Current and Emerging DMARDs for the Treatment of Rheumatoid Arthritis

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Abstract: Rheumatoid arthritis (RA) is the most prevalent form of inflammatory arthritis. It is a profoundly serious and severe disease that if it goes untreated could have severe consequences to the joints and health of the patient who carries this diagnosis. The treatment of RA has dramatically changed since the year 2000, with the discovery of the TNFis, then other biologics, and finally the JAKi. All these new medications with or without methotrexate in combination, tight control and treat to target have produced a revolution in the outcome of this disease. We reviewed and summarized the treatment options, and the most significant papers for each one of these new drugs. The reader could have a full picture with all the references of the recent publications. We also updated the biosimilar situation in RA, as well as the new drugs that will be coming to the market in the next 5 years.

Keywords: rheumatoid arthritis, DMARDs, biosimilars

Introduction

Rheumatoid arthritis (RA) is one of the most prevalent chronic inflammatory joint diseases that can lead to cartilage and bone damage as well as disability. In the last two decades, a therapeutic revolution in the treatment of RA has begun that includes treat to target strategy, tight control and new biologic and non-biologic antirheumatic drugs. With the aim to prevent structural joint damage, loss of function and maintain quality of life, patients are treated early and more aggressively.

RA is a heterogenic disease that requires the use of multiple therapies with different mode of action to achieve remission or at least low disease activity, as recommended by the EULAR guidelines and treat to target. No clear biomarker has been described that allows us to decide which drug is better for each individual patient. Two types of advance therapies are available (Table 1): bDMARDs (biological disease modifying antirheumatic drugs) which are most frequently monoclonal antibodies or receptor constructs that target a specific soluble or cell surface molecule, either a cytokine, a cytokine receptor or another cell membrane antigen. They either prevent interaction of the specific ligand with its receptor, destroy a specific cell population, or inhibit interaction between particular cell populations. They must be administered IV or SC since they are proteins. They also do not enter the cell but mediate their respective modes of action outside the cell or via the cell surface.

Target synthetic DMARDs, like the JAKi (Janus Kinase inhibitors), represent a series of intracellularly active drugs. The pathways that mediate cytokine receptor...
**Table 1** Approved Drugs for RA Treatment

| Drug         | Structure                                      | Mode of Action                          | Pivotal Studies                                      |
|--------------|------------------------------------------------|-----------------------------------------|------------------------------------------------------|
| Infliximab   | Chimeric monoclonal antibody                   | Anti TNF inhibitor                      | ATTRACT Maini 1999                                  |
|              |                                                |                                         | ASPIRE St Clair 2004                                 |
|              |                                                |                                         | BEST Yvonne 2007                                    |
| Etanercept   | Fusion protein                                 | Anti TNF inhibitor                      | ERA Barthol 2000                                     |
|              |                                                |                                         | COMET Emery 2008                                     |
|              |                                                |                                         | TEMPO Klareskog 2004                                 |
| Adalimumab   | Human monoclonal antibody                      | Anti TNF inhibitor                      | ARMADA Weinblatt 2003                                |
|              |                                                |                                         | PREMIER Breedvelt 2006                               |
|              |                                                |                                         | OPTIMA Kavanaugh 2013                                |
| Certolizumab | Pegylated human monoclonal antibody            | Anti TNF inhibitor                      | RAPID 1 Keystone 2008                                |
|              |                                                |                                         | RAPID 2 Smolen 2009                                  |
|              |                                                |                                         | FAST4WARD Fleischmann 2009                           |
| Golimumab    | Human monoclonal antibody                      | Anti TNF inhibitor                      | GO BEFORE Emery 2009                                 |
|              |                                                |                                         | GO FORWARD Keystone 2009                             |
|              |                                                |                                         | GO AFTER Smolen 2009                                 |
| Abatacept    | Human fusion protein                           | T Cell co stimulatory inhibition (CD80/CD86) | ABA in MTX RESISTENT PATIENTS Kremer 2006          |
|              |                                                |                                         | ABA in TNF REFRACTORY PATIENTS Genovese 2005        |
|              |                                                |                                         | ACQUIRE Genovese 2011                                |
| Tocilizumab  | Human monoclonal antibody                      | AntiIL 6 inhibitor receptor             | TOWARD Genovese 2008                                 |
|              |                                                |                                         | RADIATE Emery 2008                                   |
|              |                                                |                                         | AMBITION Jones 2010                                  |
|              |                                                |                                         | ACT-RAY Dougdados 2013                               |
| Sarilumab    | Human monoclonal antibody                      | AntiIL 6 inhibitor receptor             | SARIL-RA-MOBILITY Huizinga 2014                      |
|              |                                                |                                         | TARGET Fleischmann 2017                              |
| Rituximab    | Chimeric monoclonal antibody                   | B cell depletion (anti CD20)            | DANCER Emery 2006                                    |
|              |                                                |                                         | REFLEX Cohen 2006                                    |
| Tofacitinib  | Small molecule                                 | JAK1 and JAK3 inhibitor                 | ORAL Start                                          |
|              |                                                |                                         | ORAL Sync                                           |
|              |                                                |                                         | ORAL Scan                                           |
|              |                                                |                                         | ORAL Solo                                           |
|              |                                                |                                         | ORAL Standard                                       |
|              |                                                |                                         | ORAL Step                                           |
|              |                                                |                                         | Oral Strategy                                       |
| Baricitinib  | Small molecule                                 | JAK 1 and JAK 2 inhibitor               | RA Begin                                            |
|              |                                                |                                         | RA Beacon                                           |
|              |                                                |                                         | RA Bean                                             |
|              |                                                |                                         | RA Build                                            |
|              |                                                |                                         | RA Beyond                                           |
| Upadacitinib | Small molecule                                 | JAK1 inhibitor                           | SELECT Netx                                         |
|              |                                                |                                         | SELECT Beyond                                       |
|              |                                                |                                         | SELECT Monotherapy                                   |
|              |                                                |                                         | SELECT Early                                        |
|              |                                                |                                         | SELECT Compare                                      |
| Filgotinib   | Small molecule                                 | JAK1 inhibitor                           | FINCH 1                                             |
|              |                                                |                                         | FINCH 2                                             |
|              |                                                |                                         | FINCH 3                                             |
|              |                                                |                                         | FINCH 4                                             |

