

P103 Etiology of respiratory infections in non-hospitalised patients:Baseline etiological data from a randomised trial to investigate the effect of nasal sprays versus usual care in non-hospitalised patients with respiratory infections (ECRAID-Prime)

Katherine Loens¹, Lonneke Van Vught², Stefan Morreel³, Samuel Coenen³, Lisanne Vintcent², Ellen van Deuren², Frank Leus², Anouk Vanderstraeten¹, Herman Goossens¹, Chris Butler⁴, Alike van der Velden², Margareta Ieven¹ and Investigators on behalf of the ECRAID-Prime Consortium

¹Laboratory of Medical Microbiology, Vaccine & Infectious Disease Institute, University of Antwerp, Antwerp, Belgium; ²Julius Centre for Health Sciences and Primary Care, University Medical Center Utrecht, Utrecht University, Utrecht, the Netherlands; ³Department of General Practice, Vaccine & Infectious Disease Institute, University of Antwerp, Antwerp, Belgium; ⁴University of Oxford, Oxford, United Kingdom.

BACKGROUND & OBJECTIVES

COVID-19 and COVID-like-illness cause significant ill-health. Since the greatest burden is in the community, it is in primary care (PC) and community settings where early treatments will provide most benefit. ECRAID-Prime is a platform randomised, multi-country trial for the evaluation of investigational products for treatment of patients in PC with respiratory infections. The aim of the current study was to determine the baseline etiology in the recruited patient population.

METHODS

- Study design**
Platform randomised, trial comparing nasal sprays with usual care in PC patients with a respiratory tract infection (RTI) conducted in Belgium, France, Germany, Georgia, Ireland, Poland, United Kingdom (UK), and Spain.
- Participants**
795 patients (290 male and 505 female), aged 18-93 years, presenting to PC with RTI symptoms ≤7 days (**Fig.1**)
- Study duration**
Patients were enrolled during 2 consecutive winter periods (09/10/2024 and 20/03/2026)
- Procedure**
Nose/throat swabs taken at baseline (D0; N=795), D4, D7 and D14. Nucleic acids extraction using the MagMAX Viral/Pathogen II Nucleic Acid Kit on the KingFisher Flex. Analysis by the TrueMark Respiratory Panel 2.0, TaqMan Array Card (ThermoFisher Scientific) to detect presence of 21 respiratory viruses and 5 atypical bacteria.

