

Template

 Entire dispensary

Review plan (10)

[< Back](#)

Hot Flash Support

 Template by Fullscript

Updated Sep 29th, 2023

Preview**Evidence**

Evidence rating

The following protocols were developed using only **a,b,c**-quality evidence

[Learn more about our rating scale](#) 

Overview

Approximately 50.3% to 82.1% of women going through menopause experience hot flashes and night sweats. ([Ziaei 2007](#)) To help manage symptoms, 40-50% of women rely on the use of complementary therapies such as plant-based supplements. ([Franco 2016](#))

Plant compounds called phytoestrogens have been found to have similar structures and functions when compared to the hormones within humans. ([Abdali 2010](#)) Estradiol is an estrogen steroid hormone that naturally drops throughout a woman's life cycle. Decreases in estradiol levels create vasomotor instability, which contributes to hot flashes. Utilizing plant compounds is theorized to help compensate for the estradiol drops through a woman's life cycle. ([Abdali 2010](#))

Lower rates of postmenopausal symptoms have been observed in Asian women. This thought to be due to the high dietary intake of soybeans, which contain isoflavones. Isoflavones are thought to work as phytoestrogens and have a high binding affinity to

estrogen beta receptors. ([Cheng 2007](#)) This sheds light on the mechanism behind how phytoestrogens are able to reduce hot flash frequency and vaginal dryness scores. ([Franco 2016](#))

Based on the current research findings, the ingredients in the protocol below have demonstrated efficacy in improving hot flashes associated with menopause.

Isoflavones

Isoflavones

40-80 mg, total isoflavones per day (dosing varies based on isoflavone), minimum 12 weeks ([Khaodhiar 2008](#)) ([van de Weijer 2002](#)) ([Cheng 2007](#)) ([Ferrari 2009](#)) ([Mainini 2013](#))

- A meta-analysis of 17 trials found reduced frequency (20.6%) and severity (26.2%) of hot flashes when compared to placebo; further analysis found isoflavones with 18.8 mg of genistein were twice as effective for reducing the frequency of hot flashes. ([Taku 2012](#))
- Daidzein-rich isoflavone supplementation in doses of 40 mg and 60 mg improved frequency of hot flashes by 43% and 41% after eight weeks, and 52% and 51% after 12 weeks, respectively, when compared to 32% and 39% with placebo, without any alteration of endogenous sex hormones or thyroid hormones. ([Khaodhiar 2008](#))
- For acute menopausal symptoms, 60 mg of isoflavones for six weeks decreased hot flashes by 57% and night sweats by 43%, with no other changes in serum levels of lipoproteins, estradiol, and follicular stimulating hormones when compared to that of placebo. ([Cheng 2007](#)).
- Mean number of hot flushes decreased by 36.2% and 41.2 % in the treatment group compared to 24.0% and 29.3% with placebo at week six and 12, respectively. ([Ferrari 2009](#))
- Red clover isoflavones decreased Kupperman Index score at three months and number of hot flushes at one month compared to placebo, with no effects on cardiovascular risk markers. ([Mainini 2013](#))
- Hot flush frequency decreased in a pooled mean analysis when compared to that of placebo, while having no statistically significant reported side effects. ([Chen 2015](#))

St. John's wort

St. John's wort (*Hypericum perforatum*)

900 mg, three times per day, minimum of 12 weeks ([Al-Akoum 2009](#)) ([Abdali 2010](#)) or as directed ([Eatamadnia 2019](#))

- Women aged 40 to 65 with three or more hot flashes per day experienced better quality of life as shown by improvements in baseline questionnaire after three months of supplementation. ([Al-Akoum 2009](#))
- Supplementation produced a significant decrease in Kupperman Index scores, as demonstrated by a reduction in the frequency and intensity of hot flashes when compared to placebo. ([Eatamadnia 2019](#))

Black cohosh

Black cohosh (*Actaea racemosa*)

100 mg, once per day, 12 weeks ([Gao 2018](#)) ([Frei-Kleiner 2005](#))