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signal transduction JAKs are non-receptor tyrosine kinases associated with the cytoplasmic domain of type I and II cytokine receptors which are activated when these are engaged by their ligands; once phosphorylated, they phosphorylate signal transducers and activators of transcription (STATs) which then induce gene activation. They are oral small molecules that act intracellularly, in a reversible way, preventing the phosphorylation of JAKs. Many cytokines, such as interleukin (IL)-2, 6, 12, 15 and 23 as well as interferons use the JAK-STAT pathways, while others, such as IL-1, IL-17 and TNF, do not use JAK enzymes. JAK1, 2, 3 and TYK2 – function as dimers and once activated phosphorylate STATs, which subsequently induce gene transcription.

Actually, there are five molecular target families available as treatment options for RA: tumor necrosis factor (TNF) inhibitors, interleukin 6 (IL-6) receptor blockers, CD80/86 inhibitors, anti-CD20 and Janus kinases (JAK) inhibitors with multiple drugs for several of these mechanisms. Biosimilars (to some of these drugs) have been developed, becoming part of the rheumatologic armamentarium that needs to be considered, as they will increase access.

The goal of this review is to describe the current therapies, including recently approved anti-rheumatic agents, and to mention the ones that are in development (mainly in Phase 3) focusing on efficacy and emerging safety issues. Randomized controlled clinical trials that have been done to prove the efficacy and safety of drugs for the treatment of rheumatoid arthritis were selected from different databases (including PubMed, EULAR and ACR congresses). A specific search with words including clinical trials, head-to-head clinical trials, biologic DMARDs and synthetic DMARDs in rheumatoid arthritis. We also asked colleagues for specific trials that could have been important for the paper.

The Anchor Drug: Methotrexate
Methotrexate (MTX) remains the first choice in the treatment of rheumatoid arthritis, because it is effective in 25% of patients (remission), has an acceptable toxicity profile and low costs. In the management of early and established RA, MTX is recommended as a first-line drug by the European League Against Rheumatism (EULAR) and the American College of Rheumatology (ACR). It has shown to be effective in monotherapy and also is the basis for combination therapies. In early RA patients, starting MTX in monotherapy followed by the addition of anti-TNF on MTX treatment failure at 6 months had similar outcomes (clinical and functional) compared with patients who started on a combination therapy. It is important to remember that a group of patients will achieve remission only with methotrexate monotherapy, so avoiding over-treatment should always be considered.

Anti TNF Alpha Agents
Tumor necrosis factor (TNF) is a pro-inflammatory cytokine that plays an important role in joint inflammation and contributes to joint destruction. The inhibition of TNF improves the clinical manifestations of RA and reduced radiographic progression. There are 5 biologic agents targeting the TNF approved for the treatment of RA: infliximab (INF), etanercept (ETN), adalimumab (ADA), certolizumab (CMZ) and golimumab (GLM). Several clinical trials of these compounds showed excellent efficacy on RA and an acceptable risk profile.

Infliximab was the first TNF alpha inhibitor (TNFi) developed. It is a chimeric monoclonal antibody and requires intravenous application every 4–8 weeks. Several controlled trials demonstrated the efficacy of INF for early and established RA. Patients receiving the combination treatment (infliximab + MTX) showed lower radiographic progression, higher remission rates and improved efficacy compared to patients receiving MTX alone.

Etanercept is a fusion protein of the soluble TNF receptor and Fe portion of immunoglobulin, has the shortest half-life of available TNFi and is administered subcutaneously. Several trials demonstrated the efficacy of ETN in early and established RA. The combination therapy with MTX demonstrated higher clinical response rates and less radiographic progression than ETN or MTX monotherapy. These improvements were sustained during open-label extension trials.

Adalimumab is a fully human monoclonal antibody binding TNF. The clinical efficacy of this compound in combination with methotrexate was shown to be superior than MTX alone in patients with early and stabilish RA.

Certolizumab pegol is a pegylated, humanized anti-TNF Fab fragment. Its structure makes it different from other TNFi. CMZ demonstrated similar efficacy in achieving ACR disease activity measures as the other TNF inhibitors and similarly inhibits radiographic progression. Because it has minimal to no active placental transfer, analysis of pregnancy outcomes seems favorable to this drug regarding teratogenic effect and risk of fetal death.

