How do I effectively help persons with Rheumatoid Arthritis?

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Disclosures

Non-Declaration Statement: I have no relevant relationships with ineligible companies to disclose within the past 24 months. (Note: Ineligible companies are defined as those whose primary business is producing, marketing, selling, re-selling, or distributing healthcare products used by or on patients.)

Objectives

After completing this session, attendees will be able to:

- **utilize the latest diagnostic approaches when evaluating persons with rheumatoid arthritis**
- **identify the currently approved medications for rheumatoid arthritis.**
- **describe the risks, benefits and expectations of biologics in treating rheumatoid arthritis.**

In the arthritis which generally shows itself about the age of thirty-five there is frequently no great interval between the affection of the hands and feet; both these becoming similar in nature, slender, with little flesh…For the most part their arthritis passeth from the feet to the hands, next the elbows and knees, after these the hip joint. It is incredible how fast the mischief spreads.

Hippocrates

Copeman WSC. A Short History of Gout. Berkeley and Los Angeles: University of California Press; 1964.

…[when some] undifferentiated morbid condition is first described, the characters of which are so striking that it seems well-nigh impossible that they should have been long overlooked it is often suggested that the malady is one of recent development, a new disease which owes its origin to some alteration in the conditions of life…in the case of the disease now to be considered, there is no room for suggestions of this kind, for the evidence of its antiquity is derived, not from mere written descriptions, but from the impress which it has left upon the bones of its victims…

Archibald Garrod

Treatise on Rheumatism and Rheumatoid Arthritis (1890)

Garrod AE. A Treatise on Rheumatism and Rheumatoid Arthritis. London: Charles Griffin and Company; 1890.

Case

37 y.o. ♀ nurse with 6-month hx of RF/CCP+ RA on MTX 20mg po q week X 3 months with breakthrough sxs including arthralgia and synovitis. Has erosions on hands/wrists and feet x-rays.

PMH---s/p TAH/BSO

+ PPD X 5yrs, -CXR, no prior treatment

No other meds except MTX/Folic Acid

WHAT DIAGNOSIS AM I TO CONSIDER FOR THIS PERSON WITH JOINT PAIN?

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THE AMERICAN RHEUMATISM ASSOCIATION 1987 REVISED CRITERIA FOR THE CLASSIFICATION OF RHEUMATOID ARTHRITIS

FRANK C. ARNETT, STEVEN M. EDWORTHY, DANIEL A. BLOCH, DENNIS J. McSHANE, JAMES F. FRIES, NORMAN S. COOPER, LOUIS A. HEALEY, STEPHEN R. KAPLAN, MATTHEW H. LIANG, HARVINDER S. LUTHRA, THOMAS A. MEDSGER, Jr., DONALD M. MITCHELL, DAVID H. NEUSTADT, ROBERT S. PINALS, JANE G. SCHALLER, JOHN T. SHARP, RONALD L. WILDER, and GENE G. HUNDER

*Morning stiffness lasting at least 1 hour
*Arthritis of three or more joint areas
*Arthritis of hand joints
*Symmetric arthritis
*Rheumatoid nodules
*Serum rheumatoid factor
*Radiographic changes

4/7 criteria present (first four listed for at least 6 weeks)

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Arthritis & Rheumatism

An Official Journal of the American College of Rheumatology
www.arthritisrheum.org and www.interscience.wiley.com

2010 Rheumatoid Arthritis Classification Criteria

An American College of Rheumatology/European League Against Rheumatism Collaborative Initiative

Daniel Aletaha,[1] Tuhina Neogi,[2] Alan J. Silman,[3] Julia Funovits,[1] David T. Felson,[2] Clifton O. Bingham, III,[4] Neal S. Birnbaum,[5] Gerd R. Burmester,[6] Vivian P. Bykerk,[7] Marc D. Cohen,[8] Bernard Combe,[9] Karen H. Costenbader,[10] Maxime Dougados,[11] Paul Emery,[12] Gianfranco Ferraccioli,[13] Johanna M. W. Hazes,[14] Kathryn Hobbs,[15] Tom W. J. Huizinga,[16] Arthur Kavanaugh,[17] Jonathan Kay,[18] Tore K. Kvien,[19] Timothy Laing,[20] Philip Mease,[21] Henri A. Ménard,[22] Larry W. Moreland,[23] Raymond L. Naden,[24] Theodore Pincus,[25] Josef S. Smolen,[1] Ewa Stanislawska-Biernat,[26] Deborah Symmons,[27] Paul P. Tak,[28] Katherine S. Upchurch,[18] Jiří Vencovský,[29] Frederick Wolfe,[30] and Gillian Hawker[31]

