

Rheumatologic Updates for the Pain Physician



TIRR
MEMORIAL
HERMANN
Rehabilitation & Research

 **UTHealth**
The University of Texas
Health Science Center at Houston

McGovern
Medical School

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Texas Pain Society | Saturday, October 25, 2025. Houston, TX

Disclosures

☐ None

Learning Objectives

- Review epidemiology of rheumatologic disease and how it contributes to disability.
- Review pathogenesis of pain in rheumatologic diseases and how it contributes to chronic pain.
- Discuss how to obtain relevant clinical history and assessment tools.
- Highlight specific benefits and limitations of pharmacologic and interventional pain management therapies.
- Describe best practices for multimodal treatment based on evidenced-based ACR guidelines including:
 - ? Analgesics
 - ? Targeted interventions
 - ? Physical therapy
 - ? Psychology

Learning Objectives

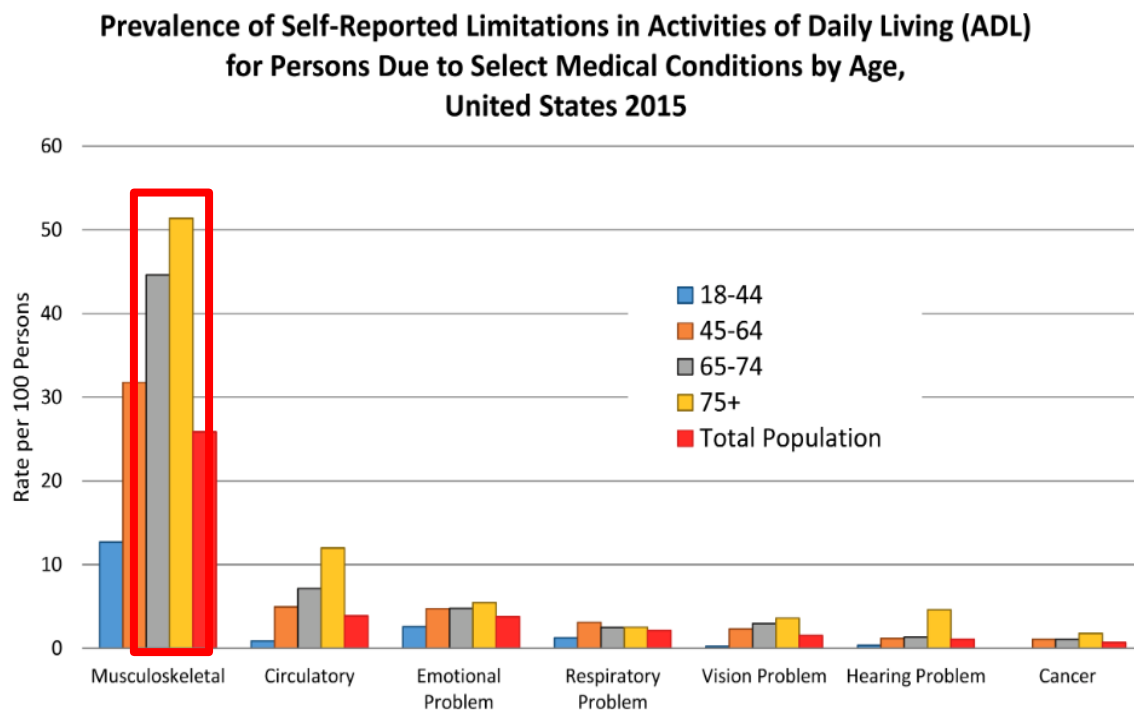
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Epidemiology: Arthritis

Rheumatologic Disease	Prevalance
Osteoarthritis	302 million worldwide. 33 million US adults
Gout	55 million worldwide 9 million Americans (4%) of adults
Rheumatoid Arthritis	18 million worldwide 1.36 million adults in the US
Lupus	3.4 million worldwide 1.5 million Americans
Psoriatic arthritis	3% global population

Despite biologic therapies, pain is often uncontrolled.
2/3 of patients with rheumatologic disease have daily pain.

Disability due to Arthritis



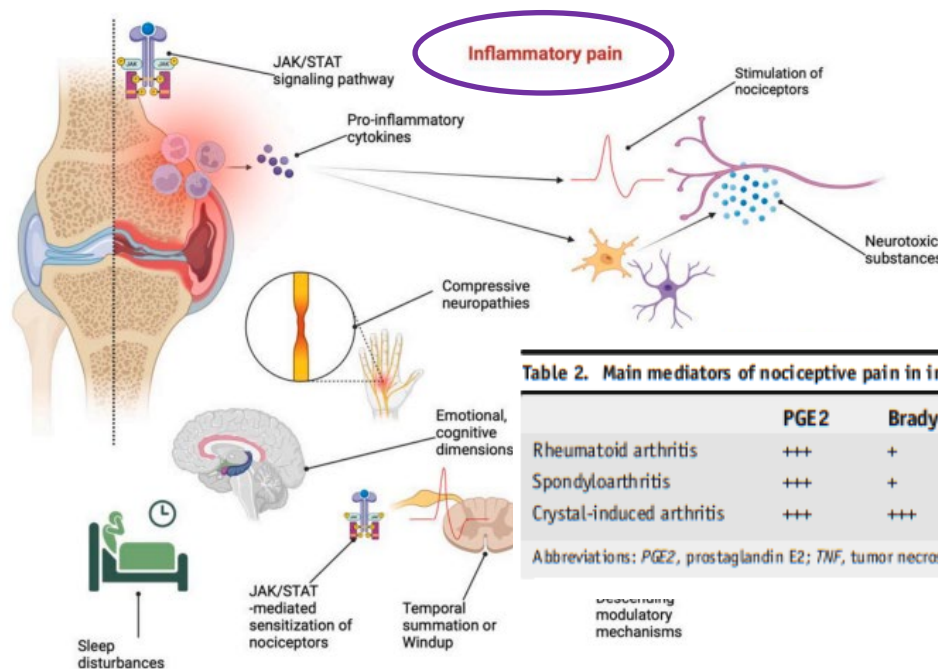
Musculoskeletal Conditions account for 34.1% of Social Security Disability (Dec 2023) contributing \$500-635 billion dollars spent annually

