Results: We enrolled 147 patients, out of them 66 patients were on monotherapy with Abatacept due to intolerance or contraindications and 81 in therapy with Abatacept plus MTX. The two cohorts appeared homogeneous in age, gender, disease duration and baseline activity indexes, with the only difference being a higher baseline Physician Global assessment (PhGA) values in monotherapy patients. During the follow-up (median duration 24±14 months), the retention rate of Abatacept treatment was 71.2% in MTX patients (median duration 27–15.6 months) and 62.1% in monotherapy patients (median duration 25.2–17.5 months). No differences between the two groups in terms of retention rate, low-disease activity and CDAI remission (log rank p=ns), Breslow p=ns) were detected.

Conclusion: In patients with RA with intolerance or contraindication to MTX use, Abatacept monotherapy could be an efficient and safe option even in the long term follow-up.


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AB0316 MULTIPLE SWITCH BETWEEN BIOSIMILARS DMARDS (BSDMARDS) IN PATIENTS WITH INFLAMMATORY ARTHRITIDES: EXPERIENCE OF A SINGLE ITALIAN CENTRE

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Background: The availability of bsDMARDs since some years represents an opportunity to improve patient access to effective biological therapy, to better accommodate restraint within healthcare budgets and to improve overall patient outcomes. Different policies are followed in different countries to implement the use of bsDMARDs. Although the latest position paper of AIFA (Agenzia Italiana del Farmaco) envisions the automatic substitution between the originator and biosimilar, until now the prescriber decision and the patient consent are strongly advised. The question around biosimilar to biosimilar switching is overlooked. Nevertheless different rules are established at regional level and in our Hospital automatic switching between originator/biosimilar and biosimilar/biosimilar was applied.

Objectives: To analyze the efficacy and safety of switch from originator to biosimilar (O/B) and/or biosimilar to biosimilar (B/B) in patients with RA, PsA and SPA.

Methods: We retrospectively analyzed in 63 patients (30 F, mean age 58.3 yr, 21 RA, 26 PsA, 16 SPA), treated with Infliximab, Etanercept and Adalimumab, disease activity (DAS28 CRP for RA, Tender/Swollen joint count for PsA, BASDAI for SPA, CRP for all) and adverse events/infections (AE). The time points considered were 3rd month before the switch and 3rd and 6th month after.

Results: 45 patients underwent single switch (35 O/B, 9 B/B) and 18 (28.5%) double switch (O/B/B). 27 B/B switch were done. No differences in disease activity were observed before and after switch (8 RA patients: mDAS28 CRP 2.86±3.23, 11 PsA patients: mTJ count 2.5 > 3.43, 8 SPA patients: mBASDAI 2.88 > 2.84). The mean number of swollen joints was very low in PsA group and we decided to exclude this variable. The CRP level was low and stable throughout the follow-up in all the groups.

Conclusion: The time points considered were 3rd month before the switch and 3rd and 6th month after.

Disclosure of Interests: None declared

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AB0317 BIOLOGIC THERAPY OPTIMIZATION IN RHEUMATOID ARTHRITIS PATIENTS IN COLOMBIA

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Background: The optimization of biological agents (bDMARD), is a strategy that has proven to be cost effective and its use can reduce the risk related to drug exposure (1–3). It is included in the EULAR management guidelines and in the consensus of the Colombian Rheumatology Association.

Objectives: To analyze optimization success of bDMARD therapies in patients with RA.

Methods: Cohort study of RA patients in a specialized multicenter institution in Colombia, followed from January 2015 to December 2019. Patients in remission or low activity for at least 6 months with bDMARD, and with at least two consecutive medical visits, were included. Optimization types were dose decrease, application interval increments, or both. Patients who had disease reactivation (DAS28-CRP >3.2) and returned to standard dose, were considered a failure. By Kaplan-Meier analysis, the optimization failure was estimated according to bDMARD type.

Results: 92 patients were included, 78.26% were women, with a median age of 57 years (IQR 50-64), a disease evolution time of 15 years (IQR 10-21), a treatment of 5.6 years (IQR 2.7-15.0), and optimization of 7.75 months (IQR 3.2-15.75). The most commonly used bDMARD therapies were etanercept 36.96%, tocilizumab 30.43% and adalimumab 16.30%. 69.39% (34) were naive for bDMARD for 12 months or more, of these, 9.43% (27) continue in sustained remission, and 55.56% (15) received combined therapy with s synthetic DMARD (sDMARD).

95.92% remained under optimization scheme without disease activity changes, and 4.08% of patients underwent definitive discontinuation of bDMARD, for sustained therapeutic objective. 8.16% (4) had relapses in the first 6 months after onset, of which 2 patients returned to standard doses. In survival analysis it was observed that patients who were optimized for anti-TNF failed faster than the non-antiTNF, although this difference was not statistically significant (Log Rank test 0.003 p value = 0.959). Of the total patients, 28 have been optimized for 12 months or more, of these, 96.43% (27) continue in sustained remission, and 55.56% (15) received combined therapy with s synthetic DMARD (sDMARD).

Conclusion: During follow-up, most patients remain in optimization strategy. In those who continued in sustained remission, more than half received sDMARD, this suggests that their use may be a determining factor in preventing disease relapses. More studies are required to evaluate this hypothesis.

Disclosure of Interests: None declared


Figure 1. Kaplan Meier

Kaplan-Meier survival estimates

0.00 0.25 0.50 0.75 1.00

0 10 20 30 40 50 60 70 80 90 100

time in years

Anti TNF No Anti TNF

Figure 1. Kaplan Meier

Conclusion: During follow-up, most patients remain in optimization strategy.

Disclosure of Interests: None declared

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