Criteria

2010 Rheumatoid arthritis classification criteria: an American College of Rheumatology/European League Against Rheumatism collaborative initiative

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Target Population of the Criteria

Two requirements:

(1) Patient with at least one joint with definite clinical synovitis (swelling)

(2) Synovitis is not better explained by "another disease"

Differential diagnoses differ in patients with different presentations.

If unclear about the relevant differentials, an expert rheumatologist should be consulted.

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2010 ACR/EULAR Classification Criteria for RA

JOINT DISTRIBUTION (0-5)	
1 large joint	0
2-10 large joints	1
1-3 small joints (large joints not counted)	2
4-10 small joints (large joints not counted)	3
>10 joints (at least one small joint)	5
SEROLOGY (0-3)	
Negative RF AND negative ACPA	0
Low positive RF OR low positive ACPA	2
High positive RF OR high positive ACPA	3
SYMPTOM DURATION (0-1)	
<6 weeks	0
≥6 weeks	1
ACUTE PHASE REACTANTS (0-1)	
Normal CRP AND normal ESR	0
Abnormal CRP OR abnormal ESR	1

≥6 = definite RA

What if the score is <6?

Patient might fulfill the criteria…

→ **Prospectively** over time (cumulatively)

→ **Retrospectively** if data on all four domains have been adequately recorded in the past

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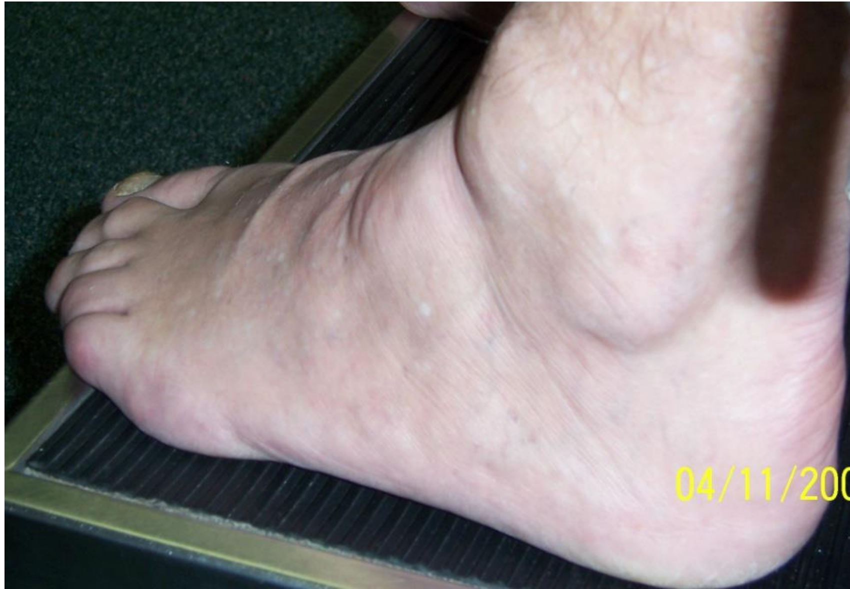
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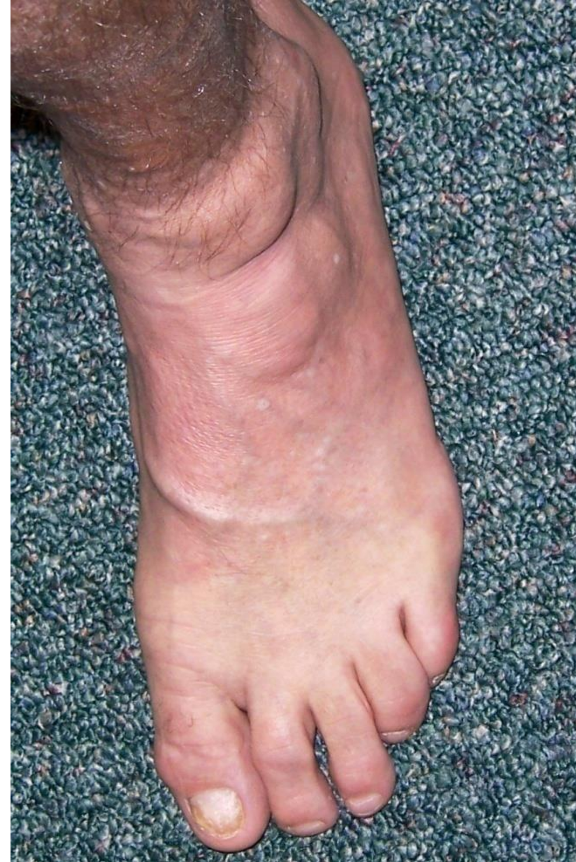