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Pain in Inflammatory Arthritis

“An unpleasant sensory and emotional experience associated with actual or potential tissue damage, or described as result of such damage” - *International Association for the Study of Pain*



Nociceptive
damage to non-neural tissue → (+) nociceptor

Table 2. Main mediators of nociceptive pain in inflammatory arthritis

	PGE2	Bradykinin	Substance P	TNF α	IL-1 β	IL-6	IL-17	ACPA	NGF
Rheumatoid arthritis	+++	+	+	+++	+++	+++	+	++	+
Spondyloarthritis	+++	+	+	+++	-	-	+++	-	+
Crystal-induced arthritis	+++	+++	+	-	+++	-	-	-	-

Abbreviations: PGE2, prostaglandin E2; TNF, tumor necrosis factor; ACPA, anti-citrullinated protein antibodies; IL, interleukin; NGF, nerve growth factor

Neurotrophic
modulatory
mechanisms

Neuropathic Pain in IA

Clin Rheumatol (2008) 27:841–844
DOI 10.1007/s10067-007-0804-x

ORIGINAL ARTICLE

A clinical, electrophysiological, and pathological study of neuropathy in rheumatoid arthritis

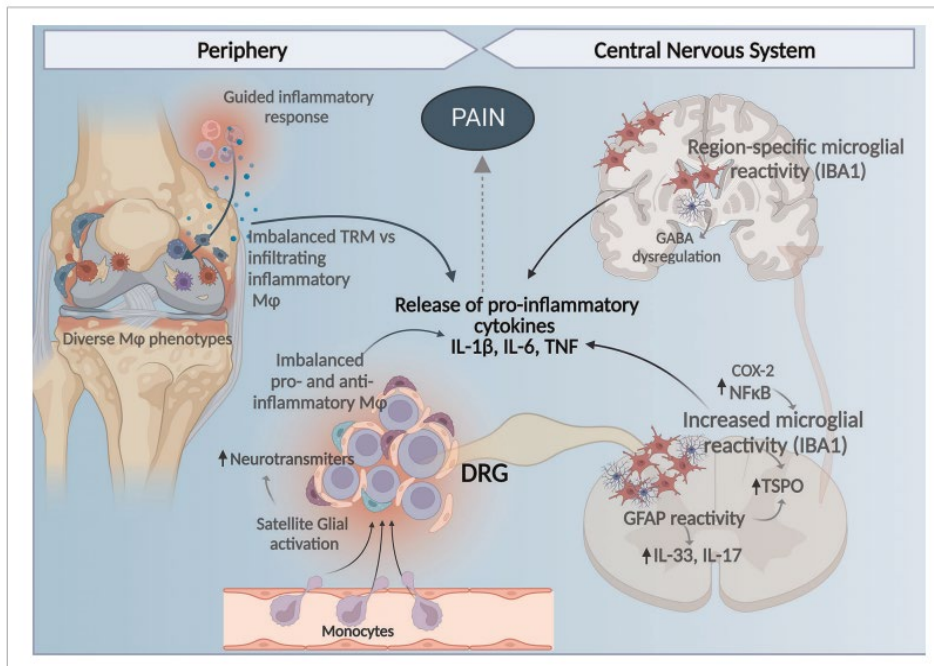
Vikas Agarwal • Ram Singh • Wiclaf •
Sandeep Chauhan • Anita Tahlan •
Chirag Kamal Ahuja • Deepak Goel • Lily Pal

Sural n. histology in patients with RA shows loss of myelinated fibers + perivascular lymphomononuclear infiltrate.

65% of patients with RA have subclinical neuropathy.

Neuropathic
Lesion or disease of somatosensory nervous system

Pro-inflammatory cytokines activate glial cells producing **free radicals, NO, chemo and cytokines** causing **neuronal damage**.



