



## Monograph

### Echinacea

#### Introduction

The Echinacea plant is a member of the Compositae family; the three species of medicinal interest being *Echinacea angustifolia*, *Echinacea purpurea*, and *Echinacea pallida*. *Echinacea angustifolia* has been used therapeutically for centuries by Native Americans as a remedy for eye conditions, snake bites, insect stings, infected wounds, eczema, enlarged glands, mumps, and rabies. It was also used as a painkiller for a variety of conditions from stomach-aches to epilepsy. In the early 20th century, Echinacea was used by a group of physicians known as the “Eclectics,” whose medicinal practice relied primarily on the use of plants and their disease-healing properties. During the Eclectic era, Echinacea was used to treat a variety of kidney and urinary tract conditions, chronic bacterial infections, and syphilis.<sup>1</sup> From the 1930s-1970s, antibiotic development resulted in a sharp decline in Echinacea use, but due to a subsequent disenchantment with the medical establishment, an herbal medicine “renaissance” in the 1980s led to renewed interest in Echinacea’s benefits. Echinacea research during the last 20 years has focused on its immune-stimulating properties. Currently, Echinacea is being used to combat bacterial, viral, protozoan, and fungal infections, as an anti-inflammatory agent, and as a possible chemopreventative agent.

#### Description and Constituents

Echinacea is also known as purple coneflower, red sunflower, thimbleweed, and Rudbeckia. It is native to much of the United States, with locations varying by species, and is usually found in open meadows or damp locations such as woods, swamps, ditches, river banks, and low-lying thickets. It has a thick, black, pungent root, narrow leaves, and a stem growing to a height of three feet.<sup>2</sup> The flowering head is orange and cone-shaped, bearing purple, rose, or white petals from June to September.<sup>3</sup>

Active constituents vary slightly according to species and include caffeic acid derivatives (primarily echinocoid), flavonoids, essential oils, polyacetylenes, alkylamides, and polysaccharides. No single constituent has been found to be primarily responsible for Echinacea’s immune-stimulating effect; rather they appear to all work together to accomplish this. Therefore, extracts standardized to a specific echinocoid concentration may not be the most beneficial, as this standardization may be at the expense of the other active constituents.<sup>2</sup>

## Myths About Echinacea

Misinterpretation of the scientific literature regarding Echinacea's effect on the immune system has led to the development of several myths regarding Echinacea's therapeutic use including: (1) Echinacea is only appropriate for short-term use because it is not desirable to stimulate the immune system continuously,<sup>4</sup> and (2) Echinacea is an immune stimulator and as such, its use may be contraindicated in "progressive conditions" such as tuberculosis, leukemia, allergies, collagen disorders, multiple sclerosis, HIV/AIDS, and autoimmune disease.<sup>5</sup> However, the Native Americans' and Eclectics' high-quality, traditional-use data is a result of decades of extensive clinical experience, and does not support the suggested limitations. King<sup>2</sup> and Ellingwood<sup>6</sup> recommended long-term use of Echinacea for a variety of chronic conditions, including tuberculosis and autoimmune-related disorders. Similarly, neither modern research data nor authoritative herbal reference sources support the suggested limitations on Echinacea use. Numerous clinical studies of Echinacea have been conducted over the last 20-30 years that overwhelmingly demonstrate its therapeutic benefit and safety, even in patients with autoimmune disorders.<sup>7,8</sup> *The British Herbal Pharmacopoeia*,<sup>9</sup> *The British Herbal Compendium*,<sup>10</sup> and *The Encyclopedia of Common Natural Ingredients Used in Food, Drugs, and Cosmetics*<sup>11</sup> list no contraindications for Echinacea. Weiss also suggests Echinacea has an excellent safety profile and no side effects.<sup>12</sup>

## Echinacea and Immune Stimulation

Echinacea's immune-stimulating properties are quite complex and are attributed to the combined effect of several of its constituents.<sup>13</sup> The Eclectic physicians discovered alcohol extracts of Echinacea directly stimulated white blood cell production and phagocytic activity.<sup>6</sup> Modern clinical and *in vitro* research has confirmed the Eclectics' observations regarding increased phagocytosis,<sup>14</sup> NK cell activity, and increased antibody-dependent cellular cytotoxicity, mediated by tumor necrosis factor-alpha (TNF- $\alpha$ ). The latter study was conducted using peripheral blood mononuclear cells from normal individuals and patients with either chronic fatigue syndrome or AIDS.<sup>15</sup> Due to its potential to stimulate TNF- $\alpha$  and interleukins 1 and 6, it has been suggested that Echinacea should not be used by AIDS patients as it may speed the course of the disease, although is not a universally held theory.<sup>3,4</sup>

*Echinacea angustifolia* also appears to have a mild antibiotic effect, probably attributable to its caffeic acid constituent, which is capable of directly inhibiting *Staphylococcus aureus*. In addition, certain polyacetylene constituents of Echinacea have been found to be bacteriostatic against *E. coli* and *Pseudomonas aeruginosa*.<sup>13</sup> *E. angustifolia* was also used by the Eclectics to treat fungal and protozoan infections, most notably malaria and *Trichomonas vaginalis*,<sup>13</sup> although current research in this area is lacking.