EXTRA-ARTICULAR MANIFESTATIONS OF RHEUMATOID ARTHRITIS

Skin	Nodules, fragility, vasculitis, pyoderma gangrenosum
Heart	Pericarditis, premature atherosclerosis, vasculitis, valve disease, and valve ring nodules
Lung	Pleural effusions, interstitial lung disease, bronchiolitis obliterans, rheumatoid nodules, vasculitis
Eye	Keratoconjunctivitis sicca, episcleritis, scleritis, scleromalacia perforans, peripheral ulcerative keratopathy
Neurologic	Entrapment neuropathy, cervical myelopathy, mononeuritis multiplex (vasculitis), peripheral neuropathy
Hematopoietic	Anemia, thrombocytosis, lymphadenopathy, Felty's syndrome
Kidney	Amyloidosis, vasculitis
Bone	Osteopenia

RA, splenomegaly, and neutropenia. This complication is seen in patients with severe, RF/ACPA-positive disease and may be accompanied by hepatomegaly, thrombocytopenia, lymphadenopathy, and fevers

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Rheumatoid Nodules

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Rheumatoid Factor

- Autoantibodies directed against Fc portion of IgG (IgM to IgG)
- 75-90% of RA patients
- Result can aid in the diagnosis, but is <u>not</u> diagnostic of RA
- RF <u>not</u> used to measure RA disease activity, but higher titers can be associated with disease severity, erosions, extra-articular manifestations, disability

Rheumatoid Factor in other diseases

CH-Chronic disease
- *hepatic (PBC)
- *pulmonary (IPF, silicosis, asbestosis)

R-Rheumatoid Arthritis

O-Other rheumatic disease
- *SLE
- *Systemic sclerosis
- *MCTD
- *Sjögren’s
- *Polymyositis
- *Sarcoid

N-Neoplasm, especially after XRT or chemo

I-Infections
- *AIDS
- *Mononucleosis
- *Parasitic infections
- *Chronic Viral
- *Hepatitis B/C
- *Chronic bacterial (SBE, syphilis, mycobacteria)

C-Cryoglobulienmia (esp with Hep C)

Anti-Cyclic Citrullinated Peptide Antibodies (anti-CCP)

- RA sensitivity 47-76%
  specificity 90-96%

- Can occur in active TB, SLE, Sjogren's, Polymyositis, Dermatomyositis, Scleroderma

- (+) CCP Ab

  more likely to have aggressive disease and progressive radiographic joint damage

Radiographic Studies

X-rays

Ultrasound

Magnetic Resonance Imaging

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X-ray Changes in RA

WHAT TREATMENTS MIGHT I CONSIDER FOR THIS PERSON?