Neuropathic Pain in IA

Ultrasound assessment of carpal tunnel in rheumatoid arthritis and idiopathic carpal tunnel syndrome

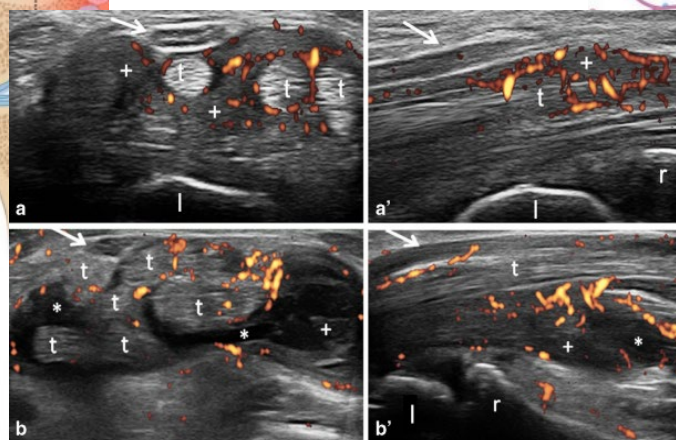
Check for updates

Gianluca Smerilli¹  • Andrea Di Matteo^{1,2}  • Edoardo Cipolletta¹  • Sergio Carloni³ • Antonella Incorvaia¹ • Marco Di Carlo¹  • Walter Grassi¹ • Emilio Filippucci¹ 

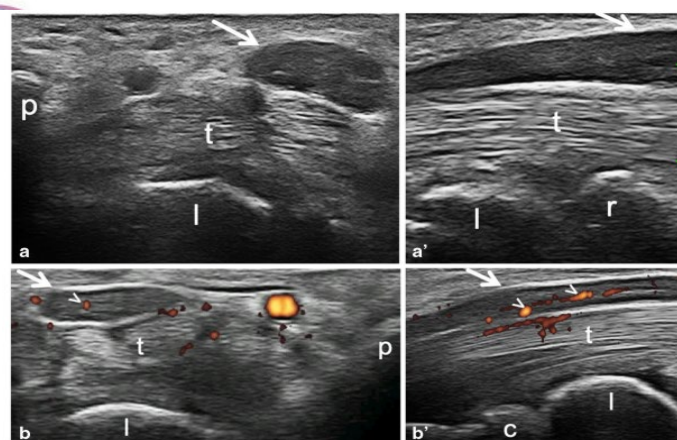
Received: 5 June 2020 / Revised: 9 July 2020 / Accepted: 13 July 2020 / Published online: 21 July 2020
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Rheumatoid Arthritis



Idiopathic CTS



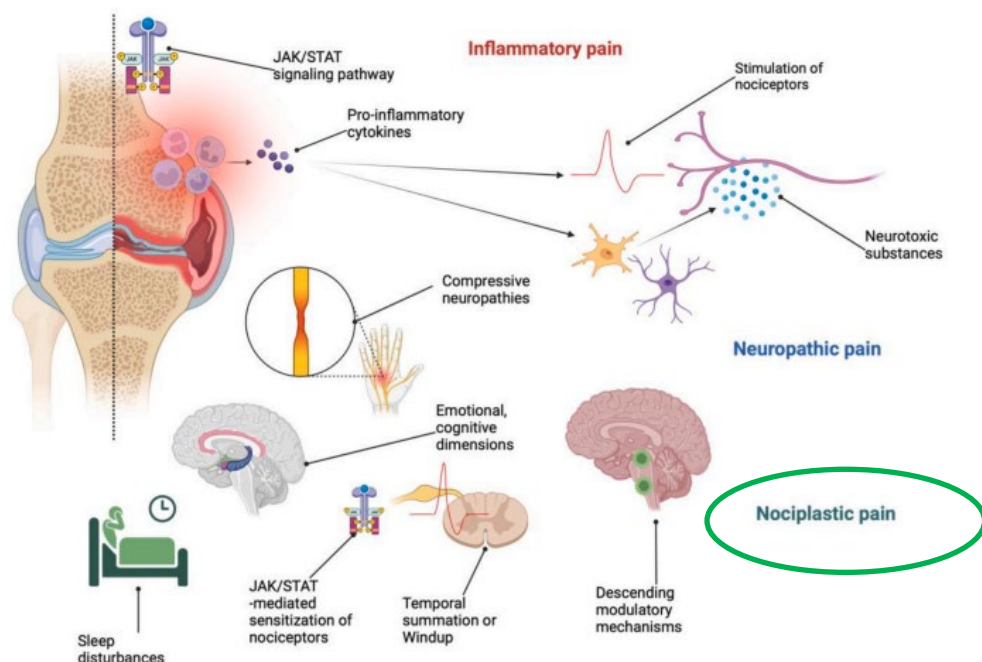
atosensory
n

Patients with RA have altered ultrasound findings including synovial hypertrophy, tenosynovitis in CTS compared to those with idiopathic CTS who demonstrate median nerve enlargement and abnormal intraneural blood flow.

Pain in the absence of synovitis or disease in remission?

Nociplastic Pain

Pain in inflammatory arthritis is complex and multifactorial.



