

# 10 AL 13 DE NOVIEMBRE 2024 GLASGOW



# ANTIRETROVIRAL TREATMENT WITH BIC/FTC/TAF: WHERE WE COME FROM AND WHERE WE ARE GOING

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## INTRODUCTION:

Since its commercialization in Spain in 2019, bictegravir/emtricitabine/tenofovir alafenamide (BIC/FTC/TAF) has been one of the preferred antiretroviral treatments (ART) in people living with HIV (PLWH), both naïve and switch patients. Its advantages include being a single tablet regimen, having a high genetic barrier, and being active against hepatitis B virus (HBV). The objective of this study is to analyse the changes in treatment from or to BIC/FTC/TAF, as well as the reasons for these changes.

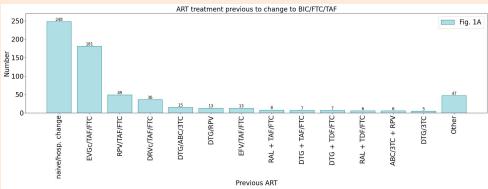
#### MATERIALS AND METHODS:

Data were analysed from all PLWH in University Hospital Son Llàtzer, in Mallorca, Spain, who at some point had been treated with BIC/FTC/TAF. We studied the previous ART as well as the subsequent treatment (if any), and the reasons for these changes.

#### RESULTS:

Since 2019, 641 PLWH have been treated with BIC/FTC/TAF (55% of all PLWH). There were 21.5% women, and 40% of patients with positive total core antibody for HBV (table 1).

The majority of patients who started BIC/FTC/TAF were naive patients or transferred from other countries without previous treatment in our hospital (248; 38.7%). There were 181 (28.2%) patients who came from regimens with another integrase inhibitor with cobicistat (EVGc/TAF/FTC) and 98 (15.3%) patients who did not have INSTI in their ART (RPV/TAF/FTC, DRVc/TAF/FTC or EFV/TAF/FTC). There were also changes from two-pill regimens (77, 12%), and some changes from dual therapies with DTG/3TC or DTG/RPV (18, 2.8%). Of the 641 PLWH treated with BIC/FTC/TAF, 575 (89.7%) maintain this treatment until now. Of the remaining 66 PLWH who switched to other treatments, most of them switched to intramuscular CAB/RPV (24; 36.4%), followed by oral dual therapies such as DTG/3TC (17; 25.8%) or DTG/RPV (4; 6%) (figure 1). The reasons for all changes are summarized in Table 2, with the discontinuation of a booster or the introduction of an integrase inhibitor being the main reasons for discontinuing BIC/FTC/TAF, and the change in route of administration or simplification being the main reasons for discontinuing BIC/FTC/TAF.



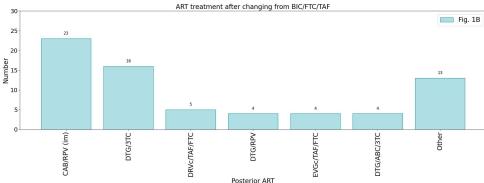


Figure 1. ART treatment before initiating BIC/FTC/TAF (Fig. 1A) and ART treatment after discontinuation of BIC/FTC/TAF (Fig. 1B)

# **CONCLUSION:**

Antiretroviral therapy with BIC/FTC/TAF is the most commonly use ART in our hospital, being a safe therapy that is generally maintained over time and suitable for PLWH co-infected with HBV. The main reason for discontinuation is the participant's desire to switch to intramuscular therapies.

**Table 1.** Characteristics of the people included in the study. Results are expressed as mean (SD) or N (%)

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Characteristic	Mean (SD) or N (%)	
Age, years	47 (SD: 12)	
Last CD4 count, cel/uL	713 (SD: 395)	
Gender, women	138 (21.5%)	
Nationality		
West Europe	452 (70.5%)	
Central/South America	127 (19.8%)	
Africa	44 (6.9%)	
Eastern Europe	12 (1.9%)	
Asia	5 (0.8%)	
North America	1 (0.1%)	
Transmission route		
Sexual intercourse-MSM	270 (42.1%)	
Sexual intercourse-MSW	221 (34.5%)	
Sharing injection material	112 (17.5%)	
Other/Unknown	38 (5.9%)	
Last HBcAb positive	249 (40.6%)	
Last HBsAg positive	23 (3.7%)	

**Table 2.** Reasons of initiation or discontinuation of BIC/FTC/TAF. Results are expressed as N (%)

2.5/1.5/1.11.11.55.11.5 4.5 6.6/1.55.55.4 45.11 (75)			
Reason of administration/change	Changes to BIC/FTC/TAF N = 641	Changes from BIC/FTC/TAF N = 66	
Naive/change of hospital	248 (38.7)	-	
Eliminate cobicistat	182 (28.4)	-	
Initiate INSTI	100 (15.6)	-	
Simplification	77 (12)	15 (22.7)	
Secondary effects	14 (2.2)	12 (18.2)	
Prevent interaction	8 (1.2)	2 (3)	
Virological failure/blips	5 (0.8)	2 (3)	
Adherence /loss to follow up	4 (0.6)	7 (10.6)	
Change administration route	-	24 (36.4)	
Other	3 (0.5)	4 (6.1)	

## References

Galan J, et al. Lancet 2017: 390: 2063-2072 Molina JM, et al. Lancet HIV 2018; 5:e357-e365

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