Goals of RA treatment

-Relieve pain

-Reduce inflammation

-Protect articular structures

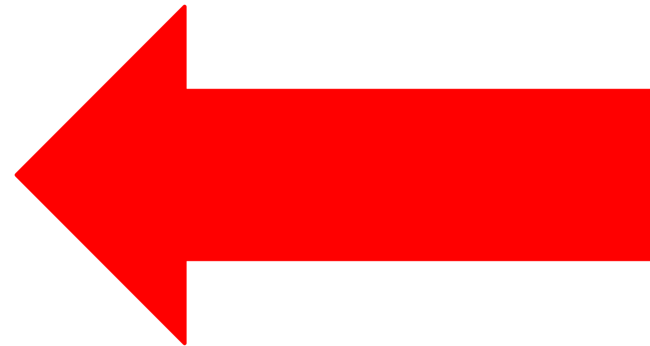
-Maintain function

-Control systemic involvement

Ruderman, Eric M., MD, Editorial Consultant. Management of Rheumatic Diseases in the Biologic Era. Coalition of Rheumatology Educators; 2008

Pharmacologic Therapy

- Nonsteroidal Anti-inflammatory Drugs
- Corticosteroids
- Hydroxychloroquine
- Sulfasalazine
- Methotrexate
- Leflunomide
- Azathioprine
- Cyclosporine



Disease modifying antirheumatic drugs (DMARDs)

Guidelines for use of glucocorticoids in RA

-Avoid use of glucocorticoids without DMARDs

-Prednisone, >10 mg/day, is rarely indicated for articular disease

-Taper to the lowest effective dose

-Use as “bridge therapy” until DMARD therapy is effective

-Remember prophylaxis against osteoporosis

WHAT ABOUT THOSE NEW TREATMENTS THAT I SEE ON TV?

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- Don’t test ANA sub-serologies without a positive ANA and clinical suspicion of immune-mediated disease.
- Don’t test for Lyme disease as a cause of musculoskeletal symptoms without an exposure history and appropriate exam findings.
- Don’t perform MRI of the peripheral joints to routinely monitor inflammatory arthritis.
- Don’t prescribe biologics for rheumatoid arthritis before a trial of methotrexate (or other conventional non-biologic DMARDs).
- Don’t routinely repeat DXA scans more often than once every two years.

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Biologics/Small Molecules

-Tumor Necrosis Factor-ά antagonists

Adalimumab (Humira)-SQ

Certolizumab (Cimzia)-SQ

Etanercept (Enbrel)-SQ

Golimumab (Simponi, Simponi Aria)-SQ, IV

Infliximab (Remicade)-IV

-Interleukin-1 receptor antagonist (IL-1)

Anakinra (Kineret)-SQ

-B cells

Rituximab (Rituxan)-IV

-T cells

Abatacept (Orencia)-SQ, IV

-Interleukin-6 receptor (IL-6R)

Tocilizumab (Actemra) SQ, IV

Sarilumab (Kevzara)-SQ

-Janus Kinase (JAK) inhibitor

Tofacitinib (Xeljanz)-PO

Baricitinib (Olumiant)-PO

Upadacitinib (Rinvoq)-PO

My Pre-Drug Questions

-Current/recurrent infxns

-Cancer (CA)

-Congestive Heart Failure (CHF)

-Chronic Obstructive Pulmonary Disease (COPD)/asthma

-Tuberculosis (TB)

  *PPD hx

  *exposure

-Multiple Sclerosis (MS)

-Hepatitis B/C

-Hyperlipidemia

Biologics/Small Molecules

Potential risks

- **Injection site/infusion reaction**
- **Infection risk (bacterial, TB/other granulomatous, opportunistic)**
- **Malignancy risk ?**
- **Demyelinating Disease, MS or Family Hx**
- **Heart failure**
- **Drug induced syndromes (ANA, dsDNA)**
- **Cytopenias**
- **Gastrointestinal perforation**

Biologics/Small Molecules

<u>Pre-drug screening</u>

-CXR

-PPD/Interferon-gamma release assays (IGRAs)