Nociplastic
 Δ sensory pathways ie descending inhibitory $\rightarrow \uparrow$ pain sensitivity & altered nociception in the absence of tissue injury

Neurogenic inflammation promotes cycle of neurogenic sensitization $\rightarrow$ CHRONIC PAIN

Learning Objectives

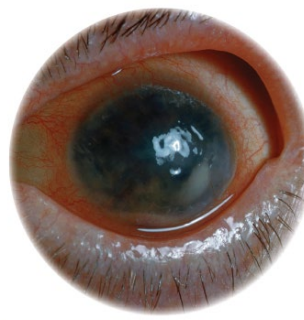
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Rheumatic Pain: Evaluation

□ History:

- ? Inciting event or trauma
- ? **Family history**
- ? **ROS: skin, bowel, renal, CV, pulm**
- ? Timing
- ? Alleviating and aggravating factors - ie transitional movements, after prolonged immobility, worse at night? Weather?

□ Physical Exam:



Clinical Photography, Central Manchester University
Hospitals NHS Foundation Trust, UK / Science Source



Rheumatic Pain: Evaluation

painDETECT PAIN QUESTIONNAIRE

Date: / / Patient: Last name: First name:

How would you assess your pain now, at this moment?
0 1 2 3 4 5 6 7 8 9 10
none max.

How strong was the strongest pain during the past 4 weeks?
0 1 2 3 4 5 6 7 8 9 10
none max.

How strong was the pain during the past 4 weeks on average?
0 1 2 3 4 5 6 7 8 9 10
none max.

Mark the picture that best describes the course of your pain:

Persistent pain with slight fluctuations ☐
Persistent pain with pain attacks ☐
Pain attacks without pain between them ☐
Pain attacks with pain between them ☐

Does your pain radiate to other regions of your body? yes ☐ no ☐
If yes, please draw the direction in which the pain radiates.

Do you suffer from a burning sensation (e.g., stinging nettles) in the marked areas?
never ☐ hardly noticed ☐ slightly ☐ moderately ☐ strongly ☐ very strongly ☐

Do you have a tingling or prickling sensation in the area of your pain (like crawling ants or electrical tingling)?
never ☐ hardly noticed ☐ slightly ☐ moderately ☐ strongly ☐ very strongly ☐

Do you have sudden pain attacks in the area of your pain, like electric shocks?
never ☐ hardly noticed ☐ slightly ☐ moderately ☐ strongly ☐ very strongly ☐

Is cold or heat (both water) in this area occasionally painful?
never ☐ hardly noticed ☐ slightly ☐ moderately ☐ strongly ☐ very strongly ☐

Do you suffer from a sensation of numbness in the areas that you marked?
never ☐ hardly noticed ☐ slightly ☐ moderately ☐ strongly ☐ very strongly ☐

Does slight pressure in this area, e.g., with a finger, trigger pain?
never ☐ hardly noticed ☐ slightly ☐ moderately ☐ strongly ☐ very strongly ☐

(To be filled out by the physician)

never ☐ hardly noticed ☐ slightly ☐ moderately ☐ strongly ☐ very strongly ☐

x=0 x=1 x=2 x=3 x=4 x=5

Total score out of 56

Development/Reference: R. Freynhagen, G. Baron, U. Giesecke, T.A. Tölle (1997) *Brain* 120, 1011-1021 (1997)
painDETECT questionnaire, version 1.00 (Pain Pharma GmbH, used with permission)

painDETECT SCORING OF PAIN QUESTIONNAIRE

Date: / / Patient: Last name: First name:

Please transfer the total score from the pain questionnaire:

Total score

Please add up the following numbers, depending on the marked pain behavior pattern and the pain radiation. Then total up the final score:

Persistent pain with slight fluctuations 0
Persistent pain with pain attacks -1 if marked, or
Pain attacks without pain between them +1 if marked, or
Pain attacks with pain between them +1 if marked

Radiating pains? +2 if yes

Final score

Screening Result
Final score

0 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32

nociceptive unclear neuropathic

A nociceptive pain component is unlikely (< 10%)
Result is ambiguous, however a neuropathic pain component can be present
A neuropathic pain component is likely (> 90%)

This sheet does not replace medical diagnostics.
It is used for screening the presence of a neuropathic pain component.

Development/Reference: R. Freynhagen, G. Baron, U. Giesecke, T.A. Tölle (1997) *Brain* 120, 1011-1021 (1997)
©2003 Pain Pharma GmbH

Today's date: / / ID No: _____

First name: Surname:

SF-36 Questionnaire

This questionnaire asks for your views about your health. For ALL questions, please tick, cross or colour the circle that most closely matches your response. There are no right or wrong answers. Please answer ALL questions.

1. In general, would you say your health is:

Poor	Fair	Good	Very good	Excellent
☐	☐	☐	☐	☐

2. Compared to one year ago, how would you rate your health general in now?

Much worse now than one year ago	Somewhat worse than one year ago	About the same as one year ago	Somewhat better than one year ago	Much better than one year ago
☐	☐	☐	☐	☐

3. The following questions are about activities you might do during a typical day. Does your health now limit you in these activities? If so, how much?