Echinacea research during the past few years has primarily focused on its therapeutic benefit in treating symptoms of the common cold. A review of several clinical studies, comprised of over 3900 patients, demonstrates that Echinacea extracts decrease the frequency, symptoms, and severity of the common cold.<sup>16-19</sup> However, other similar studies (although fewer in number) have demonstrated Echinacea use to be of no significant benefit in lessening cold and flu symptoms.<sup>20,21</sup>

## Echinacea and Inflammation

Native American and Eclectic uses of Echinacea as an anti-inflammatory agent were centered around poisonous snake bites and insect stings. Caffeic acid derivatives, high molecular weight polysaccharides, flavonoids, and essential oils found in Echinacea all possess anti-inflammatory properties.<sup>13</sup> Although current research on Echinacea's anti-inflammatory effect is minimal, animal studies using *E.*

*angustifolia* have indicated the polysaccharide constituents of the extracts possess significant anti-inflammatory activity in attenuating paw and ear edema when applied topically to the skin of mice and rats.<sup>22-24</sup>

## Echinacea and Cancer

Research on Echinacea's benefit in cancer therapy is minimal and inconclusive, but because of its immune-stimulating properties, it may prove to be a useful adjunct to conventional cancer therapies. An animal study demonstrated Echinacea's ability to enhance cellular immunity in leukemic mice, resulting in a suppressive effect on leukemia, via increased production of endogenous interferon-gamma.<sup>25</sup> In another animal study, peritoneal macrophages from immunosuppressed mice incubated with Echinacea polysaccharides showed increased production of TNF- $\alpha$  and enhanced macrophage activation. The mice in this study also exhibited restored resistance to *Listeria monocytogenes* and *Candida albicans*, lethal infections associated with their immunosuppressed state.<sup>26</sup>

## Side Effects, Contraindications, and Herb/Drug Interactions

Historical use, modern research, and herbal reference publications show Echinacea to have an excellent safety profile. However, because it is an immune stimulant, caution should be used in combining it with immunosuppressive drugs such as corticosteroids, cyclosporine, amiodarone, methotrexate, and ketoconazole.<sup>27</sup> Echinacea use has been reported to occasionally cause reversible skin reactions, and for this reason, it should be used with caution in atopic individuals.<sup>28</sup>

## Dosage

Echinacea is available in several forms, and dosages vary accordingly. Typical dosages for the various forms are:<sup>29</sup>

Dried root = 0.5 to 1.0 grams three times daily

Tincture (1:5) = 1/2 to 1 teaspoon three times daily

Dry, powdered extract (standardized to 3.5% echinacoside) = 300 mg three times daily

Liquid Extract (1:1) = 1/4 to 1/2 teaspoon three times daily

Freeze Dried = 1 to 2 capsules or tablets three times daily

## References

1. Wagner H. Herbal immunostimulants. *Z Phytother* 1996;17:79-95.
2. King J. *The American Eclectic Dispensatory*. Cincinnati, OH: Moore, Wistach, and Keys;1854.
3. Bergner P. *The Healing Power of Echinacea and Goldenseal, and Other Immune System Herbs*. Rocklin, CA: Prima Publishing; 1997.
4. Bone K. Echinacea: When should it be used? *Altern Med Rev* 1997;2:451-458.
5. Bisset NG. *Herbal Drugs and Phytopharmaceuticals*. Wichtl M Ed., (German edition). Stuttgart/Boca Raton: Medpharm Scientific Publishers/CRC Press; 1994:182-184.
6. Ellingwood F. *American Materia Medica, Therapeutics and Pharmacognosy*. Chicago, IL: Ellingwood's Therapeutist; 1919.
7. Jurcic K, Melchart D, Holzmann M, Martin P, et al. Zwei Probandenstudien zur Stimulierung der Granulozytenphagozytose durch Echinacea-Extrakt-haltige Präparate. *Z Phytother* 1989;10:67-70.
8. Parnham MJ. Benefit-risk assessment of the squeezed sap of the purple coneflower (*Echinacea purpurea*) for long-term oral immunostimulation. *Phytomed* 1996;3:95-102.
9. British Herbal Medicine Association. *British Herbal Pharmacopoeia*. Cowling: BHMA; 1983:80-81.

