

Disease Modification in Individuals with
MODERATE-TO-SEVERE
RHEUMATOID ARTHRITIS:
Optimizing Treatment Through the Finely Tuned Selectivity for JAKs



MEETING INFO

TeleECHO Presentation - Tuesday, February 8, 2022

3:00 – 4:00 PM ET

2:00 – 3:00 PM CT

1:00 – 2:00 PM MT

12:00 – 1:00 PM PT

FACULTY

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This activity is supported by an educational grant from AbbVie Inc.



Disease Modification in Individuals with **MODERATE-TO-SEVERE RHEUMATOID ARTHRITIS:** Optimizing Treatment Through the Finely Tuned Selectivity for JAKs

Agenda

I. Managing Rheumatoid Arthritis (RA)

- a. Features of rheumatoid arthritis
- b. RA pathophysiology
- c. Treatment options for RA
- d. Guideline recommended care of RA
 - i. Initial DMARD choice
 - ii. Options after inadequate response to initial therapy
- e. Treating to target

II. JAK Inhibitors for the Management of Moderate-to-Severe RA

- a. JAK inhibitor mechanism of action and selectivity
- b. Review of clinical trial data of the efficacy and safety of JAK inhibitors:
 - i. Monotherapy
 - ii. Combination therapy
 - iii. Patients with inadequate response to methotrexate (MTX IR)
 - iv. MTX naive populations
 - v. Long-term safety data

III. Application of Societal Guidelines for Moderate-to-Severe Disease – ACR & EULAR

- a. Focus on JAK inhibition and its place in management of patients
- b. Comorbid conditions and RA treatment considerations

IV. Conclusions

V. Interactive Case Studies

VI. Q & A

Disease Modification in Individuals with Moderate-to-Severe Rheumatoid Arthritis: Optimizing Treatment Through the Finely Tuned Selectivity for JAKs

FACULTY

Jon T. Giles, MD, MPH (Program Chair)

Associate Professor of Medicine

Division of Rheumatology

Columbia University, College of Physicians & Surgeons

New York, NY

PROGRAM OVERVIEW

This TeleECHO series will explore the use of JAK inhibitors for the management of rheumatoid arthritis (RA) through interactive case studies. Faculty will review emerging efficacy and safety data, discuss guideline recommended care of RA, and help clinicians identify candidates for treatment with JAK inhibitors.

TARGET AUDIENCE

This CME initiative is designed to meet the educational needs of rheumatologists, nurses and allied healthcare professionals involved in the care of patients with rheumatoid arthritis.

LEARNING OBJECTIVES

Upon the completion of this program, attendees should be able to:

- Review the members of the JAK family, involvement in RA pathology, and selectivity of JAK targets and associated effects
- Discuss the clinical trials findings for Janus Kinase inhibitors in patients with moderate-to-severe RA
- Describe the application of ACR/EULAR management recommendations to clinical practice for patients with moderate-to-severe rheumatoid arthritis

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The reviewer of this activity has nothing to disclose.

CNE Content Review

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Douglas Cox, MSN, MHA, RN

Ultimate Medical Academy/CCM – Lead Nurse Planner

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***Disease Modification in Individuals with
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Optimizing Treatment Through the
Finely Tuned Selectivity for JAKs***

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Accreditation

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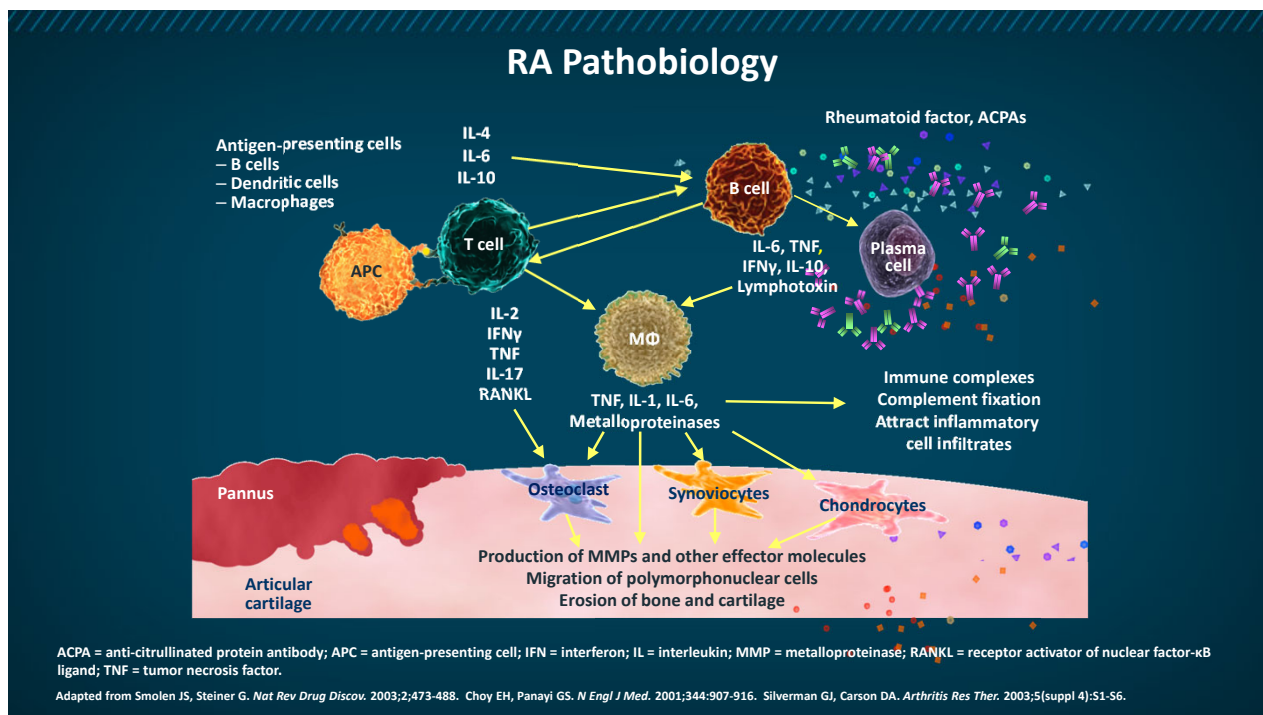
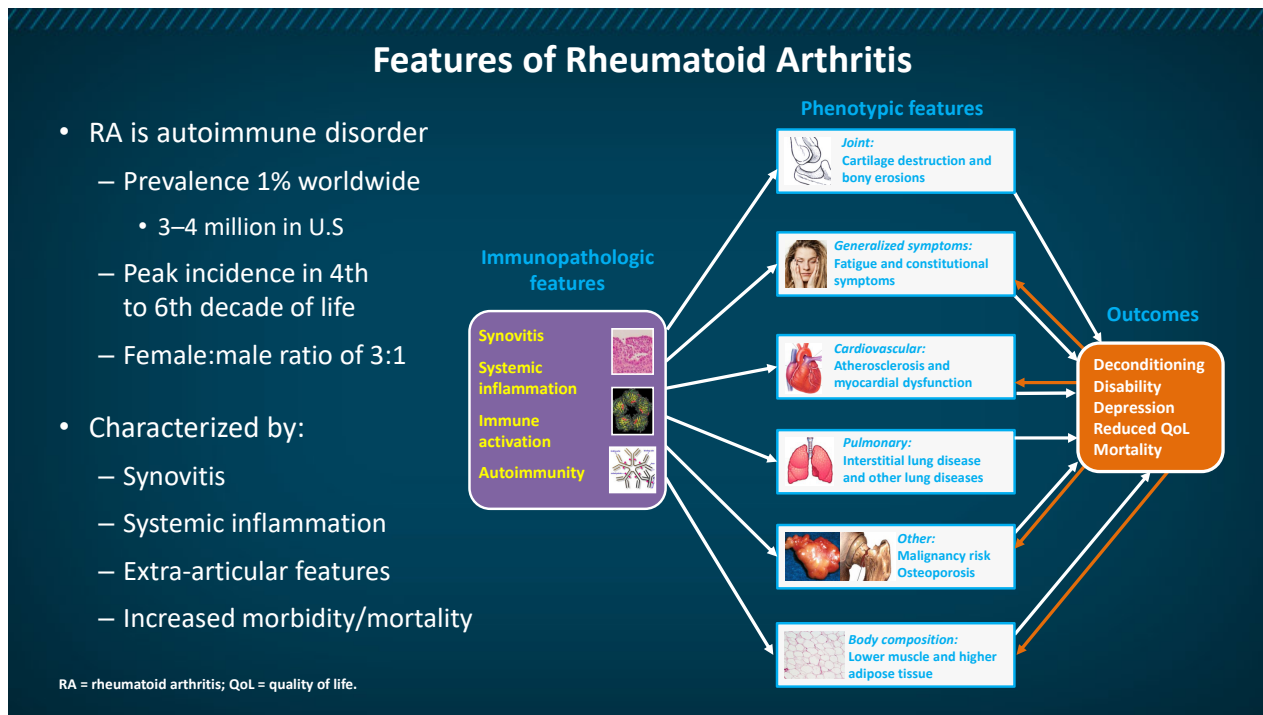
- Jon T. Giles, MD, MPH has received consulting fees from AbbVie, Bristol Myers Squibb, Lilly, Gilead, and UCB. He also states a financial relationship with Pfizer.
- During the course of this lecture, Dr. Giles may mention the use of medications for both FDA-approved and non-approved indications.