- Menopausal vasomotor symptoms reduced by 26% in pooled analysis among individuals taking black cohosh. [Shams 2010](#)
- All three treatment groups (black cohosh, estradiol valerate/progesterone, and estradiol valerate/medroxyprogesterone acetate) experienced decreased Kupperman menopause scores after 3 months; the black cohosh group experienced the lowest incidence of breast tenderness at 12.9% (vs. 36.7% and 14.3%, respectively) and vaginal bleeding at 6.5% (vs. 26.7% and 82.1%, respectively). ([Zheng 2013](#))
- In a trial of 122 menopausal women lasting 12 weeks, women with an initial Kupperman Index equal or greater than 20 experienced a 47% reduction in scores from a dried ethanolic extract group, compared to 21% in the placebo group. ([Frei-Kleiner 2005](#))
- In a trial of 120 menopausal women lasting six months, both the black cohosh and fluoxetine treatment groups experienced significantly decreased Kupperman Index and Beck's depression scale scores; however, monthly scores for hot flashes and night sweats decreased by 85% in the black cohosh group, compared to 62% in the fluoxetine group. ([Oktem 2007](#))

Rhubarb extract

Rhubarb extract – ERr 731

4 mg, once per day, 12 weeks ([Kaszkin-Bettag 2009](#)) ([Heger 2006](#)) ([Hasper 2009](#))

- Menopause Rating Scale (MRS) decreased overall as shown by a decrease in number and severity of hot flashes in treatment group when compared to placebo. ([Kaszkin-Bettag 2009](#))

- Total score for MRS II decreased as shown by a decrease in the severity of hot flashes, and an increase of menopause-specific quality of life; no adverse effects were found as indicated by a lack of findings through endometrial biopsies, bleeding, weight changes, blood pressure, and heart rate. ([Heger 2006](#))

Vitamin E

Vitamin E

400-800 IU, total per day, minimum 4 weeks ([Ziaei 2007](#)) ([Barton 1998](#))

- Severity score and frequency of hot flashes were significantly decreased when compared to placebo. ([Ziaei 2007](#))
- Supplementation reduced the frequency of hot flashes by one per day compared to the placebo. ([Barton 1998](#))

Vitex

Vitex (*Vitex agnus-castus*)

30 mg *Vitex agnus-castus* extracts for 8 weeks ([Zahra 2011](#)) ([Naseri 2019](#))

- A randomized double-blind clinical trial of 52 women found that vitex significantly reduced total menopausal symptoms, anxiety, and vasomotor dysfunction compared to placebo. ([Naseri 2019](#))
- A clinical trial of 60 postmenopausal women found that daily Vitex agnus-castus significantly reduced the frequency of hot flashes compared to placebo over 8 weeks, as measured by the Blatt-Kupperman Index. ([Zahra 2011](#))
- One study found that adding a combination of *Nigella sativa* and *Vitex agnus-castus* to citalopram significantly improved vasomotor, physical, and psychosocial symptoms in menopausal women compared to citalopram alone. ([Molaie 2019](#))

References

1. Abdali, K., Khajehei, M., & Tabatabaee, H. R. (2010). Effect of St John's wort on severity, frequency, and duration of hot flashes in premenopausal, perimenopausal and postmenopausal women: a randomized, double-blind, placebo-controlled study. *Menopause*, 17(2), 326–331. <https://pubmed.ncbi.nlm.nih.gov/20216274> (**B**)
2. Al-Akoum, M., Maunsell, E., Verreault, R., Provencher, L., Otis, H., & Dodin, S. (2009). Effects of Hypericum perforatum (St. John's wort) on hot flashes and quality of life in

