatitis, ulcers, fever, and several immunological conditions and associated remedies. This herb can be used to treat cytotoxicity and hepatotoxicity as well as to support the health of the liver and kidneys. This herbal remedy has potent antibacterial, anti-inflammatory, anti-diabetic, and immunostimulant properties. Due to the presence of several phytochemicals, this plant is probably a valuable and potent plant with the greatest results in pharmacological investigations. It is an intriguing herb with potential that can be utilized to treat a number of illnesses. It also possesses numerous additional potencies, such as anticonvulsant, antiallergic, and hepatoprotective. This review of the literature explains the plant's potency by examining experimental findings indicating *Solanumnigrum* has traditional medical value for treating ulcers, hepatitis, and numerous immunological bids in cancer, among other conditions. Additionally, it is an immunostimulant, central nervous system, and antidiabetic. It increases their use in pharmaceutical and medical procedures to treat a range of illnesses.

References

1. Potawale, S. E., Sinha, S. D., Shroff, K. K., Dhalawat, H. J., Boraste, S. S., Gandhi, S. P., & Tondare, A. D. (2008). *Solanumnigrum* Linn: A phytopharmacological review. *Pharmacologyonline*, 3, 140-63.
2. Chen, X., Dai, X., Liu, Y., Yang, Y., Yuan, L., He, X., & Gong, G. (2022). *Solanumnigrum* Linn.: an insight into current research on traditional uses, phytochemistry, and pharmacology. *Frontiers in Pharmacology*, 13, 918071
3. Jain, R., Sharma, A., Gupta, S., Sarethy, I. P., & Gabrani, R. (2011). *Solanumnigrum*: current perspectives on therapeutic properties. *Altern Med Rev*, 16(1), 78-85.
4. Gabrani, R., Jain, R., Sharma, A., Sarethy, I. P., Dang, S., & Gupta, S. (2012). Antiproliferative effect of *Solanumnigrum* on human leukemic cell lines. *Indian journal of pharmaceutical sciences*, 74(5), 451.
5. Rani, Y. S., Reddy, V. J., Basha, S. J., Koshma, M., Hanumanthu, G., & Swaroopa, P. (2017). A review on *Solanumnigrum*. *World J. Pharm. Pharm. Sci*, 6, 293-303.]
6. He, J., Zhou, C. D., Ma, B. Z., Liu, F., Liu, X., and Zhao, T. (2015). Research Progress on Chemical Constituents and Antitumor Pharmacological Activities of *Solanumnigrum*. *China Pharm*. 26 (31), 4433–4436. doi:10.6039/j.issn.10010408.2015.31.37
7. Gao, S. H., Su, Z. Z., Yang, L. J., and Li, Z. Y. (2021). Chemical Components from Stems of *Solanumnigrum* by LC-MS and NMR. *Chin. Tradit. Herb. Drugs*. 52 (5), 1263–1273. doi:10.7501/J.issn.0253-2670.2021.05.006 Pharmacological Activities of *Solanumnigrum*. *China Pharm*. 26 (31), 4433–4436. doi:10.6039/j.issn.10010408.2015.31.37
8. Wu, X. W. (2011). A virtual screening research for the antitumor activity of the ingredients in LSYQD. Henan: Master, Zhengzhou University.
9. Wang, Y., Xiang, L., Yi, X., and He, X. (2017). Potential anti-inflammatory steroidal saponins from the berries of *solanumnigrum* L. (European blacknightshade). *J. Agric. Food Chem*. 65 (21), 4262–4272. doi:10.1021/acs.jafc.7b00985