This activity is supported by an educational grant from AbbVie Inc.

Learning Objectives

- Review the members of the JAK family, their involvement in RA pathology, and selectivity of JAK targets and associated effects
- Discuss the clinical trials findings for JAK inhibitors in patients with moderate-to-severe RA
- Describe the application of ACR/EULAR management recommendations to clinical practice for patients with moderate-to-severe rheumatoid arthritis

JAK = Janus kinase; RA = rheumatoid arthritis; ACR = American College of Rheumatology; EULAR = European League Against Rheumatism.



Disease-Modifying Pharmacotherapies for RA Available in US

Oral non-biologics

Methotrexate
Sulfasalazine
Hydroxychloroquine
Leflunomide
Azathioprine
Cyclosporine
Tacrolimus
Cyclophosphamide

Prednisone

Selective cytokine inhibitors

TNF inhibitors

Etanercept
Infliximab
Adalimumab
Certolizumab
Golimumab

IL-1 inhibitors

Anakinra

IL-6 inhibitors

Tocilizumab
Sarilumab

B-cell depletion

Rituximab

T cell co-stimulation blockade

Abatacept

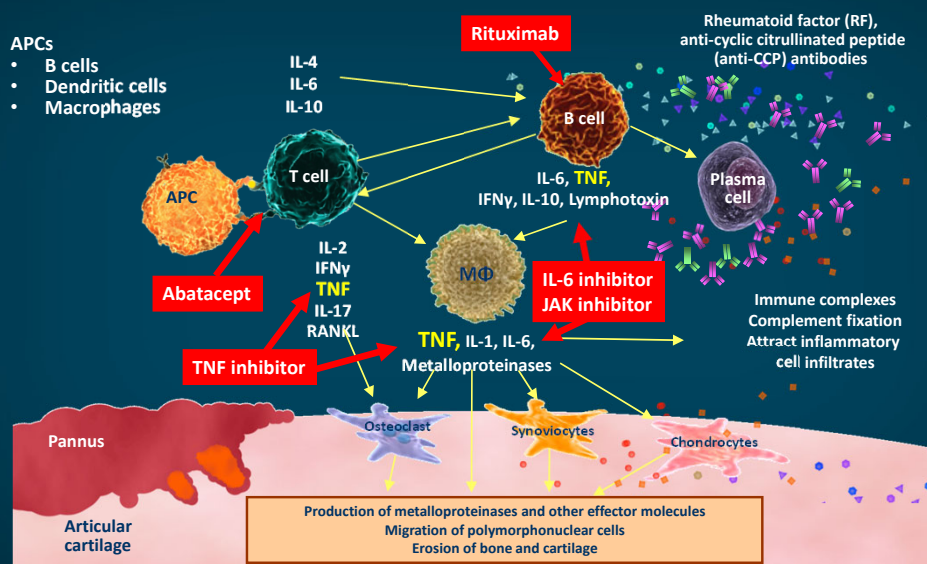
JAK inhibition

Tofacitinib
Baricitinib
Upadacitinib

US = United States; JAK = Janus kinase.

Köhler BM, et al. *J Clin Med.* 2019;8:938.

Targets of Therapy in RA



Adapted from Smolen JS, Steiner G. *Nat Rev Drug Discov.* 2003;2:473-488. Choy EH, Panayi GS. *N Engl J Med.* 2001;344:907-916. Silverman GJ, Carson DA. *Arthritis Res Ther.* 2003;5(suppl 4):S1-S6.

Initial DMARD Choice

American College of Rheumatology (ACR) 2021 RA Treatment Guideline

Initiation of DMARD treatment of patients with RA	
ACR recommendations	Certainty of evidence
Initiation of treatment in DMARD-naïve patients with MODERATE-TO-HIGH disease activity Methotrexate (MTX) monotherapy is strongly recommended over: Hydroxychloroquine or sulfasalazine bDMARD or tsDMARD monotherapy Combination of MTX + non-TNFi bDMARD or tsDMARD MTX monotherapy is conditionally recommended over: Leflunomide Dual or triple csDMARD therapy Combination of MTX + TNFi Initiation of csDMARD without short-term (<3 months) glucocorticoids is conditionally recommended over initiation of csDMARD with short-term glucocorticoids Initiation of csDMARD without longer-term (≥3 months) glucocorticoids is strongly recommended over initiation of csDMARD with longer-term glucocorticoids	Very low/low Very low/moderate Low/very low Low Moderate Low Very low Moderate
Initiation of treatment in DMARD-naïve patients with LOW disease activity Hydroxychloroquine is conditionally recommended over other csDMARDs Sulfasalazine is conditionally recommended over MTX MTX is conditionally recommended over leflunomide	Very low Very low Very low
Initiation of treatment in csDMARD-treated, but MTX-naïve, patients with MODERATE-TO-HIGH disease activity MTX monotherapy is conditionally recommended over combination of MTX + bDMARD or tsDMARD	Moderate/very low

DMARD = disease-modifying antirheumatic drug; bDMARD = biologic DMARD; tsDMARD = targeted-synthetic DMARD; csDMARD = conventional synthetic DMARD; TNFi = TNF inhibitor.

Fraenkel L, et al. *Arthritis Care Res.* 2021;73:924-939.

Failure of Initial Treatment Choice

ACR 2021 RA Treatment Guideline

Treatment modification following inadequate response to initial treatment	
Recommendations	Certainty of evidence
T2T approach is strongly recommended over usual care for patients who have not been previously treated with bDMARDs or tsDMARDs	Low
T2T approach is conditionally recommended over usual care for patients who have had an inadequate response to bDMARDs or tsDMARDs	Very low
Minimal initial treatment goal of low disease activity is conditionally recommended over goal of remission	Low
Addition of a bDMARD or tsDMARD is conditionally recommended over triple therapy for patients taking maximally tolerated doses of MTX who are not at target	Very low
Switching to bDMARD or tsDMARD of different class is conditionally recommended over switching to bDMARD or tsDMARD belonging to same class for patients taking bDMARD or tsDMARD who are not at target	Very low
Addition of/switching to DMARDs is conditionally recommended over continuation of glucocorticoids for patients taking glucocorticoids to remain at target	Very low
Addition of/switching to DMARDs (± IA glucocorticoids) is conditionally recommended over use of IA glucocorticoids alone for patients taking DMARDs who are not at target	Very low

T2T = treat-to-target; IA = intraarticular.