- perimenopausal women: a randomized pilot trial. *Menopause* , 16(2), 307–314. <https://pubmed.ncbi.nlm.nih.gov/19194342/> (C)
3. Barton, D. L., Loprinzi, C. L., Quella, S. K., Sloan, J. A., Veeder, M. H., Egner, J. R., Fidler, P., Stella, P. J., Swan, D. K., Vaught, N. L., & Novotny, P. (1998). Prospective evaluation of vitamin E for hot flashes in breast cancer survivors. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 16(2), 495–500. <https://pubmed.ncbi.nlm.nih.gov/9469333/> (C)
 4. Chen, M.-N., Lin, C.-C., & Liu, C.-F. (2015). Efficacy of phytoestrogens for menopausal symptoms: a meta-analysis and systematic review. *Climacteric: The Journal of the International Menopause Society*, 18(2), 260–269. <https://pubmed.ncbi.nlm.nih.gov/25263312/> (A)
 5. Cheng, G., Wilczek, B., Warner, M., Gustafsson, J.-A., & Landgren, B.-M. (2007). Isoflavone treatment for acute menopausal symptoms. *Menopause* , 14(3 Pt 1), 468–473. <https://pubmed.ncbi.nlm.nih.gov/17290160/> (B)
 6. Eatemadnia, A., Ansari, S., Abedi, P., & Najjar, S. (2019). The effect of Hypericum perforatum on postmenopausal symptoms and depression: A randomized controlled trial. *Complementary Therapies in Medicine*, 45, 109–113. <https://pubmed.ncbi.nlm.nih.gov/31331546/> (C)
 7. Ferrari, A. (2009). Soy extract phytoestrogens with high dose of isoflavones for menopausal symptoms. *The Journal of Obstetrics and Gynaecology Research*, 35(6), 1083–1090. <https://pubmed.ncbi.nlm.nih.gov/20025635/> (B)
 8. Franco, O. H., Chowdhury, R., Troup, J., Voortman, T., Kunutsor, S., Kavousi, M., Oliver-Williams, C., & Muka, T. (2016). Use of Plant-Based Therapies and Menopausal Symptoms: A Systematic Review and Meta-analysis. *JAMA: The Journal of the American Medical Association*, 315(23), 2554–2563. <https://pubmed.ncbi.nlm.nih.gov/27327802/> (A)
 9. Frei-Kleiner, S., Schaffner, W., Rahlfs, V. W., Bodmer, C., & Birkhäuser, M. (2005). Cimicifuga racemosa dried ethanolic extract in menopausal disorders: a double-blind placebo-controlled clinical trial. *Maturitas*, 51(4), 397–404. <https://pubmed.ncbi.nlm.nih.gov/16039414/> (B)
 10. Gao, L., Zheng, T., Xue, W., Wang, Y., Deng, Y., Zuo, H., & Sun, A. (2018). Efficacy and safety evaluation of Cimicifuga foetida extract in menopausal women. *Climacteric: The Journal of the International Menopause Society*, 21(1), 69–74. <https://pubmed.ncbi.nlm.nih.gov/29198157/> (C)
 11. Hasper, I., Ventskovskiy, B. M., Rettenberger, R., Heger, P. W., Riley, D. S., & Kaszkin-Bettag, M. (2009). Long-term efficacy and safety of the special extract ERr 731 of Rheum rhaponticum in perimenopausal women with menopausal symptoms. *Menopause* , 16(1), 117–131. <https://pubmed.ncbi.nlm.nih.gov/18978638/> (C)