	No, not limited at all	Yes, limited a little	Yes, limited a lot
a. Vigorous activities, such as running, lifting heavy objects, participating in strenuous sports	☐	☐	☐
b. Moderate activities, such as moving a table, pushing a vacuum cleaner, bowling, or playing	☐	☐	☐
c. Lifting or carrying groceries	☐	☐	☐
d. Climbing several flights of stairs	☐	☐	☐
e. Climbing one flight of stairs	☐	☐	☐
f. Bending, kneeling or stooping	☐	☐	☐
g. Walking more than a mile	☐	☐	☐
h. Walking several blocks	☐	☐	☐
i. Walking one block	☐	☐	☐
j. Bathing or dressing yourself	☐	☐	☐

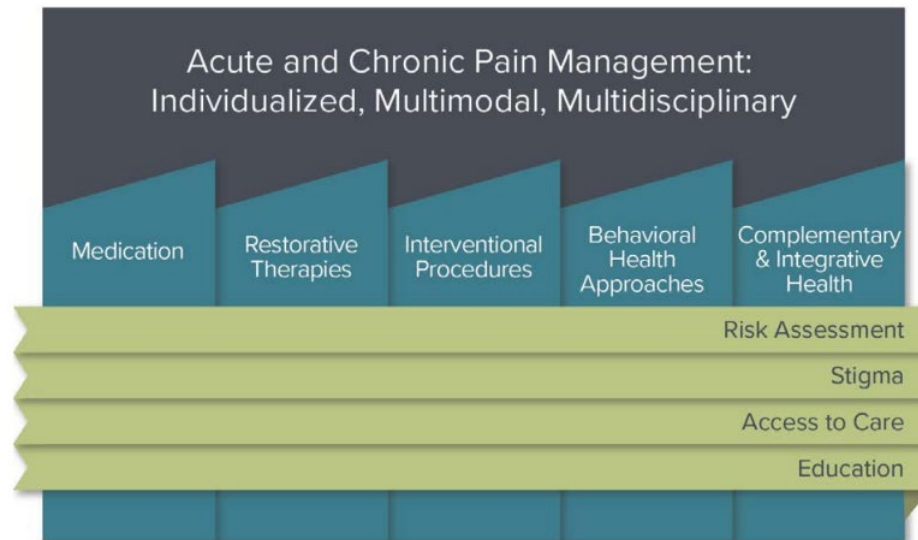
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Treatment

PAIN MANAGEMENT BEST PRACTICES INTER-AGENCY TASK FORCE REPORT

Updates, Gaps, Inconsistencies, and Recommendations



Individualized

Multimodal

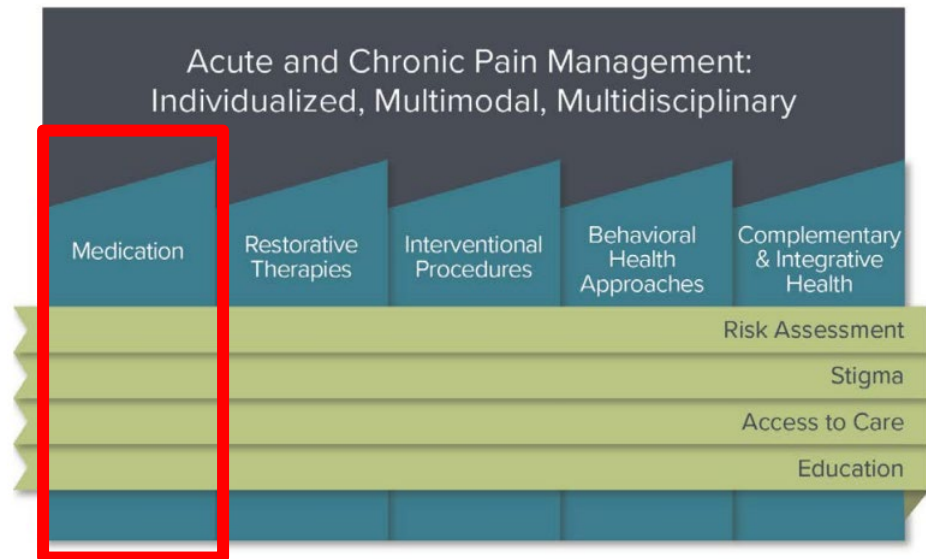
Multidisciplinary

Adapted from (2019) Pain Management Best Practices Inter-Agency Task Force.

Treatment

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CDC recommends non-pharmacologic and non-opioids as first-line therapies, when clinically appropriate.

Analgesics

Table D. Pharmacologic Treatments			
Class of Medication	Indications ^a	Magnitude of Benefit ^b	
		PAIN	FUNCTION
1 NSAIDs (topical or oral)	Low back pain, osteoarthritis, inflammatory arthritis, acute musculoskeletal (MSK) pain	Small to moderate	None to small
2 Acetaminophen	Acute MSK pain	Small	None
Antidepressants	Diabetic peripheral neuropathy, fibromyalgia	Small	None
Anticonvulsants	Diabetic peripheral neuropathy, fibromyalgia	Small to moderate	None (neuropathic pain) Small (fibromyalgia)
Opioids	Acute MSK pain, chronic pain, neuropathy	Small to no benefit ^c	Small to no benefit ^c

Adapted from (2021) AAFP Chronic Pain Management Toolkit. Pain Management Section .

Cox3 + 5HT inhibition
SNRI: duloxetine + milnacipran
Pregabalin > gabapentin (FM)

Utilize the **lowest effective dosage** for **pain relief** and **functional improvement**.