Fraenkel L, et al. *Arthritis Care Res.* 2021;73:924-939.

The Principles of Treating to Target

- T2T by measuring disease activity and adjusting therapy accordingly will result in better patient outcomes
- The primary target for treatment should be clinical remission, defined as the absence of signs and symptoms of significant inflammatory disease activity
- In some cases, low disease activity (LDA) may be an acceptable treatment goal, particularly in patients with long-standing, established disease

Smolen JS, et al. *Ann Rheum Dis*. 2010;69:631-637.

So, Can T2T Be Implemented in the Real World? A First Look...

- RA BIODAM (**B**IOmarkers of joint **DAM**age): RA registry in 10 countries^{1,2}
 - At 41% of visits, T2T protocol was not implemented¹
 - Main reason: physician felt current treatment was adequate (69%)²
- CORRONA (**C**onsortium of **R**heumatology **R**esearchers of **N**orth **A**merica)^{3,4}
 - Failure to escalate occurred at ~50% of visits
 - 2 main reasons: doctor said treatment needed more time to work (eg, 3 more months) or patient refused
- Implications for patients with persistently active disease
 - Patients and doctors feel that current treatment is “good enough”
 - Patients are risk averse to add/change therapy

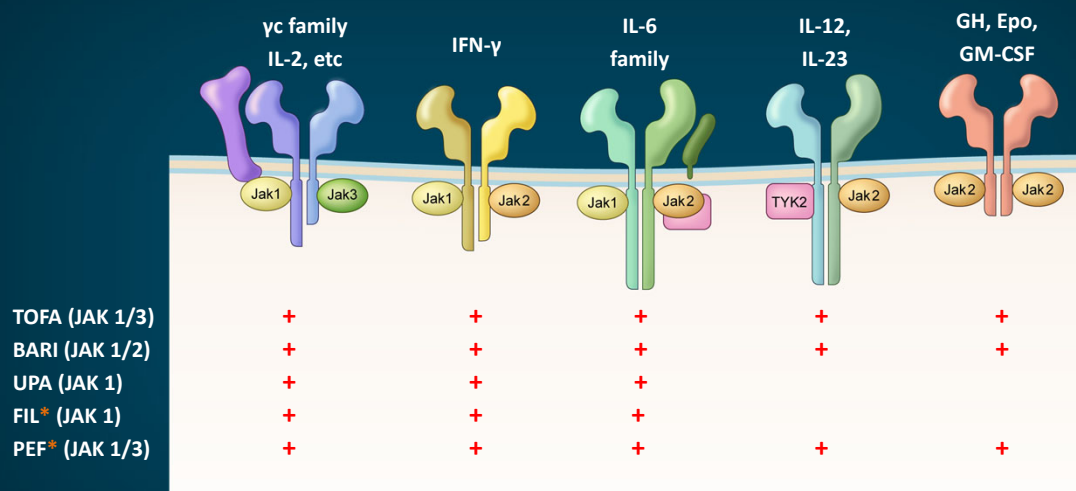
1. Ramiro S, et al. *Ann Rheum Dis*. 2020;79:453-459. 2. Maksymowych WP, et al. *Arthritis Rheumatol*. 2014;66(suppl 10):abstract 2912. 3. Harrold L, et al. *Arthritis Rheumatol*. 2015;67(suppl 10): abstract 3185. 4. Harrold LR, et al. *Arthritis Care Res (Hoboken)*. 2018;70:379-387.

Unmet Need for RA Treatments

- Rates of “low disease activity” and “remission” have improved over time
 - 5 years after symptom onset in 2009, 45% reached sustained remission (SR) compared with only 16% in 1999
 - Largely due to the availability of new treatments and adoption of T2T
- However, SR is only observed in ~25%
 - Fluctuating disease activity/flares
 - Secondary loss of response
 - Treatment resistance in a subset
 - Intolerance/side effects
- Complete treatment refractoriness is rare

Einarsson JT, et al. *Rheumatology (Oxford)*. 2020;59:205-212. Mierau M, et al. *Rheumatology (Oxford)*. 2007;46:975-979. Prince FHM, et al. *Arthritis Res Ther*. 2012;14:R68.

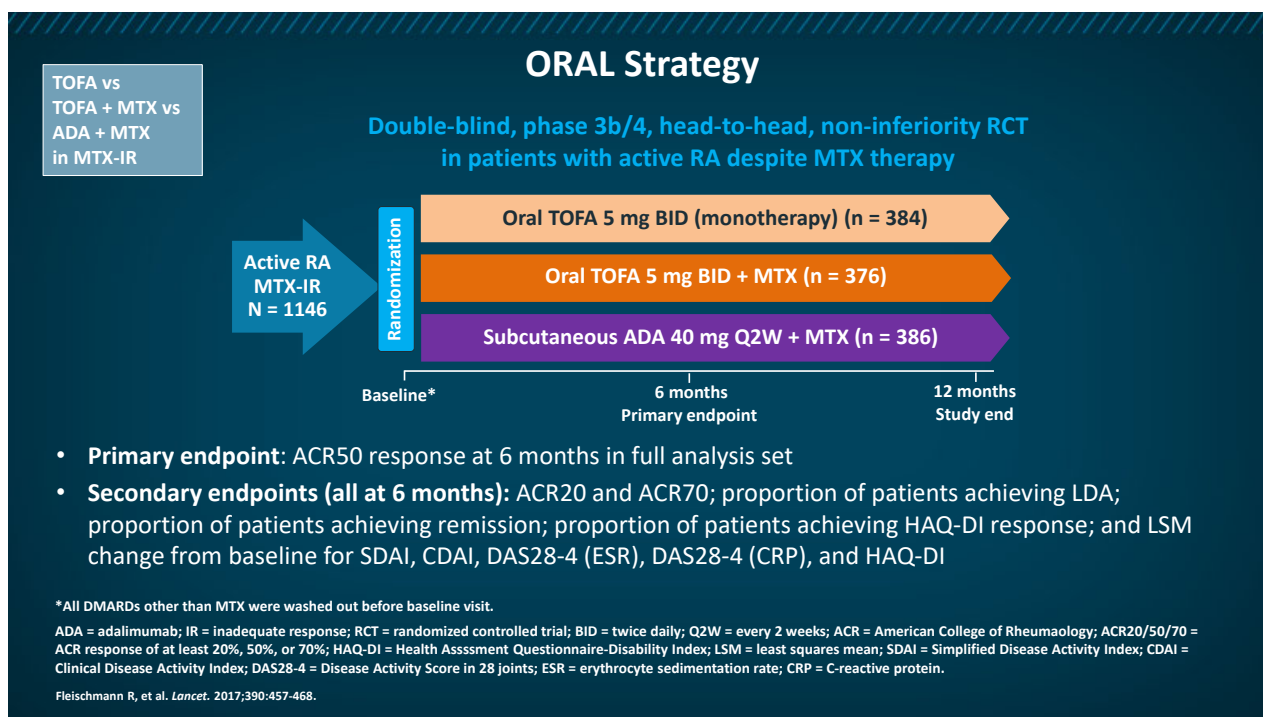
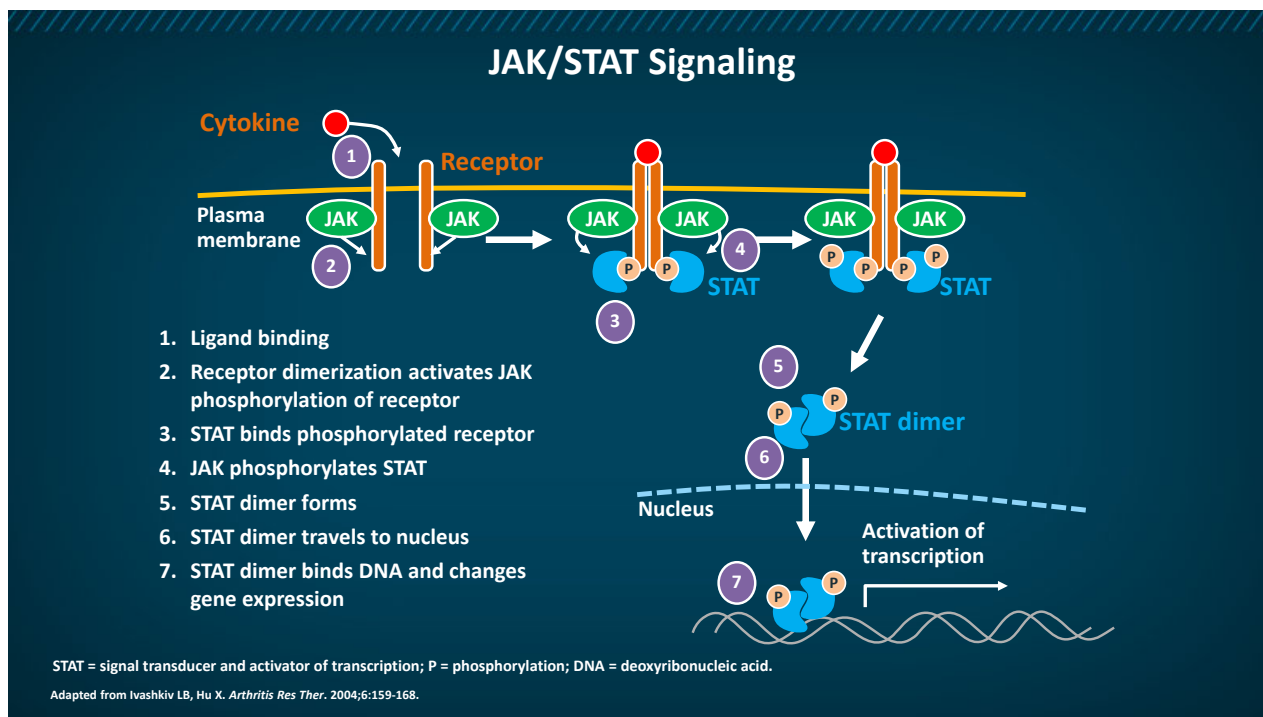
Signaling by Type I/II Cytokine Receptors and JAK Inhibitors