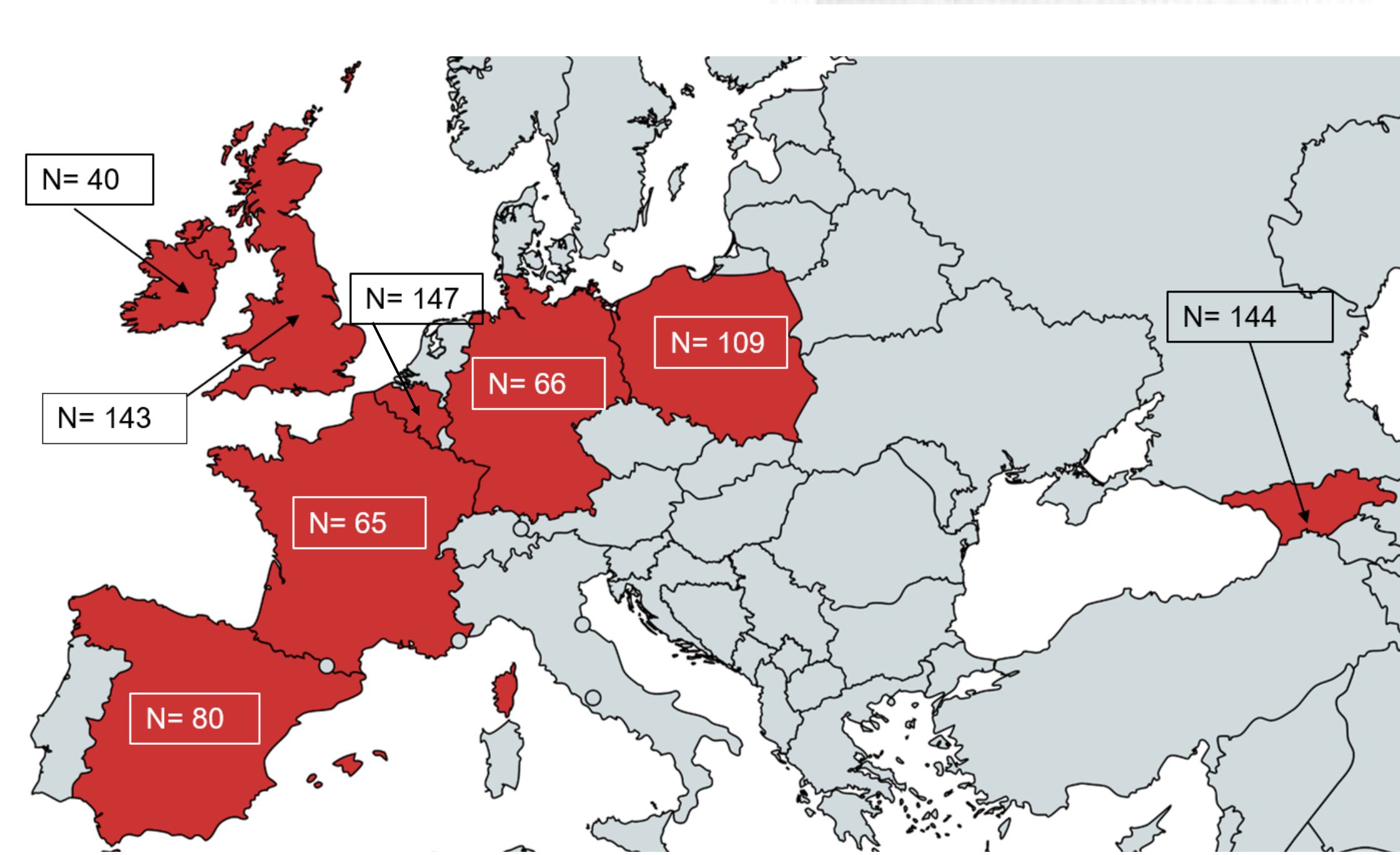
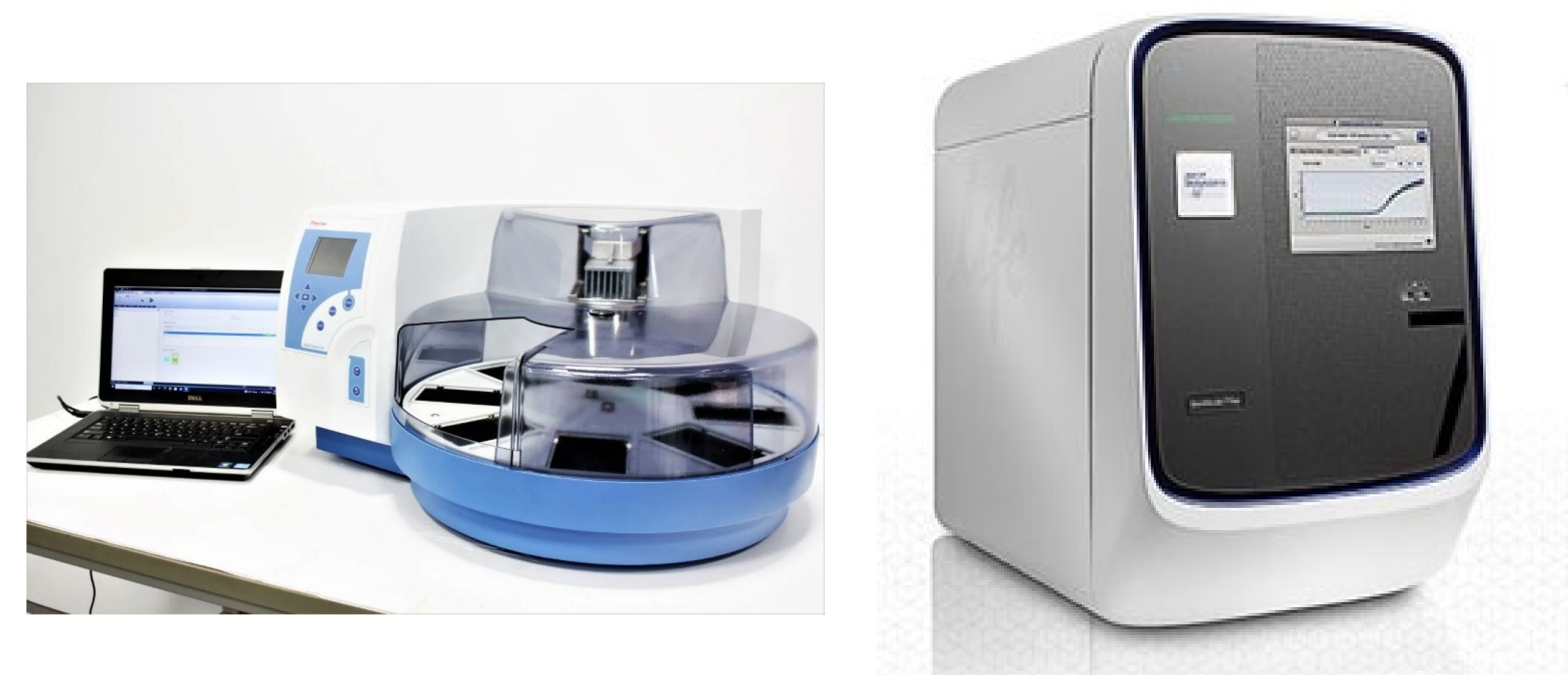