Review of publications evaluating opioid use in patients with inflammatory rheumatic disease

Christine Anastasiou and Jinoos Yazdany

Table 1. Selected studies reporting opioid use among patients with rheumatic diseases

Reference	Data source	Study design	Patients	Summary of opioid use
Rheumatoid arthritis (RA)				
Baker <i>et al.</i> [8*]	FORWARD databank	Cohort study, 1/1999–2/2019	37,868 patients with RA	27% Any opioid use
Chen <i>et al.</i> [29]	Truven Health MarketScan claims data	Retrospective observational study of US claims data, 2003–2014	181,710 patients with RA	19% Chronic opioid use
Curtis <i>et al.</i> [11]	US Medicare data	Retrospective observational study of US Medicare data, 2006–2014	240,750 patients with RA	41% Chronic opioid use, 19% Intermittent opioid use in 2014
Huang <i>et al.</i> [6*]	National Ambulatory Medical Care Survey, 2011–2016	Cross-sectional survey, 2011–2016	Estimated 4.5 million encounters with primary diagnosis of RA	Proportion of visits with opioid prescription: 24.3%
Lee YC <i>et al.</i> [14]	Corrone RA registry	Cohort study, 2002–2016	33,739 patients with RA	16.9% Chronic* opioid use in 2015
Navarro-Millán <i>et al.</i> [7*]	Medicare and Medicaid services claims data	Retrospective observational study of US claims data, 2007, 2011, 2014	43,563 patients with RA <65 years old receiving SSDI Medicare and Medicaid	63.7% Chronic opioid use in 2014
Park <i>et al.</i> [10]	IQVIA™ Health Plan Claims Data	Retrospective observational study of US claims data, 2007–2015	2,330 patients with RA	51.0% Any opioid use
Systemic lupus erythematosus (SLE)				
Birt <i>et al.</i> [30*]	IBM MarketScan Databases	Retrospective observational study of US claims data, 1/2012–5/2018	49,413 patients with SLE	52.6% Any opioid use 34.6% Chronic opioid use
Chen <i>et al.</i> [29]	Truven Health MarketScan claims data	Retrospective observational study of US claims data, 2003–2014	45,834 patients with SLE	16% Chronic opioid use
Lee J <i>et al.</i> [32*]	Single institution chart review	Retrospective observational chart review, 2013–2016	77 SLE patients who had persistent frequent ED visits	37.7% Chronic opioid use
Somers <i>et al.</i> [28]	MILES Cohort	Prospective cohort, 2/2014–9/2015	462 SLE patients	31.0% Any opioid use 21% Chronic** opioid use
Psoriasis and psoriatic arthritis (PsA)				
Chen <i>et al.</i> [29]	Truven Health MarketScan claims data	Retrospective observational study of US claims data, 2003–2014	30,307 patients with PsA	15% Chronic opioid use
Hunter <i>et al.</i> [9**]	HealthCore Integrated Research Database	Retrospective observational study of US claims data, 1/2013–7/2019	921 patients with psoriatic arthritis	33.8% Any opioid use 12 months after initiation of biologic
Loft <i>et al.</i> [34*]	Danish Skin Cohort	Prospective cohort study	4016 patients with psoriasis, 847 with concomitant PsA	13–25.6% Any opioid use within the past year
Noe <i>et al.</i> [38]	Optum Electronic Health Records Database	Retrospective study of US claims data, 1/2007–6/2017	99,830 patients with psoriasis	1.9% of opioid-naïve patients with psoriasis received an incident opioid prescription over one year.
Taylor <i>et al.</i> [35]	National Ambulatory Medical Care Survey (2006–2016) & National Hospital Ambulatory Medical Care Survey (2006–2011)	Cross-sectional survey, 2006–2016	1148 encounters for psoriasis and PsA evaluated, weighted to a US national estimate of 27 million visits	Proportion of visits with opioid prescription: 10%
Walsh <i>et al.</i> [36]	Optum Research Database	Retrospective study of US claims data, 1/2012–4/2016	1,235 patients with PsA	48.6% Any opioid use

KEY POINTS

- Chronic opioids are commonly prescribed for patients with RA, SLE, psoriasis or psoriatic arthritis, and ankylosing spondylitis, with evidence of associated adverse effects.
- Opioid use minimally decreased but remained high after biologic DMARD medications were initiated.
- There is no data reporting improved function, quality of life, or pain control with long-term use of opioids for patients with inflammatory rheumatic diseases, making this an important area for future research.