-Pneumonia vaccine

-Influenza vaccine

-Hepatitis B and C serolog

COVID-19 vaccination in Rheumatic and Musculoskeletal Disease Patients

General considerations

- Engage in shared decision-making regarding vaccination
- Recognize heterogeneity of COVID-19 and higher risk for hospitalized COVID-19, worse outcomes compared to the general population
- Should be prioritized for vaccination before the nonprioritized general population of similar age and sex
- No additional contraindications to vaccinations
- Vaccination response blunted in its magnitude and duration for those on immunomodulatory therapies compared to general population
- Theoretic risk of disease flare or worsening, but vaccine benefit outweighs risk

https://www.rheumatology.org/Portals/0/Files/COVID-19-Vaccine-Clinical-Guidance-Rheumatic-Diseases-Summary.pdf

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SPECIAL ARTICLE

2015 American College of Rheumatology Guideline for the Treatment of Rheumatoid Arthritis

JASVINDER A. SINGH,[1] KENNETH G. SAAG,[1] S. LOUIS BRIDGES JR.,[1] ELIE A. AKL,[2] RAVEENDHARA R. BANNURU,[3] MATTHEW C. SULLIVAN,[3] ELIZAVETA VAYSBROT,[3] CHRISTINE MCNAUGHTON,[3] MIKALA OSANI,[3] ROBERT H. SHMERLING,[4] JEFFREY R. CURTIS,[1] DANIEL E. FURST,[5] DEBORAH PARKS,[6] ARTHUR KAVANAUGH,[7] JAMES O'DELL,[8] CHARLES KING,[9] AMYE LEONG,[10] ERIC L. MATTESON,[11] JOHN T. SCHOUSBOE,[12] BARBARA DREVLOW,[13] SETH GINSBERG,[14] JAMES GROBER,[13] E. WILLIAM ST.CLAIR,[15] ELIZABETH TINDALL,[16] AMY S. MILLER,[17] AND TIMOTHY MCALINDON[3]

Recommendations for the Use of Vaccines in RA patients on DMARD and/or biologic therapy biologic therapy

-In early or established RA patients aged 50 and over, we conditionally recommend giving the herpes zoster vaccine before the patient receives biologic therapy or tofacitinib for their RA.

-In early or established RA patients who are currently receiving biologics, we conditionally recommend that live attenuated vaccines such as the herpes zoster (shingles) vaccine not be given.

-In patients with early or established RA who are currently receiving biologics, we strongly recommend using appropriately indicated killed/inactivated vaccines

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2022 American College of Rheumatology (ACR) Guideline for Exercise, Rehabilitation, Diet, and Additional Integrative Interventions for Rheumatoid Arthritis

- First ACR guideline for integrative interventions for RA management
- Strong recommendation for consistent exercise
- Conditional recommendations for specific exercise types, comprehensive PT/OT, several rehabilitation modalities, Mediterranean-style diet, and several additional integrative interventions
- Interprofessional teams are critical for guiding patients through their disease course
- Access to, and burden of, these interventions expected to impact patient engagement and/or adoption
- Conditional nature of most recommendations requires engaging patients in shared decision making and highlights the need for additional research

