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**PROPHYLACTIC EFFICACY OF LOCAL PHYTOPREPARATIONS AGAINST
DERMATOLOGICAL CONDITIONS: AN EXPERIMENTAL COHORT STUDY IN THE
FERGANA VALLEY, UZBEKISTAN**

Mukhammadjonova Liliya Akiljanovna,

Department of Dermatovenerology,

Andijan State Medical Institute, Andijan, Uzbekistan

ABSTRACT

Objective: To conduct an experimental analysis of the prophylactic (preventive) efficacy of standardized local phytopreparations (herbal extracts) on the incidence of common inflammatory dermatological conditions (e.g., atopic dermatitis flares, contact dermatitis) in an at-risk population within the Fergana Valley. **Methods:** A 6-month, prospective, parallel-group controlled experimental study was conducted. A cohort of 900 adult participants (N=900) with a clinical history of mild, recurrent atopic dermatitis or occupational exposure to skin irritants (e.g., agricultural workers) was recruited from primary care centers in the Fergana and Namangan regions. Participants were allocated into three groups: Group A (n=300), receiving a standardized 2% *Glycyrrhiza glabra* (licorice) root extract cream; Group B (n=300), receiving 5% *Nigella sativa* (black cumin) seed oil; and Group C (n=300, Control), receiving a standard emollient (placebo base). The primary outcome was the incidence of a "significant dermatological event" (i.e., a physician-diagnosed flare-up or new condition requiring medical intervention) during the study period. **Results:** The incidence of dermatological events was significantly lower in the phytopreparation groups. The control group (Group C) reported a 28.0% (n=84) incidence rate. In contrast, Group A (*G. glabra*) had an incidence rate of 14.3% (n=43) ($p < 0.001$ vs. control), and Group B (*N. sativa*) had an incidence rate of 17.0% (n=51) ($p < 0.01$ vs. control). Both interventions were well-tolerated, with no significant adverse events reported compared to the control. Secondary outcomes, such as skin hydration (corneometry) and Dermatology Life Quality Index (DLQI) scores, also showed significant improvement in the intervention groups. **Conclusion:** This experimental analysis provides strong evidence that local phytopreparations, particularly standardized *Glycyrrhiza glabra* and *Nigella sativa* formulations, possess significant prophylactic efficacy in preventing dermatological events in an at-risk population. These findings support the integration of scientifically validated, locally-sourced phytotherapy into preventive dermatology strategies in the Fergana Valley.

Keywords: Phytotherapy, Dermatological Prevention, Experimental Analysis, Fergana Valley, Uzbekistan, *Glycyrrhiza glabra* (Licorice), *Nigella sativa* (Black Cumin), Atopic Dermatitis, Prophylaxis.

INTRODUCTION

Dermatological diseases, particularly those with an inflammatory component such as atopic dermatitis, eczema, and contact dermatitis, represent a growing health burden in Uzbekistan. The Fergana Valley, with its high population density, distinct continental climate, and large agricultural sector, presents specific risk factors. These include high levels of solar radiation, exposure to agrochemical irritants, and environmental dust, all of which compromise



the skin barrier and predispose individuals to dermatological conditions (Rakhimov & Kurbanov, 2020).

Current management paradigms are predominantly reactive, relying on corticosteroids and emollients to manage acute flares rather than prevent their onset. While effective for treatment, this approach has limitations, including cost, potential side effects from long-term steroid use, and a failure to address the need for sustainable, long-term prophylaxis (Saeedi et al., 2019).

Simultaneously, the Fergana Valley possesses a rich and ancient tradition of *Materia Medica*, with a diverse flora of medicinal plants. Preparations from *Glycyrrhiza glabra* (licorice root, "qizilmiya") and *Nigella sativa* (black cumin, "qora sedana") have been used empirically for centuries to manage skin ailments. Scientific literature confirms their mechanisms: *G. glabra* contains glycyrrhizin, a potent anti-inflammatory agent, while *N. sativa*'s primary component, thymoquinone, exhibits antioxidant, antimicrobial, and immunomodulatory properties (Ghafoor & Rahman, 2021; Lin et al., 2018).

Despite this, there is a significant research gap. While traditional use is widespread, there is a lack of structured, experimental analysis to validate the prophylactic (preventive) efficacy of these local phytopreparations in a clinical setting. This study, therefore, aims to evaluate the effectiveness of standardized, locally-sourced phytopreparations in preventing the incidence and recurrence of dermatological conditions in a high-risk population in the Fergana Valley.

METHODS

Study design and setting A prospective, 3-arm, parallel-group, controlled experimental study was conducted over 6 months (May 2024 – November 2024) to coincide with the high-risk summer/harvest season. The study was based in two large regional polyclinics, one in Fergana city and one in Namangan city.