Golimumab is a fully humanized monoclonal antibody that has demonstrated efficacy and safety in MTX naïve MTX inadequate response and in anti-TNF failure patients.

There is only one head to head trial that compares the efficacy and safety of two different TNF inhibitors. The
### Table 2: Head-to-Head Trials with Biologic and Synthetic DMARDs in Rheumatoid Arthritis

| Study            | Design            | Drugs                  | Follow Up | Number of Patients | Primary End Point                                      | Results                                                                 |
|------------------|-------------------|------------------------|-----------|--------------------|--------------------------------------------------------|-------------------------------------------------------------------------|
| EXCELEATE        | RCT superiority   | CMZ + MTX vs ETN + MTX | 104 weeks | 915                | ACR 20 week 12                                        | CMZ + MTX is not superior to ETN + MTX                                   |
| AMPL3E           | RCT non inferiority| SC ABA + MTX vs ADA + MTX | 2 years  | 645                | ACR 20 at 1 year                                      | SC ABA + MTX is not inferior to ADA + MTX                                |
| ADACTA           | RCT superiority   | TCZ IV vs ADA          | 24 weeks  | 326                | Change DAS 28 at 24 weeks                             | TCZ monotherapy is superior to ADA monotherapy                           |
| MONARCH          | RCT superiority   | SARI vs ADA            | 24 weeks  | 369                | Change DAS 28 ERS at 24 weeks                         | SARI monotherapy is superior to ADA monotherapy                           |
| ENTRACTE         | RCT non inferiority| TCZ IV vs ETN          | 3.2 years | 3080               | Time of occurrence of first MACE                      | TCZ was not inferior to ETN                                              |
| ORAL STRATEGY    | RCT non inferiority| TOFA monotherapy       | 1 year    | 1146               | ACR 50 at 6 months                                   | TOFA + MTX was non inferior to ADA + MTX                                  |
| RA BEAN          | RCT non inferiority| PBO + MTX              | 52 weeks  | 1307               | ACR 20 at 24 weeks                                   | BARI + MTX non inferior to ADA + MTX                                     |
| SELECT COMPARE   | RCT superiority   | PBO + MTX              | 48 weeks  | 1629               | ACR 20 and DAS28 CPR < 2.6 at week 12                 | UPA + MTX superior to ADA + MTX in terms of ACR 50, DAS28 CPRs 3.2       |
| SELECT CHOICE    | RCT superiority   | UPA + csDMARDs         | 24 weeks  | 612                | Change DAS 28 CPR at week 12                         | UPA was superior to ABA                                                 |
| FINCH           | RCT non inferiority| FILGO 100 + MTX        | 52 weeks  | 1755               | ACR 20 week 12                                       | FILGO 200 non inferior to ADA based on DAS28 CPRs 3.2                   |
| NCT02092467      | RCT non inferiority| TOFA 5 mg bid          | 5 years   | 4369               | MALIGNANCY MACE                                      | Pending results                                                         |
| RA BRANCH        | RCT non inferiority| BARI 2 mg              | 5.5 years | 1300               | VTE                                                   | Pending results                                                         |

**Abbreviations:** RCT, randomised controlled trial; CMZ, certolizumab pegol; ETN, etanercept; MTX, methotrexate; SC, subcutaneous; IV, intravenous; ABA, abatacept; ADA, adalimumab; TCZ, tocilizumab; SARI, sarilumab; TOFA, tofacitinib; BARI, baricitinib; UPA, upadacitinib; csDMARDs, conventional synthetic disease modifying antirheumatic drugs; FILGO, filgotinib; ACR, American College Rheumatology response; DAS 28 CPR, disease activity score 28 c reactive protein; LDA, low disease activity; MACE, major adverse cardiovascular events; VTE, venous thromboembolism event.
EXXELERATE study (Table 2), a superiority study, showed that certolizumab pegol plus methotrexate is not superior to adalimumab plus methotrexate (ACR20 response at week 12: 69% CZM vs 71% ADA; odds ratio 0.90 [95% CI 0.67–1.20]; p=0.467). Comparisons based on indirect and retrospective data analyses have proven that the efficacy of anti-TNF seems broadly similar between the five drugs. However, some patients’ characteristics could suggest that one TNF inhibitor is more favorable over the other. Data regarding safety also seem comparable between them. A meta-analysis of randomized clinical control trials of RA patients treated with anti-TNF demonstrated higher risk of serious infection (OR, 1.42; 95% confidence interval [CI], 1.13–1.78) and treatment discontinuation due to adverse events (OR, 1.23; 95% CI, 1.06–1.43) compared with placebo and traditional disease-modifying antirheumatic drug treatments. There is also a higher risk of reactivation of latent tuberculosis (TB) and other opportunistic infections. Monoclonal antibodies seem to have a higher risk of TB infection than etanercept. There have been reports of new episodes and exacerbations of central nervous system demyelinating disorders and lupus-like syndrome during anti-TNF therapy.