*Not FDA approved.

GH = growth hormone; Epo = erythropoietin; GM-CSF = granulocyte macrophage colony-stimulating factor; TYK = tyrosine kinase; TOFA = tofacitinib; BARI = baricitinib; UPA = upadacitinib; FIL = filgotinib; PEF = peficitinib.

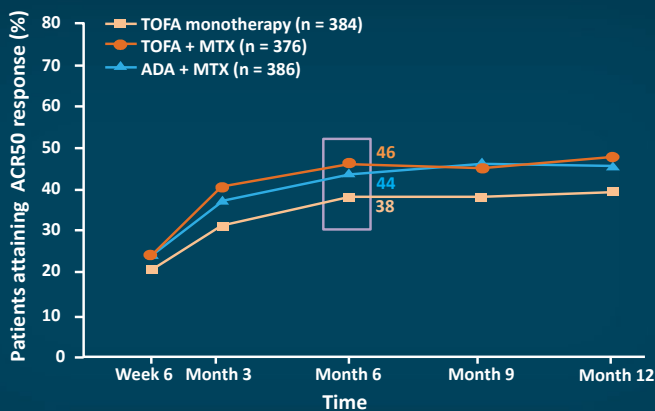
Fragoulis GE, et al. *J Allergy Clin Immunol*. 2021;148:941-952.



TOFA vs
TOFA + MTX vs
ADA + MTX
in MTX-IR

ORAL Strategy: Efficacy

Primary Endpoint: ACR50 at 6 Months



- Tofacitinib + MTX non-inferior to adalimumab + MTX
- Tofacitinib monotherapy did not meet non-inferiority criteria compared with tofacitinib + MTX
- Tofacitinib monotherapy did not meet non-inferiority criteria compared with adalimumab + MTX

Fleischmann R, et al. *Lancet*. 2017;390:457-468.

ORAL Strategy

AEs, Serious AEs, and Discontinuations

	TOFA monotherapy (n = 384)	TOFA + MTX (n = 376)	ADA + MTX (n = 386)
Total number of AEs, n	598	652	620
Patients with AEs, n (%)	226 (59%)	231 (61%)	253 (66%)
Patients with treatment-related AEs, n (%)	101 (26%)	111 (30%)	133 (35%)
Patients with serious AEs, n (%)	35 (9%)	27 (7%)	24 (6%)
Patients discontinuing due to AEs, n (%)	23 (6%)	26 (7%)	37 (10%)
Patients with severe AEs, n (%) (defined by investigator)	24 (6%)	17 (5%)	23 (6%)
Deaths, n (%)	2 (1%)	0	0
AEs of special interest, n (%)			
Serious infections	6 (2%)	10 (3%)	6 (2%)
HZ (serious and non-serious)	4 (1%)	8 (2%)	6 (2%)
HZ (serious and non-serious) in vaccinated patients	1/69 (1%)	2/75 (3%)	0/72 (0%)
Opportunistic infections (not TB)	2 (1%)	1 (< 1%)	2 (1%)
TB	0	2 (1%)	0
MACE (non-fatal)	0	0	2 (1%)
Malignancy (not non-melanoma skin cancer)	1 (< 1%)	0	0
Non-melanoma skin cancer	2 (1%)	0	1 (< 1%)

AE = adverse event; HZ = herpes zoster; TB = tuberculosis; MACE = major adverse cardiovascular event.

Fleischmann R, et al. *Lancet*. 2017;390:457-468.

Venous Thromboembolism (VTE) With JAK Inhibitors

Primary analysis: VTE events identified from inpatient claims with as-treated follow-up approach

Data source and exposure group	VTE events, n	Total PYs of follow-up	Incidence rate per 100 PYs (95% CI)	Unadjusted HR (95% CI)	Propensity score-adjusted HR (95% CI)
Truven MarketScan					
TNFi initiators (n = 32,164)	98	28,951	0.34 (0.27–0.41)	Reference	Reference
TOFA initiators (n = 1910)	8	1326	0.60 (0.26–1.19)	1.70 (0.82–3.49)	1.55 (0.75–3.18)
Medicare claims					
TNFi initiators (n = 16,091)	117	12,660	0.92 (0.76–1.11)	Reference	Reference
TOFA initiators (n = 995)	<11*	625	1.12 (0.45–2.31)	1.16 (0.54–2.49)	1.12 (0.52–2.40)
Pooled					
TNFi initiators (n = 48,255)	215	41,611	0.52 (0.45–0.59)	Reference	Reference
TOFA initiators (n = 2905)	15	1951	0.77 (0.43–1.27)	1.42 (0.84–2.40)	1.33 (0.78–2.24)

- Higher rates of VTE noted in interim analysis of large safety study for tofacitinib 10 mg BID
- EMA recommended caution for use of JAK inhibitors for people at risk for VTE
- Rates of VTE across phase 3 trials of all studied JAK inhibitors ≈ 0.5/1000 PYs