12. Heger, M., Ventskovskiy, B. M., Borzenko, I., Kneis, K. C., Rettenberger, R., Kaszkin-Bettag, M., & Heger, P. W. (2006). Efficacy and safety of a special extract of *Rheum rhaponticum* (ERr 731) in perimenopausal women with climacteric complaints: a 12-week randomized, double-blind, placebo-controlled trial. *Menopause*, 13(5), 744–759. <https://pubmed.ncbi.nlm.nih.gov/16894335/> **(B)**
13. Kaszkin-Bettag, M., Ventskovskiy, B. M., Solskyy, S., Beck, S., Hasper, I., Kravchenko, A., Rettenberger, R., Richardson, A., & Heger, P. W. (2009). Confirmation of the efficacy of ERr 731 in perimenopausal women with menopausal symptoms. *Alternative Therapies in Health and Medicine*, 15(1), 24–34. <https://pubmed.ncbi.nlm.nih.gov/19161045/> **(C)**
14. Khaodhiar, L., Ricciotti, H. A., Li, L., Pan, W., Schickel, M., Zhou, J., & Blackburn, G. L. (2008). Daidzein-rich isoflavone aglycones are potentially effective in reducing hot flashes in menopausal women. *Menopause*, 15(1), 125–132. <https://pubmed.ncbi.nlm.nih.gov/18257146/> **(B)**
15. Mainini, G., Torella, M., Di Donna, M. C., Esposito, E., Ercolano, S., Correa, R., Cucinella, G., Stradella, L., Luisi, A., Basso, A., Cerreto, F. V., Cicatiello, R., Matteo, M., & De Franciscis, P. (2013). Nonhormonal management of postmenopausal women: effects of a red clover based isoflavones supplementation on climacteric syndrome and cardiovascular risk serum profile. *Clinical and Experimental Obstetrics & Gynecology*, 40(3), 337–341. <https://pubmed.ncbi.nlm.nih.gov/24283160/> **(C)**
16. Oktem, M., Eroglu, D., Karahan, H. B., Taskintuna, N., Kuscü, E., & Zeyneloglu, H. B. (2007). Black cohosh and fluoxetine in the treatment of postmenopausal symptoms: a prospective, randomized trial. *Advances in Therapy*, 24(2), 448–461. <https://pubmed.ncbi.nlm.nih.gov/17565936/> **(C)**
17. Shams, T., Setia, M. S., Hemmings, R., McCusker, J., Sewitch, M., & Ciampi, A. (2010). Efficacy of black cohosh-containing preparations on menopausal symptoms: a meta-analysis. *Alternative Therapies in Health and Medicine*, 16(1), 36–44. <https://pubmed.ncbi.nlm.nih.gov/20085176/> **(A)**
18. Taku, K., Melby, M. K., Kronenberg, F., Kurzer, M. S., & Messina, M. (2012). Extracted or synthesized soybean isoflavones reduce menopausal hot flash frequency and severity: systematic review and meta-analysis of randomized controlled trials. *Menopause*, 19(7), 776–790. <https://pubmed.ncbi.nlm.nih.gov/22433977/> **(A)**
19. Van de Weijer, P. H. M., & Barentsen, R. (2002). Isoflavones from red clover (Promensil) significantly reduce menopausal hot flush symptoms compared with placebo. *Maturitas*, 42(3), 187–193. <https://pubmed.ncbi.nlm.nih.gov/12161042/> **(C)**
20. Zheng, T.-P., Sun, A.-J., Xue, W., Wang, Y.-P., Jiang, Y., Zhang, Y., & Lang, J.-H. (2013). Efficacy and safety of *Cimicifuga foetida* extract on menopausal syndrome in Chinese women. *Chinese Medical Journal*, 126(11), 2034–2038. <https://pubmed.ncbi.nlm.nih.gov/23769553/> **(C)**

21. Ziaei, S., Kazemnejad, A., & Zareai, M. (2007). The effect of vitamin E on hot flashes in menopausal women. *Gynecologic and Obstetric Investigation*, 64(4), 204–207.
<https://pubmed.ncbi.nlm.nih.gov/17664882/> (B)

Disclaimer: Protocols are intended solely as an informational reference tool for practicing health care professionals. The content provided is not intended to be for medical diagnosis or treatment, is not a substitute for your professional judgment, and is not meant to provide you medical or professional advice. You should evaluate and independently confirm the appropriateness of the content provided, including verifying uses, dosages, warnings and contraindications on product labels, and rely on your experience and judgment and other available resources when applying the provided content to an actual client care situation. While content has been obtained from sources believed to be reliable, we cannot and do not guarantee the accuracy, validity, timeliness or completeness of the content. We make no representation or warranty, express or implied, including, without limitation, any warranty of merchantability or of fitness for a particular purpose, and you assume full responsibility for the use of the content and products, and agree that Fullscript and its content providers are not responsible or liable for any claim, loss, injury or damage arising from your use of the information. Statements regarding dietary and other health care supplements have not been evaluated by the FDA, and are not intended to diagnose, treat, cure, or prevent any disease.

Template

Total starts at

\$198.34

Add to plan

ⓘ We won't overwrite any existing dosage information.

*These statements have not been evaluated by the Food and Drug Administration. These products are not intended to diagnose, treat, cure, or prevent any disease.

†Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

© Fullscript 2025. All rights reserved.

[Terms of Service](#) [Privacy Statement](#) [Privacy Rights Notice](#)
[Help and Support](#)

