



P110. EFFICACY OF A CORIOLUS VERSICOLOR-BASED VAGINAL GEL TO REPAIR CERVICAL MUCOSA WITH HPV LESIONS. PRELIMINARY RESULTS OF A CLINICAL TRIAL

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Context: Papilocare[®] is a non-hormonal Coriolus versicolor-based vaginal gel. In patients with HPV infection, a regression effect of the LSIL in cervix and a significant virus negativization have been observed after 1-year administration of oral Coriolus versicolor.

Objective: To evaluate the efficacy of Papilocare[®] to repair cervical mucosa in women with precancerous HPV lesions and consistent colposcopy image.

Methods: Pilot, randomized, open, parallel group, controlled clinical trial.

Patients: Currently recruiting 96 positive-HPV women age 30 to 65 with pap result of ASC-US, LSIL or AG-US and concordant colposcopy image.

Interventions: Patients are randomized to 3 groups: A) Papilocare[®] 1 cannula/day for 1 month + 1 cannula/alternate days for 5 months; B) Papilocare[®] 1 cannula/day for 3 months + 1 cannula/alternate days for 3 months; C) no treatment as usual practice.

Main outcome measures: Preliminary analysis of primary endpoint (% of patients with normal pap and concordant colposcopy image) at 3 and 6 months and the secondary endpoint (% of patients with HPV cleared) at 6 months are presented. Papilocare[®] arms (A+B) were combined for this evaluation.

Results: Data from 41 patients at 3 months and 25 at 6 months are available. At 3 months, 69.2% of patients using Papilocare[®] (n=26) negativized pap and colposcopy vs. 33.3% in control group (n=15) (p=0.048; Fisher test). At 6 months, 73% (n=15) vs 60% (n=10) of patients negativized cervical lesions for Papilocare[®] and control groups, respectively (p=ns).

The % of patients who cleared the HPV at 6 months was 56% in Papilocare[®] group vs 30% in control group (p=ns).

Conclusion: In these preliminary results, Papilocare[®] shows a significative difference in repairing HPV-cervical lesions at 3 months versus control and a positive trend in both HPV clearance and HPV-cervical lesions repairing at 6 months; these findings need to be confirmed upon study completion.

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