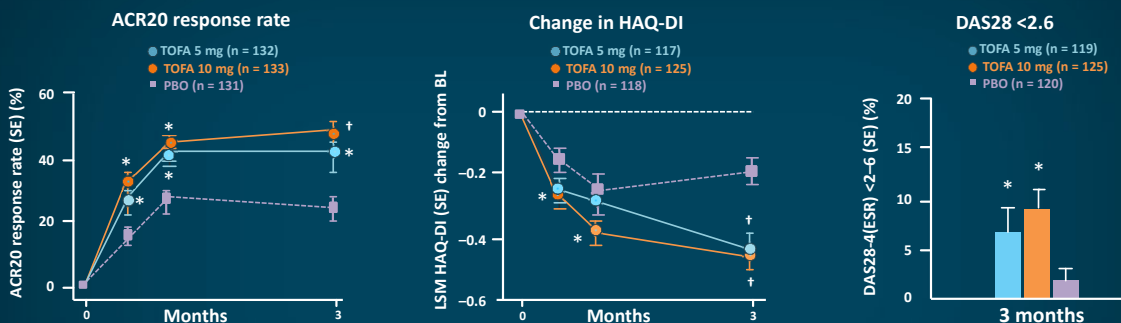
*Actual number suppressed; required by data use agreement with Medicare and Medicaid Services for counts <11.

HR = hazard ratio; CI = confidence interval; PY = person-year; EMA = European Medicines Agency.

Desai RJ, et al. *Arthritis Rheumatol.* 2019;71:892-900. Sandborn WJ, et al. *Aliment Pharmacol Ther.* 2019;50:1068-1076. EMA (www.ema.europa.eu/en/medicines/human/referrals/xeljanz). Accessed 12/21/2021.

Efficacy and Safety of Tofacitinib (JAK 1/3) + MTX in TNFi-IR Patients

Phase 3 trial outcomes for treatments at 3 months¹



- Tofacitinib has consistent, manageable safety profile across studies; no new safety signals¹
- Consistent safety through 114 months; sustained clinical efficacy through 96 months²
- 5-year analysis of AEs with tofacitinib vs bDMARDs reported in CORRONA RA registry shows higher incidence of HZ but not MACE or other serious infections³

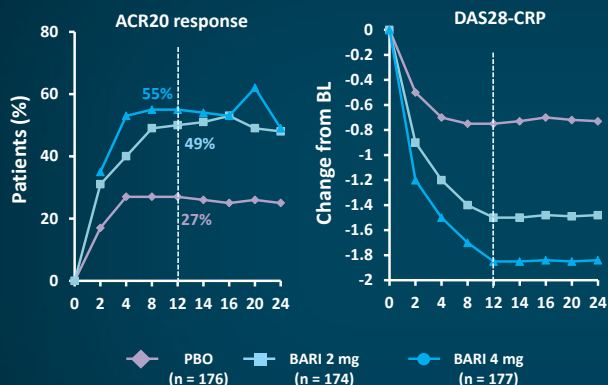
*P ≤ .05, †P < .0001 vs PBO.

PBO = placebo; SE = standard error; BL = baseline.

1. Burmester GR, et al. *Lancet.* 2013;381:451-460. 2. Wollenhaupt J, et al. *ACR/ARHP* 2017: abstract 522. 3. Kremer J, et al. *EULAR* 2019: abstract OP0028.

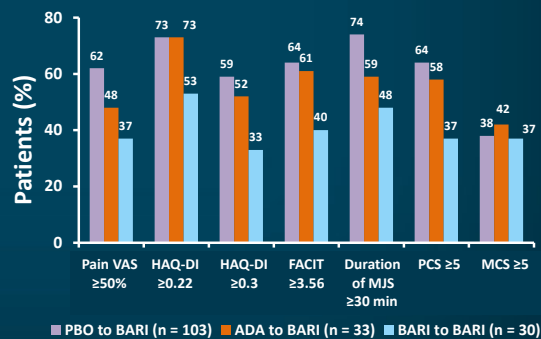
Efficacy and Safety of Baricitinib (JAK 1/2) in bDMARD-IR Patients

RA-BEACON phase 3 trial^{1,2}



RA-BEAM phase 3 trial³

Patients meeting/exceeding clinically relevant improvement thresholds at week 12 after rescue



VAS = visual analog scale; FACIT-F = Functional Assessment of Chronic Illness Therapy-Fatigue; MJS = morning joint stiffness; PCS = physical component score; MCS = mental component score; min = minute(s).

1. Genovese MC, et al. *N Engl J Med*. 2016;374:1243-1252 and supplement. 2. Genovese MC, et al. *Rheumatology (Oxford)*. 2018;57:900-908. 3. Fautrel B, et al. ACR/ARHP 2017: abstract 508.

Baricitinib

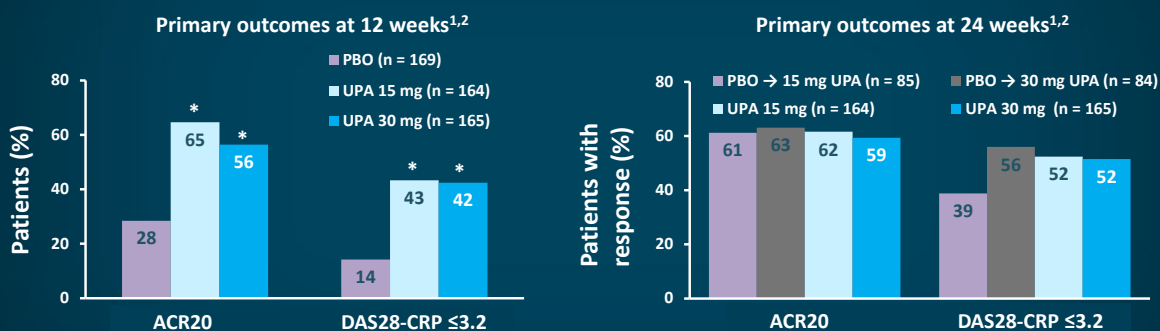
- Baricitinib led to more AEs (including infections) than placebo¹
- Most common infections were respiratory tract, urinary tract, bronchitis¹
- Rates of serious AEs through week 24 were similar among patient groups¹
- Post-hoc analysis of RA-BEACON: ORs favored use of baricitinib over placebo regardless of bDMARD history (number or type)²
- 7-year integrated safety analysis from phase 3 trials show increased risk for HZ and deep vein thrombosis/pulmonary embolism but not for malignancy³

OR = odds ratio.

1. Genovese MC, et al. *N Engl J Med*. 2016;374:1243-1252. 2. Genovese MC, et al. *Rheumatology (Oxford)*. 2018;57:900-908. 3. Genovese MC, et al. EULAR 2019: abstract THU0078.

Efficacy and Safety of Upadacitinib (JAK 1) in bDMARD-IR Patients

Phase 3 trial SELECT-BEYOND



*P < .0001 relative to PBO.

