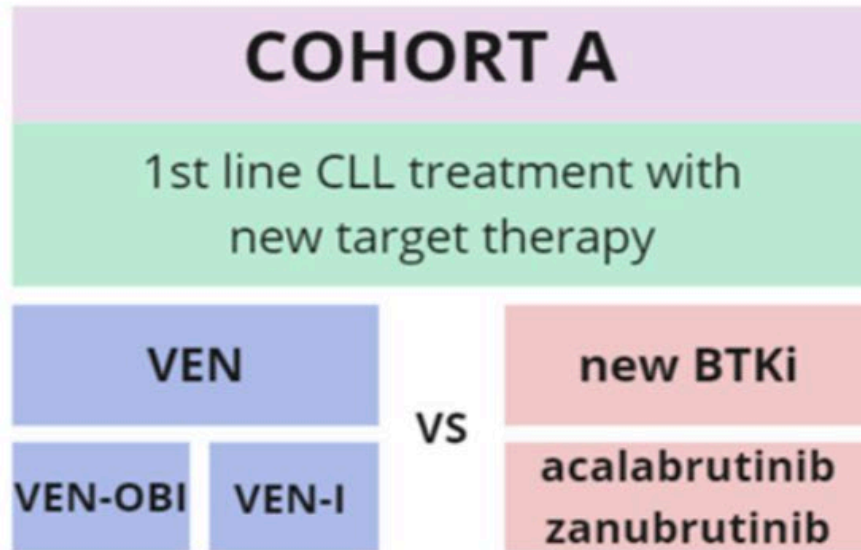


INFECTION LANDSCAPE IN PATIENTS WITH CHRONIC  
LYMPHOCYTIC LEUKEMIA TREATED WITH VENETOCLAX-BASED  
REGIMENS COMPARED TO THE SECOND-GENERATION BRUTON  
TYROSINE KINASE INHIBITORS:  
MULTICENTER, OBSERVATIONAL STUDY (**IMAGINE**) CONDUCTED  
BY ERIC, EUROPEAN RESEARCH INITIATIVE ON CLL

*Elzbieta Kalicinska, MD, PhD, Project Leader*

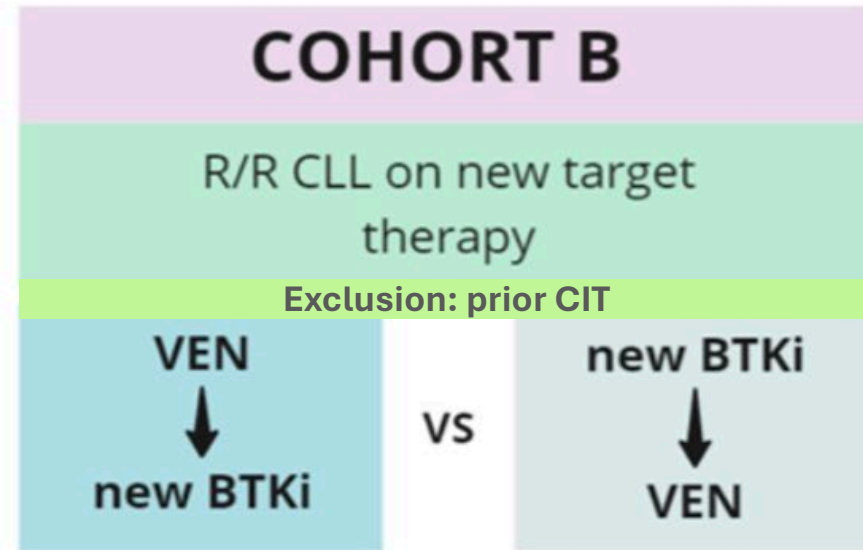
*Tomasz Wrobel, MD, PhD, Krzysztof Giannopoulos, MD, PhD,  
Marta Morawska, MD, PhD, Paula Jablonowska, MSc*

# STUDY DESIGN



24 months of active observation

Time points  
**early infections:** during the first 6 months of treatment  
**late infections:** from 6 to 24 months



24 months of active observation

Time points  
**early infections:** during the first 6 months of treatment  
**late infections:** from 6 to 24 months

# Objectives

## PRIMARY:

To compare the **cumulative incidence for severe infections (G3-G5)** in CLL patients treated with venetoclax-based regimens or second-generation BTKi in the first line and subsequent treatment lines (R/R setting) within the centers of ERIC.

## SECONDARY:

- to compare the cumulative incidence of all infections between patients treated with venetoclax-based vs new BTKi
- to evaluate **the event-free survival (EFS)**, (defined as the time from initiation of targeted therapy to the date of treatment discontinuation, first occurrence of disease progression, clinically significant infection (G3-4), or death from any cause, including death due to infections, whichever occurs first)
- to assess the impact of infections on **compliance, the timeliness of treatment administered**

# Objectives

## EXPLORATORY:

- to identify risk factors of infections to **develop the model of infectious risk score** in CLL patients treated **in the first line** with targeted therapy
- To assess whether and/or which sequence of treatment is associated with improved immune function and reduced incidence of infection in R/R CLL patients