10. British Herbal Medicine Association. *British Herbal Compendium. Vol 1*. Bournemouth: BHMA; 1992:81-83.
11. Leung AY, Forster S. *Encyclopedia of Common Natural Ingredients Used in Food, Drugs and Cosmetics. 2nd Ed.* New York-Chichester: John Wiley; 1996:216-220.
12. Weiss RF. *Herbal Medicine*. (Translated by Meuss AR from the Sixth German Edition of *Lehrbuch der Phytotherapie*). Beaconsfield: Beaconsfield Publishers Ltd; 1988:229-230.
13. Schar, D. *Echinacea, The Plant That Boosts Your Immune System*. London: Souvenir Press Ltd.; 1999.
14. Melchart D, Linde K, Worku F, et al. Results of five randomized studies on the immunomodulatory activity of preparations of Echinacea. *J Altern Complement Med* 1995;1:145-160.
15. See DM, Broumand N, Sahl L, Tilles JG. *In vitro* effects of Echinacea and ginseng on natural killer and antibody-dependent cell cytotoxicity in healthy subjects and chronic fatigue syndrome or acquired immunodeficiency syndrome patients. *Immunopharmacology* 1997;35:229-235.
16. Melchart D, Linde K, Fischer P, Kaesmayr J. Echinacea for preventing and treating the common cold. *Cochrane Database Syst Rev* 2000;CD000530.
17. Henneicke-von Zepelin H, Hentschel C, Schnitker J, et al. Efficacy and safety of a fixed combination phytomedicine in the treatment of the common cold (acute viral respiratory tract infection): results of a randomised, double blind, placebo controlled, multicentre study. *Curr Med Res Opin* 1999;15:214-227.
18. Brinkeborn RM, Shah DV, Degenring FH. Echinaforce and other Echinacea fresh plant preparations in the treatment of the common cold. A randomized, placebo controlled, double-blind clinical trial. *Phytomedicine* 1999;6:1-6.
19. Lindenmuth GF, Lindenmuth EB. The efficacy of Echinacea compound herbal tea preparation on the severity and duration of upper respiratory and flu symptoms: a randomized, double-blind placebo-controlled study. *J Altern Complement Med* 2000;6:327-334.
20. Turner RB, Riker DK, Gangemi JD. Ineffectiveness of Echinacea for prevention of experimental rhinovirus colds. *Antimicrob Agents Chemother* 2000;44:1708-1709.
21. Grimm W, Muller HH. A randomized controlled trial of the effect of fluid extract of *Echinacea purpurea* on the incidence and severity of colds and respiratory infections. *Am J Med* 1999;106:138-143.
22. Tragni E, Galli CL, Tubaro A, et al. Anti-inflammatory activity of *Echinacea angustifolia* fractions separated on the basis of molecular weight. *Pharmacol Res Commun* 1988;20:S87-S90.
23. Tubaro A, Tragni E, Del Negro P, et al. Anti-inflammatory activity of a polysaccharide fraction of *Echinacea angustifolia*. *J Pharm Pharmacol* 1987;39:567-569.
24. Tragni E, Tubaro A, Melis S, Galli CL. Evidence from two classic irritation tests for an anti-inflammatory action of a natural extract, Echinacina B. *Food Chem Toxicol* 1985;23:317-319.
25. Hayashi I, Ohotsuki M, Suzuki I, Watanabe T. Effects of oral administration of *Echinacea purpurea* (American herb) on incidence of spontaneous leukemia caused by recombinant leukemia viruses in AKR/J mice. *Nihon Rinsho Meneki Gakkai Kaishi* 2001;24:10-20.
26. Steinmuller C, Roesler J, Grottrup E, et al. Polysaccharides isolated from plant cell cultures of *Echinacea purpurea* enhance the resistance of immunosuppressed mice against systemic infections with *Candida albicans* and *Listeria monocytogenes*. *Int J Immunopharmacol* 1993;15:605-614.
27. Miller LG. Herbal medicinals: selected clinical considerations focusing on known or potential drug-herb interactions. *Arch Intern Med* 1998;158:2200-2211.
28. Bauer R, Hoheisel O, Stuhlfauth I, Wolf H. Extract of the *Echinacea purpurea* herb: an allopathic phytoimmunostimulant. *Wien Med Wochenschr* 1999;149:185-189.
29. Collins E, Berkoff N. *Everything You Need to Know About Echinacea and Immunity*. Roseville, CA: Prima Publishing; 1999:85-86.