Analgesic Harms

Nonopioid Pharmacologic Treatments for Chronic Pain

Prepared for:
Agency for Healthcare Research and Quality
U.S. Department of Health and Human Services
5600 Fishers Lane
Rockville, MD 20857
www.ahrq.gov

Table B-2. Harms

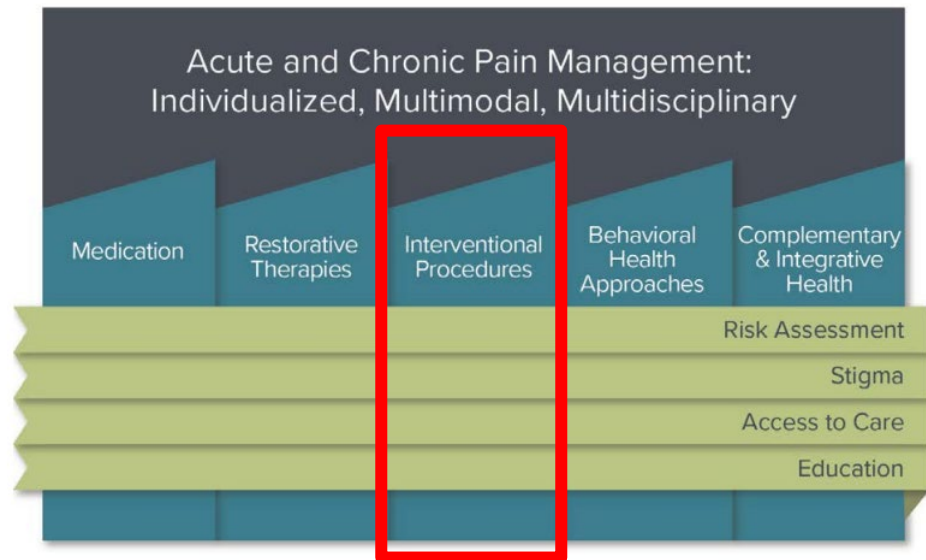
Drug(s)/Drug Class	Harms by Drug Class
All drugs	Withdrawal due to adverse events, serious adverse events, overdose, misuse, and dependence
Serotonin-norepinephrine reuptake inhibitor antidepressants	Cognitive effects, nausea, sedation
Tricyclic antidepressants	Cardiac rhythm abnormalities, cognitive effects, dry mouth, urinary retention, weight gain
Pregabalin/gabapentin anticonvulsants	Blurred vision, cognitive effects, dizziness, peripheral edema, sedation, weight gain
Oxcarbazepine/carbamazepine anticonvulsants	Cognitive effects, hyponatremia, neutropenia, sedation
NSAIDs	CV events, GI, liver dysfunction, renal dysfunction
Skeletal muscle relaxants	Dry mouth, sedation, urinary retention
Acetaminophen	Liver toxicity
Memantine	Cardiac rhythm abnormalities, cognitive effects, dizziness, sedation
Topical (any)	Application site reactions
Topical lidocaine	Cardiotoxicity, cognitive effects
Topical diclofenac	CV events, GI, liver dysfunction, renal dysfunction
Cannabis	Addiction/dependence, cognitive effects, hyperemesis, nausea, sedation

Pharmacologic Management can be **ineffective** and **limited**.

Interventional Treatments

PAIN MANAGEMENT BEST PRACTICES INTER-AGENCY TASK FORCE REPORT

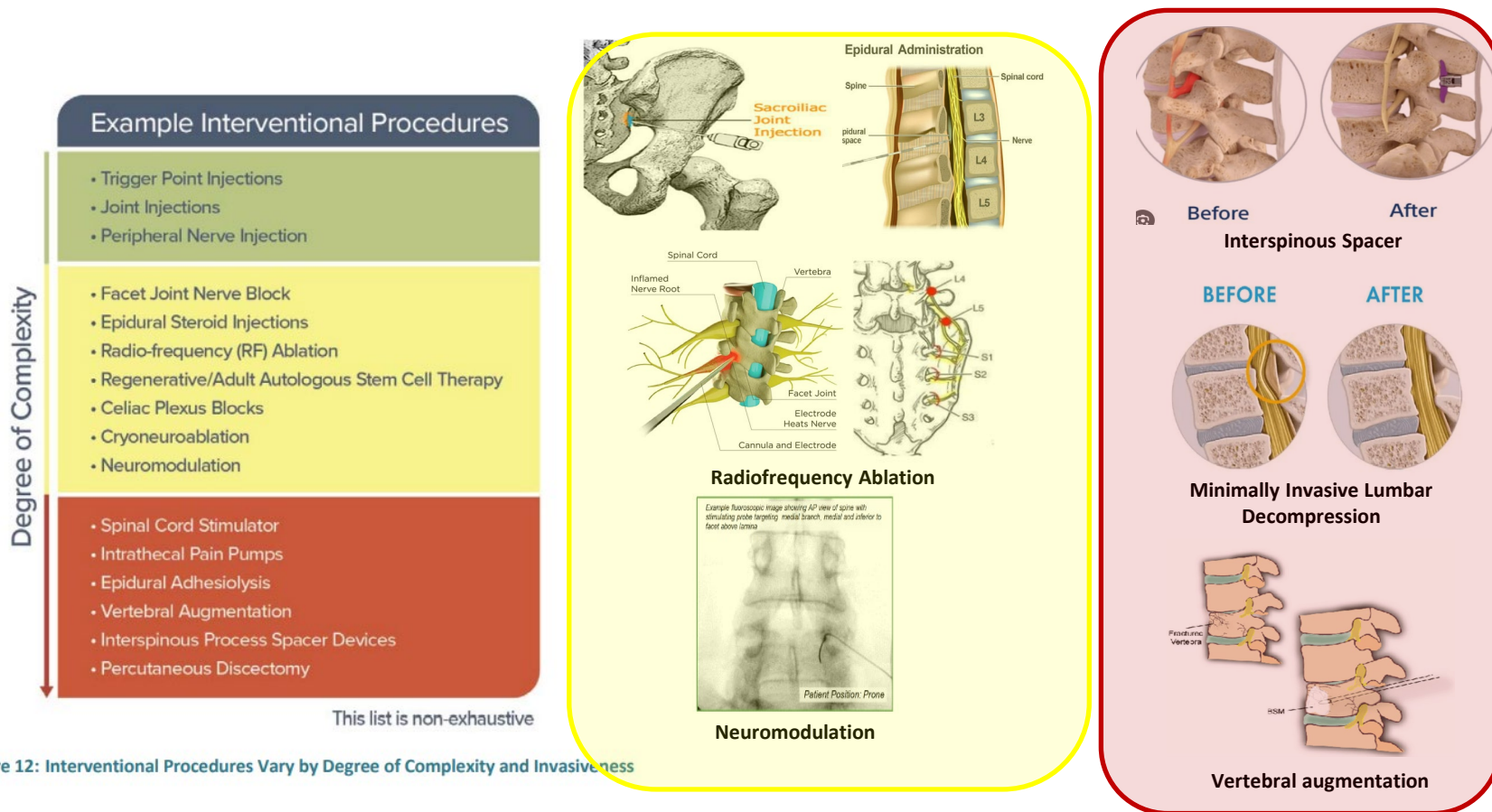
Updates, Gaps, Inconsistencies, and Recommendations