1. Genovese MC, et al. *Lancet*. 2018;391:2513-2524. 2. Genovese MC, et al. *ACR/ARHP* 2017: abstract 10L.

Upadacitinib

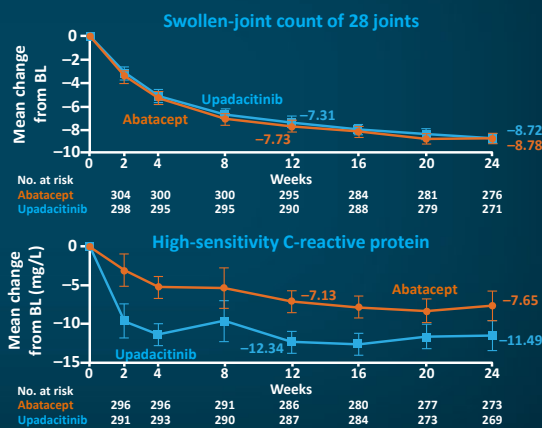
- In a post-hoc analysis, upadacitinib showed comparable efficacy whether administered with methotrexate or with other non-methotrexate csDMARDs¹
- UPA treatment also resulted in significant, clinically meaningful improvements in PROs among patients²
- Pooled safety analysis across phase 3 trials showed higher risk for serious infections and HZ but not for VTE, MACE, or malignancy (vs comparators)³
- Safety and efficacy remained consistent after 60 weeks of treatment⁴

PRO = patient-reported outcome.

1. Kremer J, et al. *Ann Rheum Dis*. 2019;78(suppl 2):A749 (abstract FRI0155). 2. Strand V, et al. *Ann Rheum Dis*. 2018;77(suppl 2):990 (abstract SAT0255). 3. Cohen SB, et al. *Ann Rheum Dis*. 2019;78(suppl 2):357 (abstract THU0167). 4. Genovese MC, et al. *Ann Rheum Dis*. 2019;78(suppl 2):360 (abstract THU0172).

Efficacy and Safety of Upadacitinib vs Abatacept in bDMARD-IR Patients

- Phase 3 trial SELECT-CHOICE
- Assessed non-inferiority (primary outcome) and superiority (secondary outcome) of oral upadacitinib 15 mg daily vs IV abatacept monthly for change in DAS28-CRP
 - Stable background non-biologic DMARDs continued
 - Primary endpoint measured at week 12
- Upadacitinib was non-inferior and superior to abatacept at week 12
 - Primary driver was change in CRP
- Remission at week 12 higher for upadacitinib vs abatacept
 - 30.0% vs 13.3% ($P < .001$)
- Adverse events higher for upadacitinib
 - Serious infection, MACE, VTE
 - HZ not higher, but low frequency at week 12

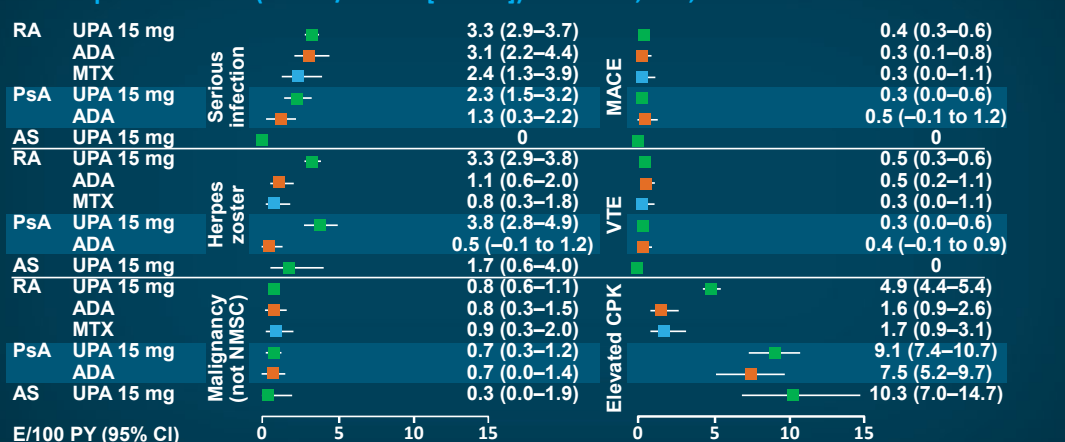


IV = intravenous.

Rubbert-Roth A, et al. *N Engl J Med.* 2020;383:1511-1521.

Longer Term Safety of Upadacitinib from Open-Label Trial Extensions Across Indications and Compared with Adalimumab and MTX

AEs of special interest (events/100 PY [95% CI]) across RA, PsA, and AS



4298 patients received ≥ 1 dose UPA 15 mg (n = 3209 RA; n = 907 PsA; n = 182 AS)

PsA = psoriatic arthritis; AS = ankylosing spondylitis; NMSC = non-melanoma skin cancer; CPK = creatine phosphokinase.

Burmester G, et al. *ACR* 2021, abstract 1691.

ORAL Surveillance Safety Trial: Tofacitinib vs TNF Inhibitors

Randomized, Open-Label Phase 4 Non-inferiority Trial

Inclusion/exclusion criteria

- Moderate/severe RA MTX-IR
- Age ≥50 years
- ≥1 CVD risk factor
- No current/prior malignancy

R
1:1:1
N = 4362

Study start **Protocol amendment** **Study completed**

MAR 2014 FEB 2019 JUL 2020

TOFA 5 mg BID + MTX (n = 1455)
TOFA 10 mg BID + MTX (n = 1456) TOFA 10 mg → 5 mg BID
TNFi + MTX (n = 1451)

ADA 40 mg Q2W (North America) or ETN 50 mg QW (RoW)

- Co-primary outcomes: MACE and malignancy (combined TOFA doses vs TNFi)
 - Non-inferiority margin set at 1.8
- After a signal for VTE was detected for TOFA 10 mg BID in Feb 2019, patients on this dose were changed to 5 mg BID
- The study was completed once ≥1500 patients were followed for ≥3 years, ≥103 MACE were reported, and ≥138 malignancies excluding NMSC were reported

CVD = cardiovascular disease; ETN = etanercept; QW = each week; RoW = rest of world.
Ytterberg SR, et al. ACR 2021, abstract 831

ORAL Surveillance Safety Trial: Tofacitinib vs TNF Inhibitors

HRs (95% CI) and IRs (95% CI) for adjudicated MACE and malignancies

◆ TOFA 5 mg BID ◆ TOFA 10 mg BID ◆ Combined TOFA doses ◆ TNFi

MACE

HR = 1.24 (0.81–1.91)

HR = 1.43 (0.94–2.18)

HR = 1.15 (0.77–1.71)

HR = 1.33 (0.91–1.94)

HR = 0.98

HR = 0.73

Malignancies (excluding NMSC)

HR = 1.47 (1.00–2.18)

HR = 1.48 (1.00–2.19)

HR = 1.00 (0.70–1.43)

HR = 1.48 (1.04–2.09)

HR = 1.13

HR = 1.13

HR = 1.13

HR = 0.77

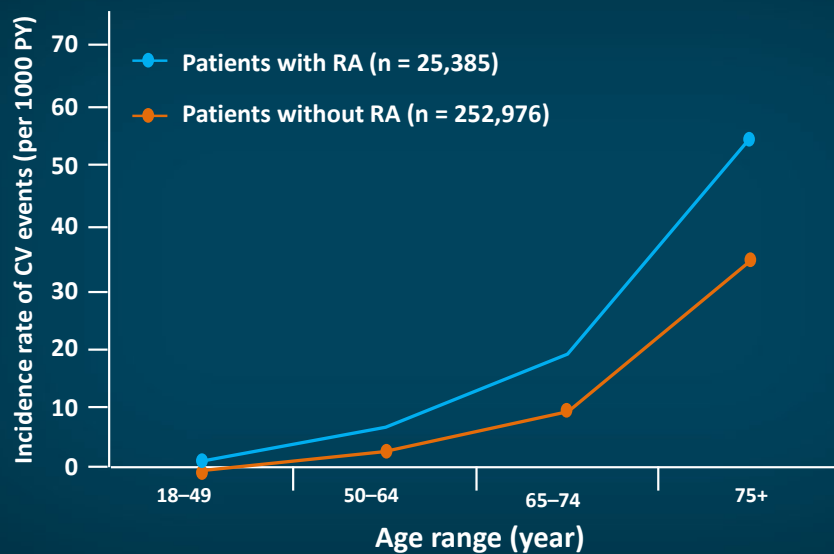
IR (95% CI)