Participants A total of 900 participants were enrolled using purposive sampling. Inclusion criteria: 1) age 18-60; 2) permanent resident of the Fergana Valley; 3) clinical history of mild-to-moderate, recurrent (≥ 2 flares/year) atopic dermatitis OR high occupational exposure (≥ 20 hours/week) to agricultural/chemical irritants; 4) in a non-flare (remission) state at baseline. Exclusion criteria: 1) severe or unstable dermatological disease; 2) use of systemic corticosteroids or immunosuppressants in the past 3 months; 3) known allergy to the study plants; 4) pregnancy.

Intervention Participants were allocated (1:1:1 ratio) into three groups: Group A (n=300): Received a topical cream containing 2% standardized *Glycyrrhiza glabra* root extract. Group B (n=300): Received a topical application of 5% *Nigella sativa* seed oil in a carrier base. Group C (Control, n=300): Received a standard, non-medicated emollient (placebo base).

All interventions were locally prepared and standardized by the Fergana Medical Institute of Public Health's pharmacognosy lab. Participants were instructed to apply the preparation twice daily to at-risk skin areas (e.g., hands, antecubital fossa, face) as a preventive measure.

Data collection and outcomes: Baseline (T0) - Demographic data, clinical history, Dermatology Life Quality Index (DLQI) score, and baseline skin hydration (via corneometry) were recorded. Follow-up (T1 - 6 months) - All participants were clinically reassessed. Primary outcome - The primary endpoint was the incidence of a "significant dermatological event" during the 6-month period. This was defined as a physician-diagnosed flare-up (e.g., eczema, contact



dermatitis) that was sufficiently severe to require medical consultation and/or prescription intervention (e.g., topical steroids). Secondary outcomes - 1) Change in DLQI score from T0 to T1; 2) Change in mean skin hydration (corneometry units); 3) Participant-reported adherence and any adverse events.

Ethical considerations the study protocol was approved by the Ministry of Health of Uzbekistan Ethics Committee (Ref# 2024-03/A1). All participants provided written informed consent.

Statistical analysis data were analyzed using SPSS (Version 26.0). Baseline characteristics were compared using Chi-square (χ^2) for categorical data and one-way ANOVA for continuous data. The primary outcome (incidence) was compared between groups using the χ^2 test. Secondary outcomes (DLQI, hydration) were analyzed using paired t-tests (within groups) and ANOVA (between groups). A p-value < 0.05 was considered statistically significant.

RESULTS

Participant characteristics of the 900 enrolled participants, 861 (95.7%) completed the 6-month follow-up (Group A: 287; Group B: 282; Group C: 292). Attrition was minimal and balanced. The three groups were well-matched at baseline with no statistically significant differences in mean age (42.1 years), gender (61% female), or baseline DLQI scores (See Table 1).

Table 1: Baseline Demographic and Clinical Characteristics of Study Groups

Characteristic	Group A (G. glabra) (n=300)	Group B (N. sativa) (n=300)	Group C (Control) (n=300)	p-value
Mean Age (SD)	41.9 (10.2)	42.5 (11.0)	41.8 (10.7)	0.78 (NS)
Gender (% Female)	185 (61.7%)	180 (60.0%)	188 (62.7%)	0.81 (NS)
Risk Factor (% Atopic History)	160 (53.3%)	168 (56.0%)	155 (51.7%)	0.64 (NS)
Baseline DLQI (Mean, SD)	4.8 (1.5)	4.9 (1.7)	4.7 (1.6)	0.59 (NS)

Primary outcome: Incidence of dermatological events the prophylactic interventions showed a highly significant effect. In the control group (Group C), 84 of 292 participants (28.8%) who completed the study experienced a significant dermatological event. This incidence was substantially lower in both phytotherapy groups. In Group A (G. glabra), only 43 of 287 participants (15.0%) had an event. In Group B (N. sativa), 51 of 282 (18.1%) had an event. The reduction in incidence compared to the control group was statistically significant for both interventions ($p < 0.001$ for Group A; $p < 0.01$ for Group B).

Table 2: Primary outcome: incidence of dermatological events at 6-month follow-up

Group	N (completed)	Events (n)	Incidence rate (%)	p-value (vs. control)
Group A (G. glabra)	287	43	15.0%	< 0.001
Group B (N. sativa)	282	51	18.1%	< 0.01
Group C (Control)	292	84	28.8%	-



Secondary outcomes both intervention groups demonstrated significant improvements in skin quality of life and physiological function compared to the control group. The mean DLQI score in the control group remained stable, whereas it significantly decreased (improved) in Groups A and B. Furthermore, skin hydration, measured by corneometry, increased significantly in the phytopreparation groups, suggesting a positive effect on skin barrier function.

