# EFFICACY OF A MULTI-INGREDIENT *CORIOLUS VERSICOLOR*-BASED VAGINAL GEL IN HIGH-RISK HPV INFECTED PATIENTS: RESULTS OF 6 DIFFERENT STUDIES

#### Authors:

Cortés J<sup>1</sup>, Dexeus D<sup>2</sup>, Palacios S<sup>3</sup>, Serrano L<sup>4</sup>, Gajino C<sup>5</sup>, Marín E<sup>6</sup>, Criscuolo A<sup>7</sup>, Riera M<sup>8</sup>, Gaslain Y<sup>9</sup>

<sup>1</sup>Private Practice, Palma, Spain, <sup>2</sup>Clínica Ginecológica Women's, Barcelona, Spain, <sup>3</sup>Instituto Palacios de Salud y Medicina de la Mujer, Madrid, Spain, <sup>4</sup>Policlínico HM Gabinete Velázquez, Madrid, Spain, <sup>5</sup>Hospital Materno Infantil Teresa Herrera, A Coruña, Spain, <sup>6</sup>Hospital Álvaro Cunqueiro, Vigo, Spain, <sup>7</sup>Tor Vergata University Hospital, Rome, Italy, <sup>8</sup>Institut Català de la Salut, Barcelona, Spain, <sup>9</sup>Procare Health Iberia, Barcelona, Spain

#### INTRODUCTION:

Papilocare® is a patented multi-ingredient vaginal gel which includes *Coriolus versicolor* and other active ingredients. It is approved in Europe with the indication of re-epitelization for the cervical transformation zone for the prevention and co-adjuvant treatment of HPV-induced lesions.

## **OBJECTIVES:**

The main objective is to evaluate the consistency of the efficacy of a multi-ingredient *Coriolus versicolor*-based vaginal gel on high-risk HPV (HR-HPV) clearance in 6 different studies.

- Coruña study: Retrospective one-cohort. Primary endpoint (PE), HR-HPV clearance assessed in 86 medical patients' records.
- Hospitalet study: Retrospective one-cohort. PE,
  Composite efficacy variable (patients with normal cytology and/or HPV clearance) in 91 HR-HPV patients.
- Roma study: Retrospective controlled. PE, HR-HPV clearance in 183 patients.
- Paloma trial: SE, HR-HPV clearance in 91 patients.
- PapilOBS study: SE, HR-HPV clearance in 192 patients.

#### **METHODS:**

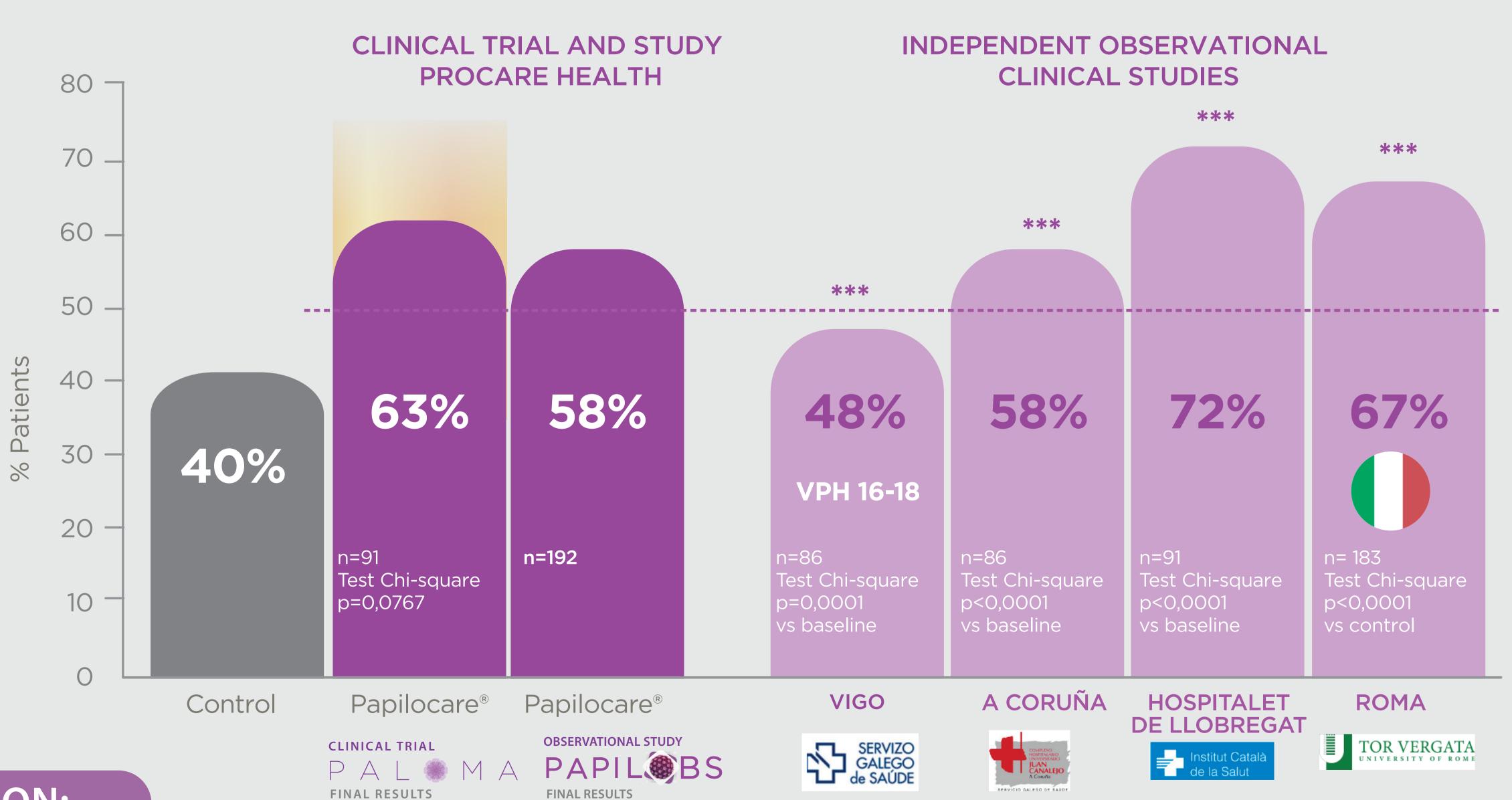
Results from 4 independent observational studies (6 month-treatment period with Papilocare®) were compared to results from a randomized, open, parallel, controlled trial (Paloma: NCT04002154) and an observational, multicenter, prospective, one-cohort study (PapilOBS: NCT04199260).

• Vigo study: Prospective one-cohort. Secondary endpoint (SE), HPV clearance in 86 patients infected by HPV 16 and/or 18.

#### **RESULTS:**

48% of patients cleared 16-/18-HPV in Vigo study. 58% of reduction was observed in the number of HR-HPV patients (Coruña) and 72.5% normalized cytology and/or cleared HR-HPV (Hospitalet). 67% HR-HPV clearance was observed (treated group) vs 37.2% (control group), in the Roma study. In the Paloma trial, HR-HPV clearance reached 63% (treated group) vs 40% (control group). 57.4% HR-HPV clearance was observed in the PapilOBS study.

### HR-HPV CLEARANCE AT 6 MONTHS RESULTS IN DIFFERENTS STUDIES



### **CONCLUSION:**

Papilocare® has shown significant consistent rates of efficacy with a 64% of HR-HPV clearance in weighted average in 6 different studies involving more than 700 patients. These data reinforce the beneficial effect observed in HR-HPV patients.