	◆ TOFA 5 mg BID	◆ TOFA 10 mg BID	◆ Combined TOFA doses	◆ TNFi
n	47	51	98	37
N	1455	1456	2911	1451
PY	5166	4872	10,038	5045
IR (95% CI)	0.91	1.05	0.98	0.73
Noninferiority margin: upper CI	<1.8			
NNH vs TNFi	567	319	412	ref
IR (95% CI)	1.13	1.13	1.13	0.77
NNH vs TNFi	276	275	275	ref

- Combined tofacitinib doses were not non-inferior to TNFi for either MACE or malignancy
- No difference between BID tofacitinib doses
- Most MACE and malignancies occurred in patients >65 years and/or ever smokers
- MACE difference for tofacitinib vs TNFi primarily for MI
- Malignancy difference for tofacitinib vs TNFi primarily for lung cancer, lymphoma, and NMSC

IR = incidence rate; NNH = number needed to harm; MI = myocardial infarction.
Ytterberg SR, et al. ACR 2021, abstract 831. Charles-Schoeman C, et al. ACR 2021, abstract 958. Curtis J, et al. ACR 2021, abstract 1940.

RA Is an Independent Risk Factor for Cardiovascular Events*



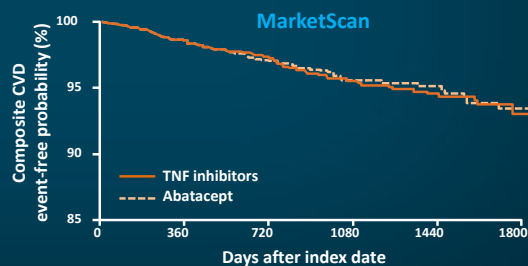
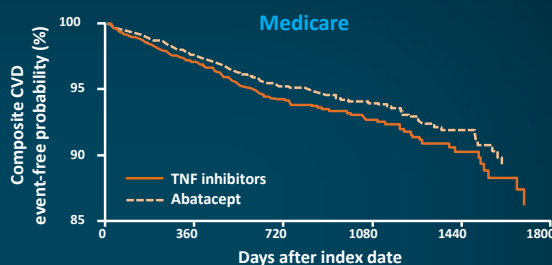
*MI and stroke.

CV = cardiovascular.

Solomon DH, et al. *Ann Rheum Dis.* 2006;65:1608-1612.

DMARDs and CVD Events in RA

- Methotrexate and TNF inhibitors are most studied
- Observational data only
- Comparison group problematic
- MTX use associated with 28% reduction in all CVD events across 8 cohort studies (HR = 0.72 [95% CI, 0.57–0.91], $P = .007$)¹
- TNFi use associated with 30% reduction in CVD events across 16 cohort studies (HR = 0.70 [95% CI, 0.54–0.90], $P = .005$)¹
- Abatacept similar to etanercept from claims data²
- Tocilizumab similar to etanercept in head-to-head randomized clinical trial³



1. Roubille C, et al. *Ann Rheum Dis.* 2015;74:480-489. 2. Kang EH, et al. *J Am Heart Assoc.* 2018;7:e007393. 3. Giles JT, et al. *Arthritis Rheumatol.* 2020;72:31-40.

RA Treatments and Comorbid Disease Considerations from the ACR Treatment Guideline

RA treatments and comorbid disease considerations	
Recommendations	Certainty of evidence
Previous serious infection Addition of csDMARDs is conditionally recommended over addition of a bDMARD or tsDMARD for patients with a serious infection within previous 12 months who have moderate-to-high disease activity despite csDMARD monotherapy Addition of/switching to DMARDs is conditionally recommended over initiation/dose escalation of glucocorticoids for patients with a serious infection within previous 12 months who have moderate-to-high disease activity	Very low Very low
Pulmonary disease Methotrexate is conditionally recommended over alternative DMARDs for treatment of inflammatory arthritis for patients with clinically diagnosed mild and stable airway or parenchymal lung disease who have moderate-to-high disease activity	Very low
Heart failure Addition of a non-TNFi DMARD or tsDMARD is conditionally recommended over addition of a TNFi for patients with New York Heart Association (NYHA) class III or IV heart failure and an inadequate response to csDMARDs Switching to a non-TNFi bDMARD or tsDMARD is conditionally recommended over continuation of a TNFi for patients taking a TNFi who develop heart failure	Very low Very low

Fraenkel L, et al. *Arthritis Care Res.* 2021;73:924-939.

RA Treatments and Comorbid Disease Considerations from the ACR Treatment Guideline (continued)

RA treatments and comorbid disease considerations	
Recommendations	Certainty of evidence
Hepatitis B infection Prophylactic antiviral therapy is strongly recommended over frequent monitoring alone for patients initiating rituximab who are hepatitis B core antibody positive (regardless of hepatitis B surface-antigen status) Prophylactic antiviral therapy is strongly recommended over frequent monitoring alone for patients initiating any bDMARD or tsDMARD who are hepatitis B core antibody positive and hepatitis B surface antigen positive Frequent monitoring alone is conditionally recommended over prophylactic antiviral therapy for patients initiating a bDMARD other than rituximab or a tsDMARD who are hepatitis B core antibody positive and hepatitis B surface antigen negative	Very low Very low Very low
Lymphoproliferative disorder Rituximab is conditionally recommended over other DMARDs for patients who have a previous lymphoproliferative disorder for which rituximab is an approved treatment and who have moderate-to-high disease activity	Very low

Fraenkel L, et al. *Arthritis Care Res.* 2021;73:924-939.

Conclusions

- RA is a chronic inflammatory disease with significant morbidity and disability if left untreated
- Advances in our understanding of RA pathogenesis have led to the development of new and more effective therapies; most recently, these have included newly developed agents that block JAK signaling
- Optimal approach to therapy needs to consider comorbidities, quality of life, T2T, tailoring regimens when needed, and promoting strategies that improve adherence

Case Study: New Mother

Case Study: Patient History

- Tracey is a 35-year-old woman who gave birth to her second child 4 months ago.
- Not long after the delivery, she began noticing morning stiffness in her hands and wrists, and she felt like she was walking “on rocks” when she took her first steps out of bed in the morning.
- She attributed the symptoms to the fatigue and lack of sleep that accompany caring for an infant, but the symptoms worsened over time.
- Her primary care doctor evaluated her and ordered some lab tests after observing some joint swelling in her fingers and wrists.
- She is seropositive for anti-CCP and RF, and her inflammatory markers are moderately elevated.
- She is referred to a rheumatologist for her first evaluation.

Case Study: Question #1

What are some of the challenges for treating Tracey?

Case Study: Considerations

- Patients like Tracey are not uncommon since RA often presents in women of childbearing age. The immediate post-partum period is a common time at which RA initially presents.
- In initiating therapy, the physician needs to consider treatments that are compatible with breastfeeding.
- Additional consideration should be made for whether Tracey is thinking of having more children and her timeframe for having them.

Case Study: Question #2

- Tracey is ready to stop breastfeeding now but wants to have another child in the next 6 to 12 months.
- What initial RA treatment options are appropriate for Tracey today?
 - A. Start methotrexate, as recommended as first-line therapy in the ACR guidelines
 - B. Hold off on any treatment until Tracey has her next child
 - C. Treat with prednisone only during this time
 - D. Start a TNF inhibitor
 - E. Start a JAK inhibitor

Case Study: Treatment Considerations

- Although methotrexate is typically used first-line in RA, it is class X for conception, pregnancy, and breastfeeding. It must be discontinued at least 3 months before attempting conception, which does not fit well into Tracey's timeline.
- Not treating Tracey's RA until after she delivers her next child is not ideal, since you do not know how long it will take for her to conceive. If there is a long period before conception, she would have joint symptoms and potentially experience joint damage.
- The same is true for prednisone use, which is associated with multiple side effects with prolonged use.
- TNF inhibitor therapy is an option, since it can be used through conception. Most rheumatologists will stop therapy when pregnancy is confirmed since RA tends to abate during pregnancy.
- Little to no data are available about the effect of JAK inhibitors on pregnancy outcomes or their safety during breastfeeding.

