

## Background & Objective






- BV is a common vaginal infection with a prevalence of 29% among US women ages 14-49 years.<sup>1</sup>
- More than 50% of patients have recurrent BV within 6-12 months of initial treatment.<sup>2,4</sup>
- This study summarized patient characteristics, treatment patterns, HCRU, and costs among commercial insured patients diagnosed with BV.

## Study Design & Methods

- Retrospective cohort study using the Merative™ MarketScan® Commercial Database.
- Patients were 12-49 years old with an incident BV diagnosis (ICD-10: N76.0 or N76.1 “vaginitis”) from 01 JAN 2017 through 30 SEP 2020 and ≥1 BV med (metronidazole, clindamycin, secnidazole, tinidazole).
- Baseline and follow-up periods were one year before and after incident diagnosis, respectively.
- Recurrent BV was defined as ≥2 treatment courses.
- Analyses included descriptive statistics, generalized linear model with gamma distribution (cost), and Poisson regression (treatment courses) using SAS.
- Models included: geographic region, age group, pregnancy status, preterm labor, post procedural gynecological infection, pelvic inflammatory disease, any sexually transmitted infection, female infertility, and Charlson Comorbidity Index\*.

\*Includes myocardial infarction, congestive heart failure, peripheral vascular disease, cerebrovascular disease, dementia, chronic pulmonary disease, connective tissue-rheumatic disease, peptic ulcer disease, mild liver disease, diabetes w/o complications, diabetes w/ complications, paraplegia and hemiplegia, renal disease, cancer, moderate or severe liver disease, metastatic carcinoma.

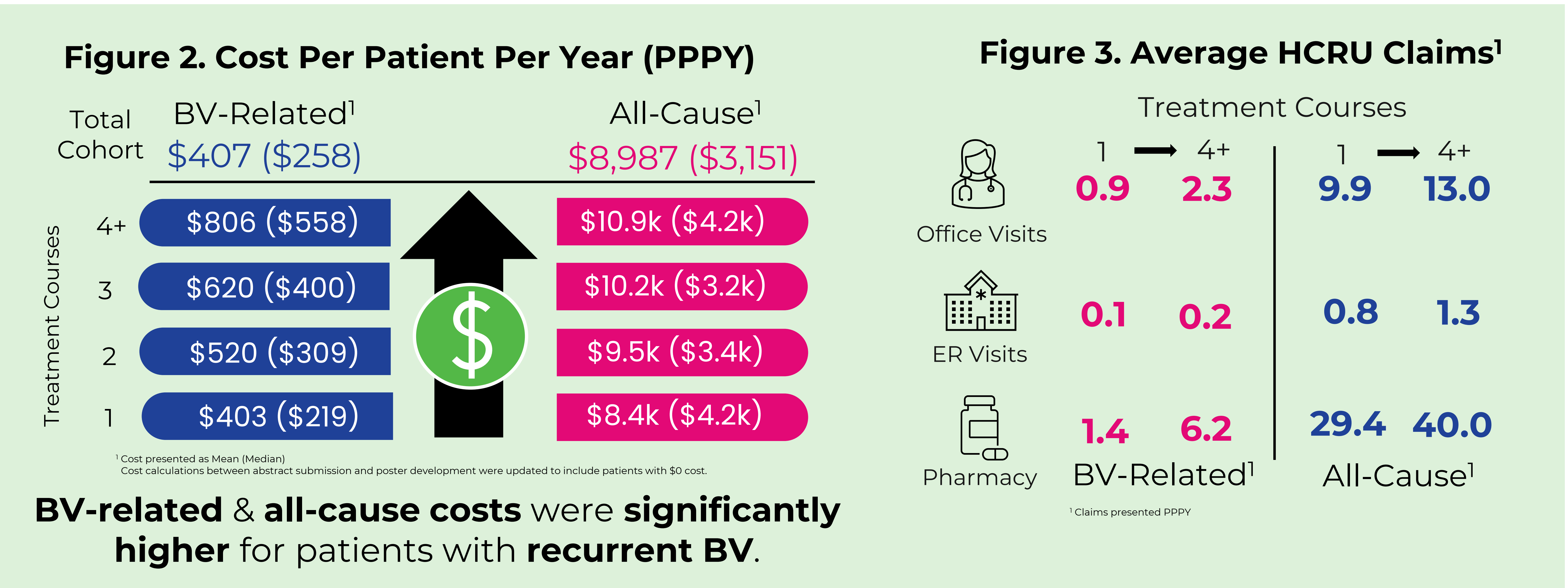
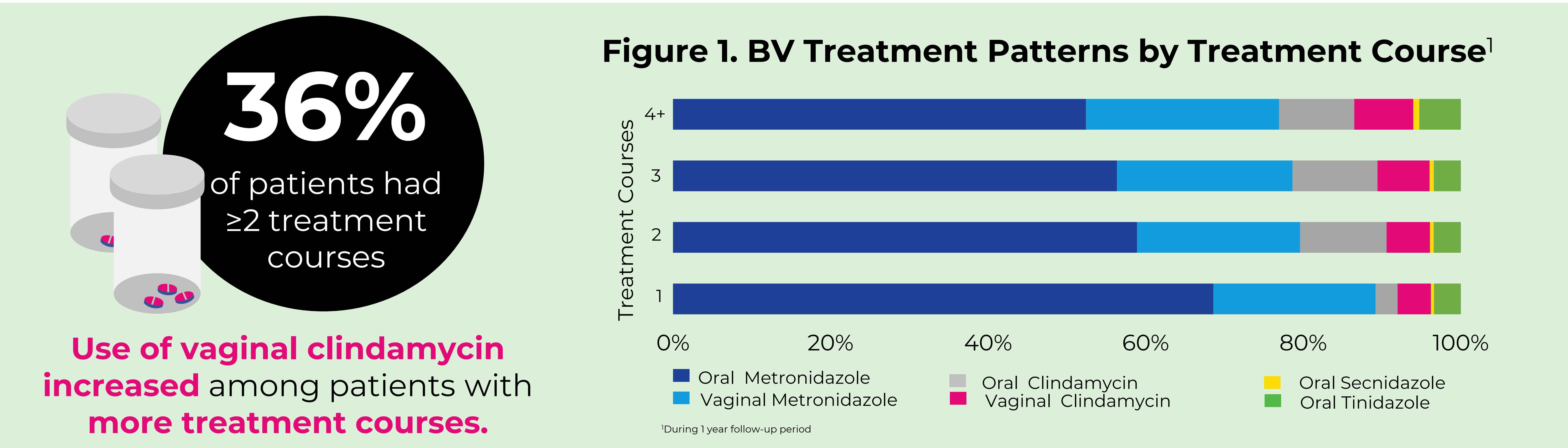
## Baseline Population Demographics