Figure 1: Number of participants per country



RESULTS

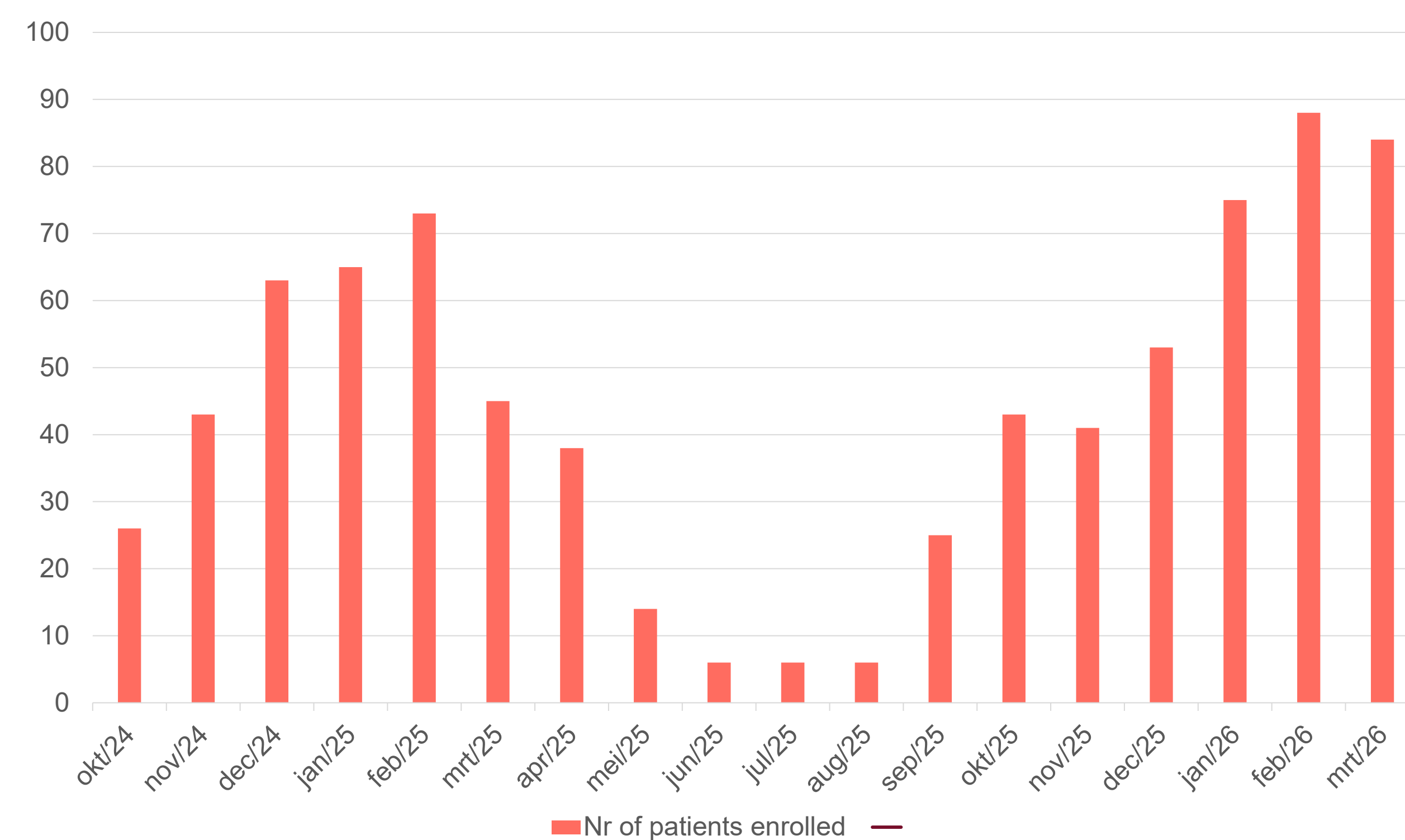


Figure 2: Per month participant enrollment

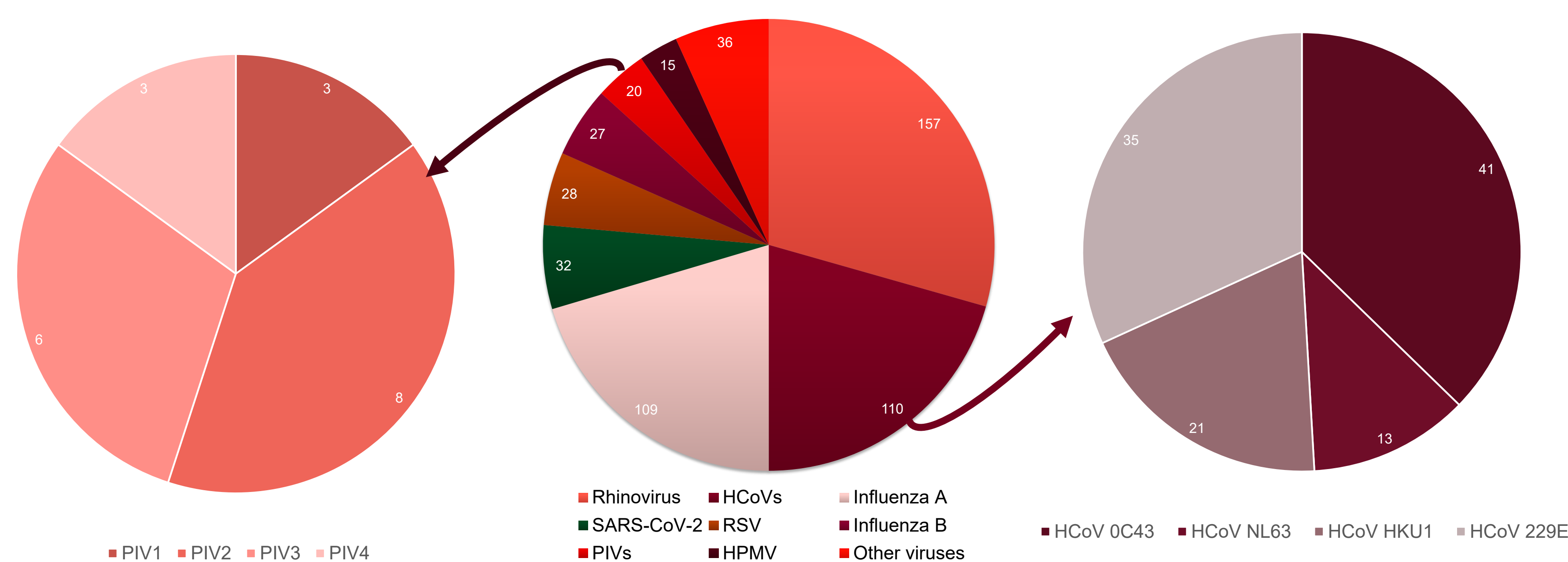


Figure 3: Cumulative positive results per virus October 2024 to March 2026

- Samples from 794/795 patients, enrolled from 10/2024 till 03/2026, in Belgium, France, Germany, Georgia, Ireland, Poland, UK and Spain (Fig. 1), respectively, were analysed (males n=209, females n=505, 18 - 93 years old).
- Respiratory viruses were detected in 489 (61.6%) of cases: human rhinovirus (157), influenza A/B (136), seasonal coronaviruses (110, predominantly HCoV OC43 at 37.3%), human parainfluenza viruses (HPIV, 20, predominantly HPIV-2 at 40.0%), severe acute respiratory syndrome-Coronavirus-2 (SARS-CoV-2; 32), RSV (28), human metapneumovirus (HMPV, 15), EV-D68 (17), HBoV (11) and HADE (8) (**Fig. 3**).
- M. pneumoniae* (n=3) and *C. pneumoniae* (n=1) infections were also detected.
- Viral coinfections occurred in 44 (5.5%) of positive samples, with influenza and rhinovirus (75.0%, 33/44) being most frequent involved (**Fig. 4**).