**T Cell Costimulatory Blocking (CD80/86)**

Abatacept is a fully human fusion protein that inhibits the second signal required for T-cell activation (by binding to CD80 and CD86 costimulatory antigens). It was first developed in EV formulation and later subcutaneously. Both treatment options showed comparable efficacy and safety in patients with RA. Abatacept reduces disease activity in MTX inadequate response patients. At 1 year, abatacept responses compared with placebo for ACR 20, 50 and 70 were 73.1% aba vs 39.7% pbo, 48.3% aba vs 18.2% pbo and 28.8% aba vs 6.1% pbo, respectively. Abatacept also produced significant clinical and functional benefits in patients who failed anti-TNF treatment. Several studies demonstrated the reduction of radiographic joint damage in patients treated with abatacept. The ATTEST study evaluated abatacept and infliximab vs placebo in RA patients with inadequate response to MTX. After one year, adverse events, serious infections and discontinuations due to AE were lower with abatacept than infliximab, showing a more acceptable safety and tolerability profile for this drug. The AMPLE trial (head to head comparing abatacept and adalimumab both combined with MTX) showed similar efficacy based on clinical functional and radiographic outcome. Even though the frequency of AE was similar in both groups, there were less discontinuations due to AEs and serious infections and fewer local injection site reactions with abatacept, favoring this drug (Table 2).

Like other biologics, serious infections were reported in patients with abatacept, but the frequency was low overall (3.0% vs 1.9% in abatacept- versus placebo-treated patients, respectively). There have been reports of acute infusion adverse effects (9.8% vs 6.7% in the abatacept versus placebo groups, respectively) but were mostly mild-to-moderate in intensity. There is a label warning of abatacept and patients with chronic obstructive pulmonary disease because more frequent respiratory adverse events were reported in this population.

**Anti-Interleukin 6**

There are two interleukin 6 receptor antagonist (anti-IL6) drugs approved for the treatment of RA. Tocilizumab (TCZ) is a fully humanized monoclonal antibody directed against the IL-6 receptor that can be administrated intravenous or subcutaneous, and sarilumab, a human monoclonal antibody directed against the alpha subunit of the IL-6 receptor complex. Both drugs demonstrated to be effective for patients with inadequate response to csDMARDs (conventional synthetic disease modifying antirheumatic drugs) and to TNF inhibitors. The ACT-RAY study showed that there was no relevant superiority of TCZ + MTX compared to TCZ monotherapy regarding clinical and radiographic responses in MTX-IR patients. As a consequence, anti-IL6 have been recommended by EULAR guidelines in patients on monotherapy. Both drugs have been compared with an anti-TNF treatment in monotherapy. The MONARCH study, a double-blind head-to-head superiority trial, compared Sarilumab with adalimumab. Sarilumab was superior to adalimumab in terms of the change from baseline in DAS28-ESR (−3.28 vs −2.20; p<0.0001). The ADACTA trial showed the superiority of tocilizumab compared to adalimumab (DAS28 TCZ group (−3.3) vs ADA group (−1.8), difference −1.5, 95% CI −1.8 to −1.1; p=0.0001). Some side-effects of anti-IL6 inhibitors are more prevalent from the other available biologics like neutropenia, elevation of liver function tests (hepatic transaminases and bilirubin) and elevations of total cholesterol, triglycerides, and high-density lipoprotein levels. GI perforations have also been reported more frequently with IL6 inhibitors in the Rabbit and other registries. In the MONARCH and ADACTA trials, the incidence of infections was similar between anti IL6 inhibitors and ADA. In a head-to-head non-inferior trial of Etanercept vs Tocilizumab, there was no difference in the rate of cardiovascular events independently of the higher rate of cholesterol elevation with Tocilizumab (Table 2).
Rituximab (CD20)

Rituximab is chimeric murine-human monoclonal antibody directed against CD20 that produces depletion of B cells. Its efficacy has been shown in patients who failed to respond to DMARDs. In the DANCER study significantly more patients who received rituximab in two 500 mg or two 1000 mg infusions with MTX achieved ACR 20 response rates at week 24 (55% and 54%, respectively), compared with placebo (28%, P < 0.001). Also, in patients with an inadequate response to anti-TNF therapies, a single course of rituximab (two 1000 mg infusions 15 days apart) with concomitant MTX demonstrated significant improvements in disease activity. It seems that seropositive rheumatoid factors patients respond better to rituximab than seronegative patients. The most frequent adverse event with Rituximab is infusion reaction. Even though this drug produces a prolonged B cell depletion, the risk for serious infection was similar between the placebo and RXT groups. Fulminant reactivation of hepatitis B has been reported after rituximab treatment, use of this drug in patients with hepatitis B positive serology is contraindicated. Progressive multifocal leukoencephalopathy, a very rare, but often fatal complication, has been described rarely in RA patients treated with RTX.

Small Molecules: Janus Kinase Inhibitors

The Janus Kinase inhibitors (JAKi) are the newest class of drug license for the treatment of RA. There are four different types of JAKs proteins: JAK1, JAK2, JAK3, and Tyk2 (tyrosine kinase) and so far, four different JAKs inhibitors are approved for the treatment of RA: Tofacitinib, Baricitinib, Upadacitinib and Filgotinib (penfiticitinib, a 5th one, is only approved in Japan and South Korea). All seem to share a similar efficacy and safety profile for patients with RA.

Tofacitinib

Tofacitinib is nonselective JAKi and was the first approved for the treatment of RA. Inhibits JAK1, JAK2, JAK3 and, to a lesser extent, TYK2. The Phase III tofacitinib trial program included 7 randomized controlled studies that demonstrated the efficacy, safety and prevention of structural damage in different populations of early and stablish RA patients.