-  **140,826** with incident BV
-  **55%** US South Region
-  **13%** BV-associated complication
-  **11%** Charlson Comorbidity Index ≥1
-  **3%** Pregnant at baseline

# Treatment Patterns, Healthcare Resource Utilization (HCRU), and Direct Costs for Bacterial Vaginosis (BV) in US Commercially Insured Populations

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## Predictors of Cost & Recurrent BV<sup>+</sup>

- Having a comorbidity (CCI = ≥1), female infertility, and post procedural gynecological infection at baseline increased average costs from \$13k-14.5k U.S dollars.
- Top predictors of more treatment courses at baseline:
  - having any STI (Rate Ratio [95% CI]): 1.06 [1.05-1.08];
  - 30-39 years: 1.06 [1.03-1.08];
  - 18-29 years: 1.06 [1.04-1.08];
  - 40-49 years: 1.04 [1.01-1.06];
  - gynecological procedure infection: 1.05 [1.00-1.09]

## Key Assumptions & Limitations

- Patients with non-bacterial vaginitis who received BV medication may have been included.
- Data reflect treatment dispensed from pharmacy claims, and not real-world compliance which is variable. We did not assess use of over-the-counter BV drugs or behavioral factors.
- Findings are not generalizable to patients with non-commercial insurance.

## Conclusions

- Recurrent BV significantly (p < 0.01) increased costs at follow-up.
- Patients with more treatment courses used more HCRU for any cause.
- Baseline comorbidities significantly predicted (p < 0.01) increased cost and treatment courses.
- Providers should consider switching treatment earlier for patients over age 18 with BV-associated sequelae at baseline.

## Disclosure & References

This study was funded by Organon. Watkins, Yong, Tangirala and Collins and Li are employees of Organon. All authors contributed to and approved the poster.

- Koumans E, Sternberg, M, Bruce C, et al. Sex Transm Dis. 2007;34(11):864–869.
- Bradshaw CS, Sobel JD. J Infect Dis. 2016;214(Suppl 1):S14–S20.
- Workowski KA, Bolan GA; Centers for Disease Control and Prevention. MMWR Recomm Rep. 2015;64(RR-03):1-137.
- Bradshaw CS, Morton AN, Hocking J, et al. J Infect Dis. 2006;193:1478-1486.