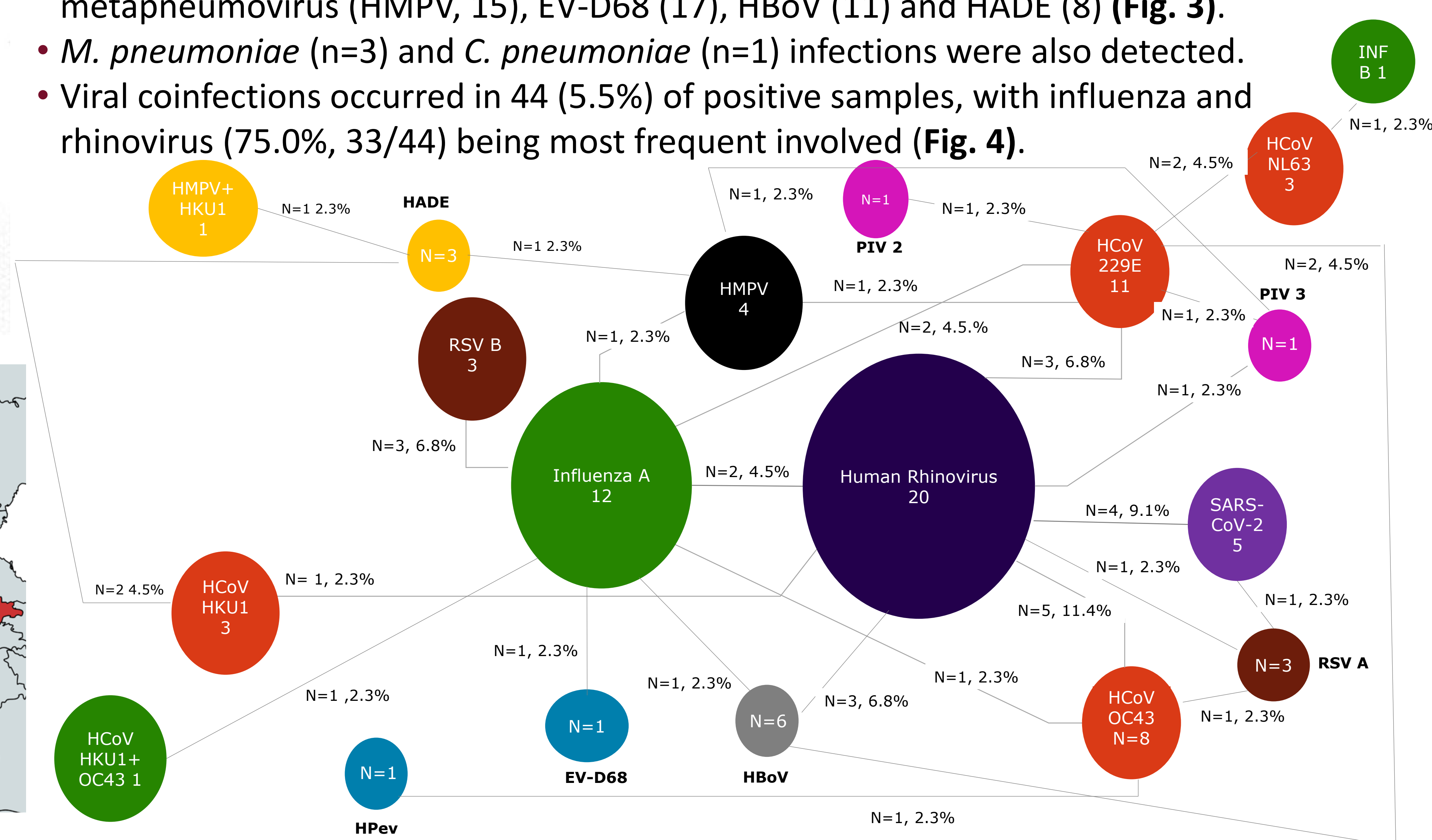


Figure 4: Viral coinfections in primary care patients with respiratory tract infections

CONCLUSION

In two out of three patients presenting with COVID-like-illness to primary care, a respiratory virus was detected. Influenza, Human Rhinovirus and seasonal coronaviruses were most often detected. In 5.5% multiple respiratory viruses were detected.

This study builds on European Clinical Research Alliance on Infectious Diseases – Primary care adaptive platform trial for pandemics and epidemics (ECRAID-Prime) which has received funding from the European Union's Horizon 2020 research and innovation programme under grant agreement No. 101046109.