The ORAL START was a MTX naïve trial that compared MTX to tofacitinib. The primary end point ACR 70 response rates were 25.5% in the 5 mg twice a day (bid), 37.7% in the 10 mg bid and 12% in the MTX (P<0.0001 for both comparisons). Tofacitinib had significantly less radiographic progression than MTX (Sharp score 0.2 point in 5 mg bid group, <0.1 point in 10 mg bid group and 0.8 point in MTX group P < 0.0001 for both comparisons).

The ORAL Sync study evaluated the efficacy and safety of Tofacitinib in combination with csDMARDs. ACR 20 response rates at 6 months were 21.2% for tofacitinib 5 mg bid compared to the combined placebo group (P < 0.01).

The ORAL Scan trial compared in MTX-IR patients tofacitinib 5 and bid 10 mg bid with placebo. The inhibition of structural damage by changes in total modified Sharp/van der Heijde from baseline was only statistically significant for the 10 mg bid group compared to placebo (0.06 vs 0.47 P< 0.05, respectively).

The ORAL SOLO trial assessed the efficacy and safety of tofacitinib monotherapy in MTX-IR patients. Tofacitinib groups showed significantly higher percentage of ACR 20 response rates compared to placebo (59.8% in the 5 mg bid tofacitinib group and 65.7% in the 10 mg bid tofacitinib group vs 26.7% in the combined placebo groups, P<0.001 for both comparisons).

The ORAL STANDARD trial compared the efficacy and safety of tofacitinib with placebo and with Adalimumab (as a control) in MTX-IR patients. The study demonstrated that tofacitinib was significantly superior to placebo and was numerically similar to adalimumab in efficacy (ACR 20) response rates for tofacitinib 5 mg bid 51.5%, 10 mg bid 52.6%, adalimumab 47.2% and placebo 28.3% (P<0.001 for all comparisons). This study was not a head-to-head trial to demonstrate superiority or non-inferiority between tofacitinib and adalimumab.

ORAL STEP compared the efficacy of tofacitinib with methotrexate in patients with an inadequate response to at least one prior TNF inhibitor. The primary end points demonstrated that tofacitinib plus methotrexate was superior than placebo in this treatment refractory population (ACR 20 41.7% tofacitinib 5 mg bid, 48.1% tofacitinib 10 mg bid vs 24.4% placebo, p=0.0024 and p<0.0001, respectively). Also, there were statistically significant improvements in ACR 50 and 70 response rates, and PROs for tofacitinib groups.

ORAL Strategy, a phase 3b/4 head to head, non-inferiority trial assessed the efficacy of tofacitinib monotherapy, tofacitinib plus methotrexate, and adalimumab plus methotrexate in MTX-IR patients. ACR 50% response rate (primary end point) was 38% for tofacitinib monotherapy, 46% for tofacitinib plus MTX and 44% for adalimumab plus MTX, demonstrating the non-inferiority of tofacitinib plus MTX vs Ada plus MTX. However, tofacitinib monotherapy was inconclusive to either combination (Table 2).
Tofacitinib is approved in a dose regime of 5 mg bid and in 2016, an extended-release formulation of 11 mg daily was approved later on.

**Baricitinib**

Baricitinib inhibits JAK1 and JAK2 and to a much lesser extent TYK2. Four randomized double-blind placebo-controlled phase III clinical trials assessed the efficacy of this drug as mono or in combination therapy in patients with RA. There is also a long extension study trial, for the patients that completed these pivotal trials. The five trials achieved the primary endpoint (ACR 20 improvement criteria) at week 12 and 24, and also major secondary endpoints (ACR 50 and 70, DAS28 response and patients reported outcomes) were accomplished by baricitinib vs placebo. RA BEGING evaluates baricitinib as monotherapy or combined with MTX compared to MTX monotherapy in patients who had received no or minimal (limited exposure) csDMARDs and who were naive to biologic DMARDs. Baricitinib monotherapy was superior to MTX monotherapy (primary endpoint), with a higher ACR20 response rate (77% versus 62%; \( P \leq 0.01 \)).

Similar results were observed for combination therapy. Radiographic progression showed a statistically significant reduction in structural damage for baricitinib plus MTX compared to MTX monotherapy, but not for the monotherapy.

RA BEACON assessed the efficacy in bDMARDs-IR patients, including at least one anti-TNF inhibitor. At week 12, ACR 20 responses (primary end point) were 55% for baricitinib 4 mg and 49% for baricitinib 2 mg compared with 27% for placebo group (\( P < 0.001 \)).

RA BEAN included MTX IR patients. Study population was randomized to PBO, baricitinib 4 mg and Adalimumab 40 mg on background MTX. Comparisons between baricitinib and adalimumab were controlled for multiplicity with respect to ACR20 response and change from baseline in DAS28-CRP at week 12. Baricitinib plus MTX was non-inferior to adalimumab plus MTX for the ACR20 response, with a margin of 12% (70% vs 61% for adalimumab), and was therefore considered to be significantly superior to adalimumab (\( P = 0.01 \)). Radiographic progression was significantly lower for baricitinib compared with placebo (at 52 weeks change from baseline in mTSS was 0.71 vs 1.8, respectively).

RA BUILD compared in MTX-IR patients baricitinib 2 and 4 mg once a day to placebo. A statistically significant reduction in structural joint damage (radiographic outcome) from baseline to week 24 was observed for baricitinib 2 and 4 mg compared with placebo.