10. Ohno, M., Murakami, K., El-Aasr, M., Zhou, J.-R., Yokomizo, K., Ono, M., et al. (2012). New spirostanol glycosides from *Solanum nigrum* and *S. jasminoides*. *J. Nat. Med.* 66, 658–663. doi:10.1007/s11418-012-0637-
11. Sharma, B. K., Iyer, D., and Patil, U. K. (2012). Bioactivity guided fractionation in experimentally induced hyperlipidemia in rats and characterization of phytoconstituent from *Solanum nigrum*. *J. Herbs, Spices Med. Plants* 18,
12. Xiang, L., Wang, Y., Yi, X., and He, X. (2018). Anti-inflammatory steroidal glycosides from the berries of *Solanum nigrum* L. (European black nightshade). *Phytochemistry* 148, 87–96. doi:10.1016/j.phytochem.2018.01.019
13. Wu, X. W. (2011). A virtual screening research for the antitumor activity of the ingredients in LSYQD. Henan: Master, Zhengzhou University.
14. Yang, Y. (2014). The therapeutic basis and hepatoprotective activities of *Solanum nigrum* L. Jiangsu: Master, Nanjing agricultural University
15. Zhao, Y. (2010). Chemical constituents of two *Solanum* species, microbial transformation and biological activities. Shandong: Doctor, Shandong University
16. Wang, L. Y., Wang, N. L., and Yao, X. S. (2007). Non-Saponins from *Solanum nigrum* L. *Zhong Yao Cai* 30 (7), 792–794. doi:10.13863/j.issn1001-4454
17. Gao, S. H., Su, Z. Z., Yang, L. J., and Li, Z. Y. (2021). Chemical Components from Stems of *Solanum nigrum* by LC-MS and NMR. *Chin. Tradit. Herb. Drugs*. 52(5), 1263–1273. doi:10.7501/j.issn.0253-2670.2021.05.006
18. Zhou, X. L. (2006). Study on anticancer active component of *Solanum nigrum*. Liaoning: Doctor, Shenyang Pharmaceutical University.
19. Li, J. H., Li, S. Y., Shen, M. X., Qiu, R. Z., Fan, H. W., and Li, Y. B. (2021). Anti-Tumor Effects of *Solanum nigrum* L. Extraction on c6 High-Grade Glioma. *J. Ethnopharmacol.* 274 (1), 114034. doi:10.1016/j.jep.2021.114034
20. Elhag, R. A. M., El Badwi, S. M. A., Bakhiet, A. O., & Galal, M. (2011). Hepatoprotective activity of *Solanum nigrum* extracts on chemically induced liver damage in rats. *Journal of Veterinary Medicine and Animal Health*, 3(4), 45-50.
21. Mani, R. K., Paramashree, J. B., Bharathi, D. R., & Ahmed, S. S. (2022). The traditional and pharmacological properties of *Solanum nigrum*: a review. *International Journal of Indigenous Herbs and Drugs*, 49-55.
22. Perez G, R. M., Perez L, J. A., Garcia D, L. M., & Sossa M, H. (1998). Neuropharmacological activity of *Solanum nigrum* fruit. *Journal of ethnopharmacology*, 62(1), 43-48.]
23. Ding, X., Zhu, F., Yang, Y., & Li, M. (2013). Purification, antitumor activity in vitro of steroidal glycoalkaloids from black nightshade (*Solanum nigrum* L.). *Food chemistry*, 141(2), 1181-1186.]
24. Mohyuddin, A., Kurniawan, T. A., Khan, Z. U. D., Nadeem, S., Javed, M., Dera, A. A., ... & Saeed, S. (2022). Comparative insights into the antimicrobial, antioxidant, and nutritional potential of the *Solanum nigrum* complex. *Processes*, 10(8), 1455.
25. Muto, H., Nakatogawa, H., & Ito, K. (2006). Genetically encoded but nonpolypeptide prolyl-tRNA functions in the A site for SecM-mediated ribosomal stall. *Molecular cell*, 22(4), 545-552.
26. Ajayi, E. I. O. (2011). A review of the pharmacological aspects of *Solanum nigrum* Linn. ... and Molecular Biology....
27. Amer, M., Jayar, A. F., & Aljohani, M. S. Critical Review of the Level of Perspectives' Towards Work Risks among Health Care Providers.
28. Ahmed, A. H., & Ramzy, R. M. (1998). Seasonal variation in molluscicidal activity of *Solanum nigrum* L. *Journal of the Egyptian Society of Parasitology*, 28(3), 621-629.
29. Dua, V. K., Pandey, A. C., Raghavendra, K., Gupta, A., Sharma, T., & Dash, A. P. (2009). Larvicidal activity of neem oil (*Azadirachta indica*) formulation against mosquitoes. *Malaria Journal*, 8(1), 124.
30. Ahmed, A. S., Billings, B. K., Morton, R. M., & Stanford-Harris, M. (2002). The role of accounting conservatism in mitigating bondholder-shareholder conflicts over dividend policy and in reducing debt costs. *The accounting review*, 77(4), 867-890.
31. An, L., Tang, J. T., Liu, X. M., & Gao, N. N. (2006). Review about mechanisms of anti-cancer of *Solanum nigrum*. *Zhongguo Zhongyao zazhi = Zhongguo Zhongyao zazhi = China journal of Chinese materia medica*, 31(15), 1225-6.]
32. Kaushik, D., Jogpal, V., Kaushik, P., Lal, S., Saneja, A., Sharma, C., & Aneja, K. R. (2009). Evaluation of activities of *Solanum nigrum* fruit extract. *Archives of Applied Science Research*, 1(1), 43-50.
33. Ravi, V., Saleem, T. S. M., Maiti, P. P., Gauthaman, K., & Ramamurthy, J. (2009). Phytochemical and pharmacological evaluation of *Solanum nigrum* Linn. *Afr J Pharm Pharmacol*, 3(9), 454-457.
34. Zakaria, Z. A., Sulaiman, M. R., Morsid, N. A., Aris, A., Zainal, H., Mohd Pojan, N. H., & Hanan Kumar, G. (2009). Antinociceptive, anti-inflammatory and antipyretic effects of *Solanum nigrum* aqueous extract in animal models. *Methods and findings in experimental and clinical pharmacology*, 31(2), 81
35. Potawale, S. E., Sinha, S. D., Shroff, K. K., Dhalawat, H. J., Boraste, S. S., Gandhi, S. P., & Tondare, A. D. (2008). *Solanum nigrum* Linn: A phytopharmacological review. *Pharmacologyonline*, 3, 140-63.
36. Sankaran, M. (2012). Protective effect of *Solanum nigrum* fruit extract on the functional status of liver and kidney against ethanol induced toxicity. *Journal of Biochemical Technology*, 3(4), 339-343
37. Wannang, N. N., Anuka, J. A., Kwanashie, H. O., Gyang, S., & Auta, A. (2008). Anti-seizure activity of the aqueous leaf extract of *Solanum nigrum* Linn (solanaceae) in experimental animals. *African Health Sciences*, 8(2).