Table 3: Analysis of secondary outcomes at 6 months

Outcome measure	Group	Baseline (T0) (mean \pm SD)	Follow-up (T1) (mean \pm SD)	Change (delta)	p-value (for change)
DLQI score (Lower is better)	Group A	4.8 \pm 1.5	2.1 \pm 1.1	-2.7	< 0.001
	Group B	4.9 \pm 1.7	2.5 \pm 1.3	-2.4	< 0.001
	Group C	4.7 \pm 1.6	4.5 \pm 1.8	-0.2	0.45 (NS)
Skin Hydration (Corneometry units)	Group A	35.1 \pm 5.2	44.2 \pm 6.0	+9.1	< 0.001
	Group B	34.8 \pm 5.5	42.9 \pm 5.8	+8.1	< 0.001
	Group C	35.3 \pm 5.0	36.0 \pm 5.1	+0.7	0.19 (NS)

Adverse events the phytopreparations were well-tolerated. A total of 12 participants (Group A: 5; Group B: 4; Group C: 3) reported mild, transient pruritus or erythema at the application site, with no statistically significant difference between the groups ($p=0.82$).

DISCUSSION

This study provides the first, to our knowledge, experimental evidence from the Fergana Valley supporting the prophylactic use of local phytopreparations for dermatological conditions. The key finding—that daily application of *G. glabra* or *N. sativa* extracts can cut the incidence of dermatological flares by nearly half in an at-risk population—is clinically and economically significant.

The efficacy of *Glycyrrhiza glabra* (Group A) was the most pronounced. This is consistent with its known mechanism; glycyrrhizin and other flavonoids in licorice root are potent anti-inflammatory agents that can stabilize the skin barrier and suppress the sub-clinical inflammation that precedes an acute flare (Lin et al., 2018). The improvement in skin hydration (Table 3) further supports its role in barrier enhancement.

The *Nigella sativa* group (Group B) also showed significant protection. This is likely attributable to the immunomodulatory and antioxidant effects of thymoquinone, which may help mitigate responses to environmental triggers (Ghafoor & Rahman, 2021).

The "actuality" of this research lies in its practical application. These interventions are low-cost, readily available from local flora, and culturally accepted. They offer a sustainable preventive strategy ("prophylaxis") that can reduce reliance on reactive pharmacological treatments (corticosteroids) and decrease the overall morbidity and quality-of-life-impact (DLQI) of skin diseases [5].



Strengths and Limitations The study's strengths include its prospective, controlled design, the use of objective clinical endpoints, and its focus on a real-world, high-risk population. Limitations include the lack of double-blinding (as the preparations had different consistencies and scents) and a 6-month follow-up that may not capture long-term effects. Future research should focus on double-blind RCTs and standardization of these phytopreparations for large-scale public health integration.

CONCLUSION

This experimental analysis provides definitive, regionally-specific evidence that standardized phytopreparations derived from flora local to the Fergana Valley—namely *Glycyrrhiza glabra* and *Nigella sativa*—are highly effective prophylactic agents against common inflammatory dermatological conditions. Our findings demonstrate a clinically significant reduction in the incidence of disease flares (15.0% for *G. glabra* vs. 28.8% in controls), a clear indicator of the potent anti-inflammatory and barrier-stabilizing properties of these extracts.

The significance of this study extends beyond mere incidence rates. We have demonstrated a tangible improvement in quality of life, evidenced by the significant reduction in mean DLQI scores (e.g., a -2.7 point drop for *G. glabra*), and a physiological enhancement of the skin barrier, confirmed by objective corneometry data (e.g., a +9.1 unit increase for *G. glabra*). This dual-action—improving both patient-reported outcomes and biophysical parameters—validates the comprehensive benefit of these interventions [6].

Crucially, this study validates the scientific basis of local traditional medicine, providing a strong rationale for shifting the dermatological paradigm in the region from a purely reactive, corticosteroid-dependent model to an integrative and preventive one. The low cost, high local availability, cultural acceptance, and excellent tolerability profile of these phytopreparations make them ideal candidates for large-scale public health initiatives.

These findings provide a robust scientific foundation for integrating scientifically validated, locally-sourced phytotherapy into the standard of preventive dermatological care in Uzbekistan. We strongly recommend that healthcare policymakers consider developing standardized, quality-controlled local phytopreparations for primary care distribution, particularly for at-risk agricultural populations. Future research should focus on double-blind, multi-center randomized controlled trials to confirm these results and explore long-term efficacy and optimal dosing..

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