The long extension trail RA BEYOND included a substudy population for the assessment of a step-down dose strategy. Patients who were on baricitinib 4 mg for at least 15 months and who had achieved sustained LDA or remission were re-randomized to continue with 4 mg or stepping down to 2 mg. Most patients in both regimens (standard or step-down) were still in low disease activity or remission, but the step-down group had statistically significant increase in tender and swollen joint count, physician global assessment, DAS28-CRP, clinical disease activity index (CDAI), and SDAI scores. This demonstrated that the 4 mg dose is the most effective and that stepping down strategy is a valid option but not for all patients.

**Upadacitinib**

Upadacitinib is selective for JAK1 74-fold over JAK2. Six global phase III randomized controlled clinical trials (SELECT phase III program) evaluated the efficacy and safety of upadacitinib covering different RA subpopulations. The approved dose for RA patients is 15 mg once daily.

The SELECT NEXT study included active RA patients with an inadequate response to csDMARDs and the SELECT BEYOND trial patients with inadequate response or intolerance to bDMARDs. The SELECT MONOTHERAPY study showed the efficacy of upadacitinib monotherapy in clinical and functional outcomes vs methotrexate. In these studies, patients were randomized to upadacitinib 15 or 30 mg or placebo for at least 12 weeks. Overall, results showed a rapid statistically significant improvement in the ACR20 response, and in the ACR50 and ACR70 responses with upadacitinib 15 and 30 mg. Several patients reported outcomes like quality of life, physical function, fatigue, and duration of morning stiffness were also significantly improved in upadacitinib arms.

The SELECT EARLY study compared the clinical efficacy of upadacitinib monotherapy vs MTX monotherapy, in MTX-naive patients. Significantly, more patients receiving upadacitinib 15 mg and 30 mg vs MTX achieved both primary end points: ACR50 responses at week 12 (52.1% and 56.4% vs 28.3%) and DAS28[CRP] <2.6 at week 24 (48.3% and 50.0% vs 18.5%).

The SELECT COMPARE study evaluated the efficacy of upadacitinib as compared to PBO or adalimumab (ADA) in MTX IR patients. Patients were randomized to upadacitinib 15 mg, placebo, or ADA (40 mg every other week) while continuing to take a stable background dose of MTX. This study was designed and powered for superiority against placebo, noninferiority and superiority against ADA. Upadacitinib was superior to ADA based on the ACR50.
response rate (45% vs 29%, respectively, p ≤ 0.001), change in pain severity score (mean change –32.1 upadacitinib group vs –25.6 ADA group; P ≤ 0.001), and change in the Health Assessment Questionnaire Disability Index (mean change –0.60 upadacitinib group vs –0.49 ADA group; P ≤ 0.01). At week 26, more patients receiving upadacitinib than those receiving PBO or ADA achieved low disease activity or remission (P ≤ 0.001). The non-inferiority of upadacitinib compared to ADA was met for DAS28-CRP score of ≤3.2 (45% versus 29%, respectively). The SELECT-CHOICE was the other head-to-head, double-blind study in bDMARD-IR patients comparing the efficacy and safety of upadacitinib 15 mg to ABA IV, each in combination with stable background csDMARDs. The primary endpoint was the non-inferiority comparison of upadacitinib vs ABA in the change from baseline in DAS28(CRP) at Week 12. The results demonstrated that upadacitinib was non-inferior to ABA for the primary end point (change from baseline in DAS28(CRP) P < 0.001) and was superior to ABA for change from baseline in DAS28(CRP) (P < 0.001) and proportion of patients achieving DAS28(CRP) <2.6 remission (P < 0.001) at Week 12. A significant difference in the proportion of patients achieving DAS28(CRP) <2.6 was also maintained at Week 24.

The impact of upadacitinib on structural joint damage was assessed during SELECT-EARLY and SELECT-COMPARE, showing in both studies that upadacitinib significantly reduced progression of joint damage as determined by significantly lower change from baseline in modified Total Sharp Score (mTSS) compared with MTX arms. No difference was seen compared to ADA in the mTSS.

**Filgotinib**

Filgotinib is a JAK1 selective 30 fold more selective versus JAK2. The FINCH program includes four clinical phase III trials conducted also in different RA patient types that.

The FINCH 1 study included MTX-IR comparing: filgotinib 200 mg or filgotinib 100 mg once daily, subcutaneous adalimumab 40 mg every 2 weeks, or matching placebo all with background MTX up to week 52. The primary end point was met by filgotinib in both doses compared to placebo at week 12 (ACR 20%: 76.6% filgotinib 200, 69.8% filgotinib 100, 49.9% placebo P < 0.001 and 70.8% ADA). Non-inferiority of filgotinib 200 mg compared to ADA was met based on DAS28-CRP ≤3.2 but the response between patients treated with filgotinib 100 mg vs adalimumab was numerically similar (49.7% filgotinib 200, 38.8% filgotinib 100 mg, 43.4% ADA). Radiographic progression measured by change from baseline in mTSS vs placebo at week 24 was significantly lower in filgotinib 200 mg or 100 mg vs placebo (P<0.001).