When to Refer

Uncertain diagnosis

Confusing Lab Results

Uncomfortable with DMARDS or Biologic Use

Patient not responding

Erosions or other radiographic changes

Side effects

Lessons for Practice

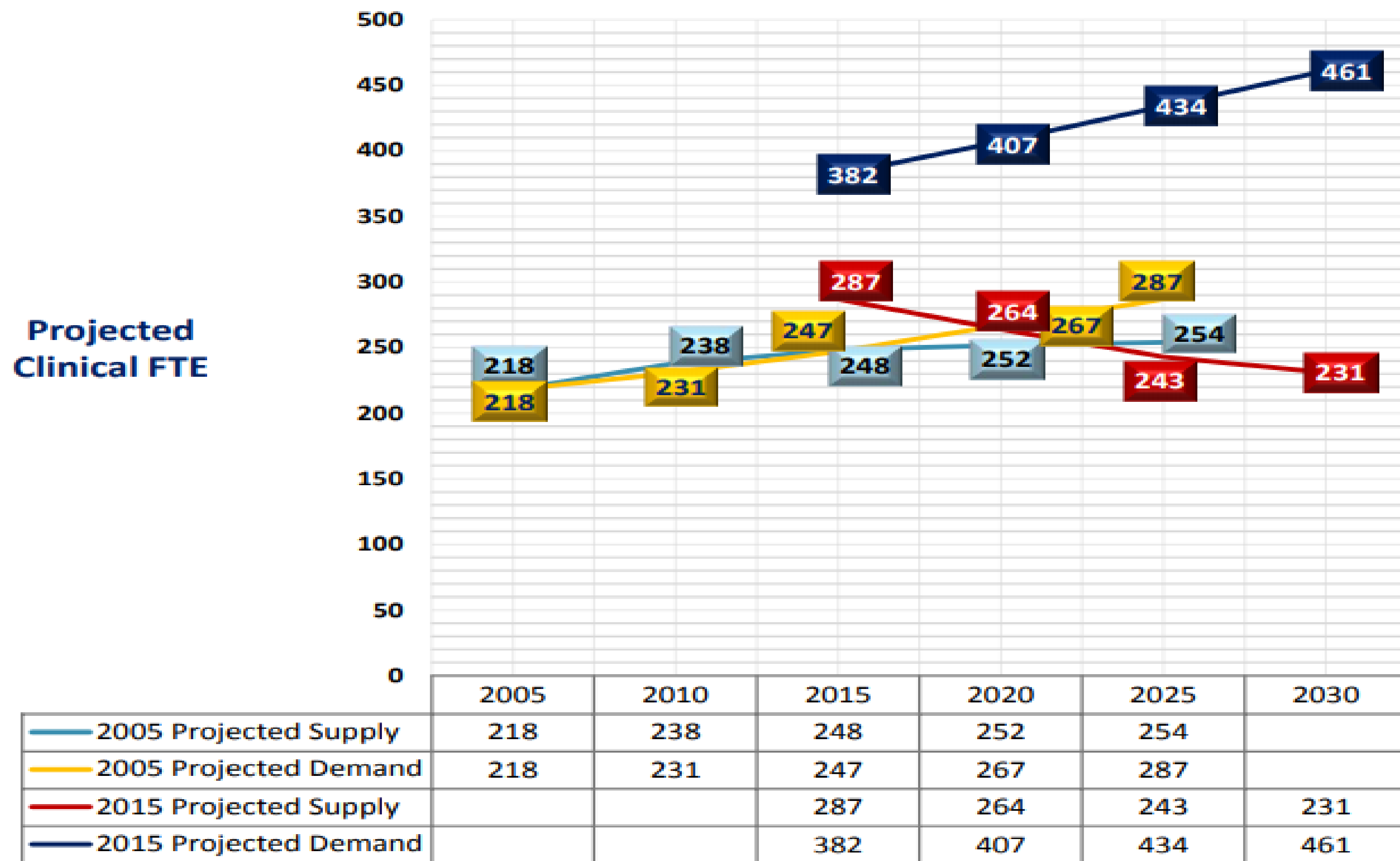
-Rheumatoid Arthritis is a systemic, inflammatory condition. Generally, early diagnosis lends itself to a better prognosis.

-There are multiple pharmacologic treatment options to care for those with rheumatoid arthritis.

-Biologic DMARDS require a thorough prescreening process and appropriate, ongoing monitoring while taking these powerful medications.

Figure 2. Comparison of Projected Supply and Demand of Adult Rheumatology Workforce

2015 Workforce Study of Rheumatology Specialists in the United States: Final Report
http://www.rheumatology.org/portals/0/files/ACR-Workforce-Study-2015.pdf



	2005	2010	2015	2020	2025	2030
2005 Projected Supply	218	238	248	252	254	
2005 Projected Demand	218	231	247	267	287	
2015 Projected Supply			287	264	243	231
2015 Projected Demand			382	407	434	461

Figure E-7. Comparison of Projected Supply and Projected Demand of Pediatric Rheumatologists

Note. Data from 2005 workforce study (2005 to 2025); Data from the 2015 workforce study (2015 to 2030).

2015 Workforce Study of Rheumatology Specialists in the United States: Final Report
(http://www.rheumatology.org/portals/0/files/ACR-Workforce-Study-2015.pdf)

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