Case Study: Tracey

Case Study: Birth of Third Child

- Tracey started taking a TNF inhibitor after the birth of her second child to control her newly diagnosed RA. She has had a very favorable response.
- After 10 months of treatment, she became pregnant with her third child and stopped the TNF inhibitor. Her RA was in remission during pregnancy, and she gave birth to a healthy baby girl after an uneventful pregnancy.
- Soon after the birth, Tracey's RA returned with more joints involved than originally. In addition to the fingers, wrists, and toes, she now has swollen ankles and knees, and her left elbow is so stiff and swollen that she cannot extend it.
- When her rheumatologist restarted the TNF inhibitor that she was taking before her pregnancy, it did not seem to work at all this time.
- She does not plan to have any more children.

Case Study: Question #1

**What are Tracey's
treatment options?**

Case Study: Considerations

- As before, the post-partum period has revved up her RA disease activity. Now the symptoms are more severe than previously, and they interfere with her ability to function and care for her family.
- Methotrexate is an appropriate option since she has no immediate pregnancy plans and has no contraindications.
- Tracey starts with a low dose of prednisone. She feels better initially, but once the prednisone is tapered, she still has significant joint swelling and stiffness, even after prednisone has been increased to 25 mg per week.
- In addition, Tracey reports that the day or two after she takes methotrexate, she feels exhausted and has headaches. She tells you that she cannot continue taking a medication that causes her to be out of commission for 2 days out of the week.

Case Study: Question #2

What change would you make to Tracey's treatment regimen?

- A. Change her current oral methotrexate to subcutaneous injected methotrexate at the same dose
- B. Restart prednisone and continue it indefinitely
- C. Start a different TNF inhibitor than the one she was taking before her second pregnancy
- D. Start a non-TNF inhibitor biologic
- E. Start a JAK inhibitor

Case Study: Treatment Considerations

- Changing oral MTX to subcutaneous form can help with some symptoms, such as gastrointestinal intolerance, but it usually does not help with fatigue and headache. In fact, it can make those symptoms worse.
- Long-term corticosteroid therapy carries significant risk of toxicity and is not likely to help with the MTX side effects very much.
- Tracey responded to a TNF inhibitor before but failed to respond when it was restarted after her second pregnancy. She might respond to a second TNF inhibitor, but there is no guarantee.
- A non-TNF inhibitor biologic is an option, but each of them requires 3 months or more to reach full effectiveness, and most work better in combination with MTX or an additional non-biologic DMARD.
- The JAK inhibitor class is effective with or without concomitant non-biologic DMARDs, and responses are typically observed in weeks rather than months.

Thank You!

Q&A

**Disease Modification in Individuals with Moderate-to-Severe Rheumatoid Arthritis:
Optimizing Treatment Through the Finely Tuned Selectivity for JAKs**

Resource	Address
Burmester G, Cohen S, Winthrop K, et al. Long-term safety profile of upadacitinib in patients with rheumatoid arthritis, psoriatic arthritis, or ankylosing spondylitis. Presented at: ACR Convergence; November 9, 2021; Abstract 1691.	https://acrabstracts.org/abstract/long-term-safety-profile-of-upadacitinib-in-patients-with-rheumatoid-arthritis-psoriatic-arthritis-or-ankylosing-spondylitis/
Burmester GR, Blanco R, Charles-Schoeman C, et al. Tofacitinib (CP-690,550) in combination with methotrexate in patients with active rheumatoid arthritis with an inadequate response to tumour necrosis factor inhibitors: A randomised phase 3 trial. <i>Lancet.</i> 2013;381:451-460.	https://pubmed.ncbi.nlm.nih.gov/23294500/
Desai RJ, Pawar A, Weinblatt ME, Kim SC. Comparative risk of venous thromboembolism in rheumatoid arthritis patients receiving tofacitinib versus those receiving tumor necrosis factor inhibitors: An observational cohort study. <i>Arthritis Rheumatol.</i> 2019;71:892-900.	https://pubmed.ncbi.nlm.nih.gov/30552833/
Fleischmann R, Mysler E, Hall S, et al. Efficacy and safety of tofacitinib monotherapy, tofacitinib with methotrexate, and adalimumab with methotrexate in patients with rheumatoid arthritis (ORAL Strategy): A phase 3b/4, double-blind, head-to-head, randomised controlled trial. <i>Lancet.</i> 2017;390:457-468.	https://pubmed.ncbi.nlm.nih.gov/28629665/
Fraenkel L, Bathon JM, England BR, et al. 2021 American College of Rheumatology Guideline for the Treatment of Rheumatoid Arthritis. <i>Arthritis Care Res (Hoboken).</i> 2021;73:924-939.	https://pubmed.ncbi.nlm.nih.gov/34101387/
Fragoulis GE, Brock J, Basu N, McInnes IB, Siebert. The role for JAK inhibitors in the treatment of immune-mediated rheumatic and related conditions. <i>J Allergy Clin Immunol.</i> 2021;148:941-952.	https://pubmed.ncbi.nlm.nih.gov/34450118/
Genovese MC, Fleischmann R, Combe B, et al. Safety and efficacy of upadacitinib in patients with active rheumatoid arthritis refractory to biologic disease-modifying anti-rheumatic drugs (SELECT-BEYOND): A double-blind, randomised controlled phase 3 trial. <i>Lancet.</i> 2018;391:2513-2524.	https://pubmed.ncbi.nlm.nih.gov/29908670/
Genovese MC, Kremer J, Zamani O, et al. Baricitinib in patients with refractory rheumatoid arthritis. <i>N Engl J Med.</i> 2016;374:1243-1252.	https://pubmed.ncbi.nlm.nih.gov/27028914/

<p>Köhler BM, Günther J, Kaudewitz D, Lorenz HM. Current Therapeutic Options in the Treatment of Rheumatoid Arthritis. <i>J Clin Med.</i> 2019;8:938.</p>	<p>https://pubmed.ncbi.nlm.nih.gov/31261785/</p>
<p>Rubbert-Roth A, Enejosa J, Pangan AL, et al. Trial of upadacitinib or abatacept in rheumatoid arthritis. <i>N Engl J Med.</i> 2020;383:1511-1521.</p>	<p>https://pubmed.ncbi.nlm.nih.gov/33053283/</p>
<p>Smolen JS, Aletaha D, Bijlsma JW, et al. Treating rheumatoid arthritis to target: Recommendations of an international task force. <i>Ann Rheum Dis.</i> 2010;69:631-637.</p>	<p>https://pubmed.ncbi.nlm.nih.gov/20215140/</p>
<p>Ytterberg SR, Bhatt DL, Mikuls T, et al. Safety and efficacy of tofacitinib vs TNF inhibitors in RA patients aged 50 years or older with one or more cardiovascular risks: Results from a phase 3b/4 randomized safety trial. Presented at: ACR Convergence; November 7, 2021; Abstract 831.</p>	<p>https://acrabstracts.org/abstract/safety-and-efficacy-of-tofacitinib-vs-tnf-inhibitors-in-ra-patients-aged-50-years-or-older-with-one-or-more-cardiovascular-risks-results-from-a-phase-3b-4-randomized-safety-trial/</p>