The FINCH 2 study evaluated the efficacy of filgotinib vs placebo in bDMARDs-IR patients. The primary end point, ACR 20 response rates at week 12 was 66% (95% CI, 58.0–74.0%) and 57.5% (95% CI, 49.4–65.7%) of patients with filgotinib 200 mg and 100 mg, respectively, vs 31.1 (95% CI, 23.3–38.9%) for placebo (significant difference for both doses of filgotinib vs placebo P<.001).

The FINCH 3 trial included MTX naive, randomised to filgotinib 200 mg once daily plus MTX, filgotinib 100 mg plus MTX, filgotinib 200 mg monotherapy and MTX ≤ 20 mg weekly. At week 24, significantly more patients in the filgotinib 200 mg + MTX (81.0%; P<0.001) and filgotinib 100 mg + MTX (80.2%; P<0.05) arms achieved an ACR20 response compared to MTX monotherapy (71.4%) (primary end point). Filgotinib 200 mg monotherapy did not reach a significant difference compared to MTX on ACR20, although it did on ACR 50/70 at week 24. There was less radiographic progression as measured by change in mTSS from baseline at week 52 in patients receiving filgotinib 100/200 mg plus MTX and filgotinib 200 mg monotherapy vs MTX monotherapy.

The FINCH 4 is a study that is evaluating the long-term safety and tolerability of filgotinib in participants who have completed one of the parent studies of filgotinib in RA. The study is still active and partial results are expected soon.

The safety data from JAK inhibitors comes from the clinical development programs and post-marketing surveillance. All JAKi produce changes in laboratory parameters that may differ between the drugs in relation of the selectivity for each JAK. There have been changes in blood cell counts (risk of lymphopenia), haemoglobin levels, liver transaminase, creatine kinase, cholesterol and creatine. A higher risk of infection was reported with JAKi, and the most common serious infection reported for tofacitinib and baricitinib was pneumonia. There is difference between JAKi and other bDMARDs regarding the risk of herpes zoster infection, which is higher with these drugs and is most marked in Japanese and Korean ethnicity patients. Concerns have been raised for a potential risk of thromboembolic events (pulmonary embolism and deep vein thrombosis) with these drugs that are still under evaluation.
Table 3  Up to Date 15 Biosimilars Have Been Approved by the FDA for Rheumatic Diseases

| Commercial Name | Compound Name   | Date Approved | Reference Product     |
|-----------------|-----------------|---------------|-----------------------|
| Ibrili           | Rituximab-arx    | December 2020 | Rituxan (rituximab)   |
| Hulio            | Adalimumab-fkjq  | July 2020     | Humira (adalimumab)   |
| Avsola           | Infliximab-axxq  | December 2019 | Remicade (infliximab) |
| Abrilada         | Adalimumab-afzb  | November 2019 | Humira (adalimumab)   |
| Hadlima          | Adalimumab-bwwd  | July 2019     | Humira (adalimumab)   |
| Ruxience         | Infliximab-pvvr  | July 2019     | Rituxan (rituximab)   |
| Eticovo          | Etanercept-ykro  | April 2019    | Enbrel (etanercept)   |
| Truxima          | Rituximab-abb    | November 2018 | Rituxan (rituximab)   |
| Hyrimoz          | Adalimumab-adaz  | October 2018  | Humira (adalimumab)   |
| Ixfi             | Infliximab-qbx   | December 2017 | Remicade (infliximab) |
| Cytezo           | Adalimumab-adbm  | December 2017 | Humira (adalimumab)   |
| Renflexis        | Infliximab-abda  | May 2017      | Remicade (infliximab) |
| Amjevira         | Adalimumab-atso  | September 2016| Humira (adalimumab)   |
| Erelzi           | Etanercept-szzi  | August 2016   | Enbrel (etanercept)   |
| Inflectra        | Infliximab-dyyb  | April 2016    | Remicade (infliximab) |

Biosimilars

A biosimilar is an agent that presents a similar molecular structure of the active substance of an already approved agent, the reference product and is intended to be used in the same way as the reference product. It must have similar biological activity and quality characteristics and should have no significant safety nor efficacy differences. To get a biosimilar to be approved, it must undergo a thorough full development process that involves a series of comparability exercises to establish biosimilarity to the reference product and at least one randomized controlled trial (Tables 3 and 4). The aim of biosimilarity clinical trials is not to establish efficacy per se, which has already been established in clinical trials conducted with the reference product, but to demonstrate equivalent clinical performance of the biosimilar in relation to the reference product. The same quality manufacturing standards that apply to the original biologic also apply to the biosimilar. When biosimilarity is demonstrated in one indication, this can be extrapolated to other approved indications of the reference product.

The approval of biosimilars can help the health care systems worldwide to make substantial savings. If patients receiving a reference biological product are switched to biosimilars, and if biological-naive patients are started on biosimilars rather than reference products, it could save economic resources as long as the cost is much lower. Thus, improving the access of the population to biologic treatment in case it is needed and provide an earlier usage.

Emerging Therapies

There are up to date some promising molecules with different mechanism of actions in the pipeline to be approved.

Otilimab is a human monoclonal antibody that inhibits granulocyte-macrophage colony-stimulating factor (GM-CSF), a key driver in immune-mediated inflammatory conditions. A phase 2b, dose-ranging, multicenter, placebo-controlled study (Baroque study) was done. A total 222 patients who were receiving stable methotrexate were randomly assigned to six different groups subcutaneous placebo or otilimab 22.5 mg, 45 mg, 90 mg, 135 mg, or 180 mg, plus methotrexate, once weekly for 5 weeks, then every other week until week 50. Otilimab plus methotrexate was well tolerated and, despite not achieving the primary endpoint of DAS28-CRP remission, there were improvements compared with placebo in disease activity scores. Patients reported significant improvement in pain and physical function, supporting further clinical development of otilimab in...
Table 4 Adding to This List the EMA Has Approved 16 More Biosimilars

| Commercial Name | Date Approved | Reference Product |
|-----------------|---------------|-------------------|
| Remsima         | September 2013| Rituximab         |
| Iraldi          | September 2013| Infliximab        |
| Nepexto         | May 2015      | Etanercept        |
| Ritemvia        | July 2017     | Rituximab         |
| Blitzima        | July 2017     | Rituximab         |
| Halimazo        | July 2018     | Adalimumab        |
| Idacio          | April 2019    | Adalimumab        |
| Asparity        | February 2020 | Adalimumab        |
| Zessly          | May 2018      | Infliximab        |
| Hyrimoz         | May 2016      | Adalimumab        |
| Flixabi         | May 2016      | Infliximab        |
| Ruxience        | April 2020    | Rituximab         |
| Kromays         | April 2019    | Adalimumab        |
| Riximyo         | June 2017     | Rituximab         |
| Solymbic        | March 2017    | Adalimumab        |
| Riexathon       | June 2017     | Rituximab         |
| Benepali        | January 2016  | Etanercept        |

rheumatoid arthritis. The phase III clinical program (named "ContRAst") is ongoing. It compares otilimab against placebo (ContRAst 1) and against two treatments with different modes of actions: tofacitinib (ContRAst 2) and sarilumab (ContRAst 3). The program also enrolls a broad range of difficult-to-treat patients who have had an inadequate response to or have been unable to tolerate currently available treatments. Patients who complete the pivotal studies may be eligible to participate in a long-term extension study to further evaluate the efficacy and safety of otilimab for up to 4 years.

Olokizumab is a monoclonal antibody that targets interleukin (IL) 6. There are currently 4 clinical trials to evaluate the treatment of moderate to severe active rheumatoid arthritis (RA) in adults for whom methotrexate is inadequate or had presented inadequate response to anti-TNF alpha blockers. The CREDO 1, CREDO2 and CREDO 3 (core studies) evaluate the efficacy and safety of different regimens of subcutaneous Olokizumab compared to placebo (CREDO 1 and 3) or adalimumab (CREDO 2) CREDO 4 evaluates the long-term safety, tolerability and efficacy of two dosing regimens of Olokizumab (OKZ), in subjects with Rheumatoid Arthritis (RA) who previously completed 24 weeks of blinded treatment in one of the core studies.

ABX464 is a small molecule that produces a specific and selective induction of miR-124 in immune cells. miR-124 is a crucial modulator of inflammation and innate immunity that could provide therapeutic restitution of physiological pathways lost in inflammatory diseases. The Phase 2a ongoing study, ABX464-301, investigates the safety and tolerability of ABX464 in combination with methotrexate in patients with moderate-to-severe active RA. Patients enrolled in the study had an inadequate response to methotrexate or and to one or more anti-tumor necrosis factor alpha (TNFα) therapies. Patients who complete the ABX464-301 trial, have the possibility to roll over into a Phase 2a open-label study, ABX464-302, aiming at the evaluation of the one-year safety and efficacy of ABX464 as maintenance therapy in RA.

Iscalimab (CFZ533) is a fully human, aglycosylated non-depleted monoclonal antibody that blocks CD154-CD40 pathway activation that is being developed as an immunosuppressive agent. The CD40-CD154 costimulatory pathway is essential for the generation of T cell-dependent antibody responses (TDAR), germinal center (GC) formation, and memory B cell differentiation. In macrophages and dendritic cells, it regulates their activation and differentiation as well as antigen presentation to T cells. Due to its mechanism of action, Iscalimab shows to be a promising therapy in transplant rejection and autoimmune diseases. The first Phase 1 randomized, double-blind, placebo-controlled, parallel-group, 2-part study clinical trial was conducted between January 2013 and February 2017, to assess pharmacokinetics, pharmacodynamics, safety, and tolerability of ascending intravenous doses of iscalimab. In this trial, there was a cohort of RA between 18 and 65 years old. Iscalimab showed to be safe and well tolerated at single doses up to 30 mg/kg IV with no evidence of increased risk of infection or thromboembolic complications. There are ongoing phase 2 studies in Graves’ disease, liver transplant rejection, lupus nephritis, myasthenia gravis, renal transplant rejection, Sjögren’s syndrome, systemic lupus erythematosus and type 1 diabetes mellitus.

Conclusion
The better understanding of the pathogenesis of RA has allowed the development of a great number of effective therapies for the treatment of this disease. Even though the
menu is wide, there is still a lack of reliable tools for predicting which patients will respond better to a given drug. The increasing numbers of head-to-head trials is essential in order to answer the question of which therapy is more effective and safer for each individual patient. Also, more trials dealing with tapering or stopping therapy are still needed to help optimize the use of the existing drugs. Remission of LDA persistence remains a goal to achieve.

Despite the advances in therapies, the most important strategy is the early diagnosis and treatment, and the frequent assessments of disease activity with adjustments in therapy for achieving clinical remission or low disease activity in most patients.

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