Individualized
Multimodal
Multidisciplinary

Adapted from (2019) Pain Management Best Practices Inter-Agency Task Force.

Interventional Treatments



Interventional Treatments

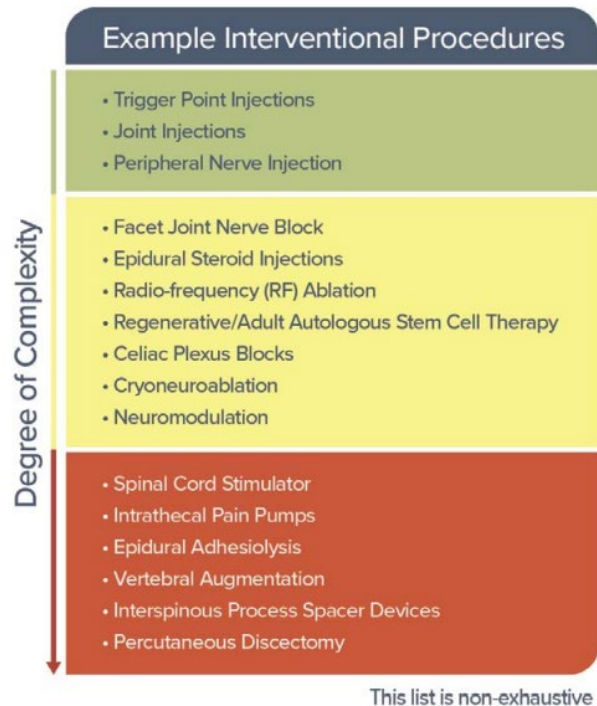


Figure 12: Interventional Procedures Vary by Degree of Complexity and Invasiveness

- ☐ **Fewer side effects** in comparison to pharmacologic interventions
- ☐ **Combine** interventions with **therapy** for **synergistic** benefit
- ☐ Interventions **limit** the **need** for **pharmacologic** interventions or **surgery**

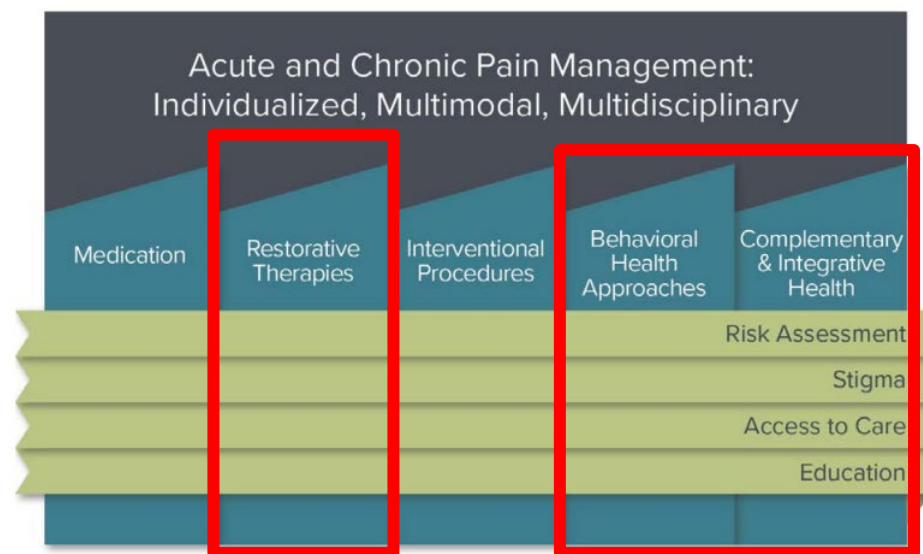
Interventions offer a direct, targeted approach to both diagnosing and treating a pain generator however requires complex decision making.

PRP in systemic inflammation?

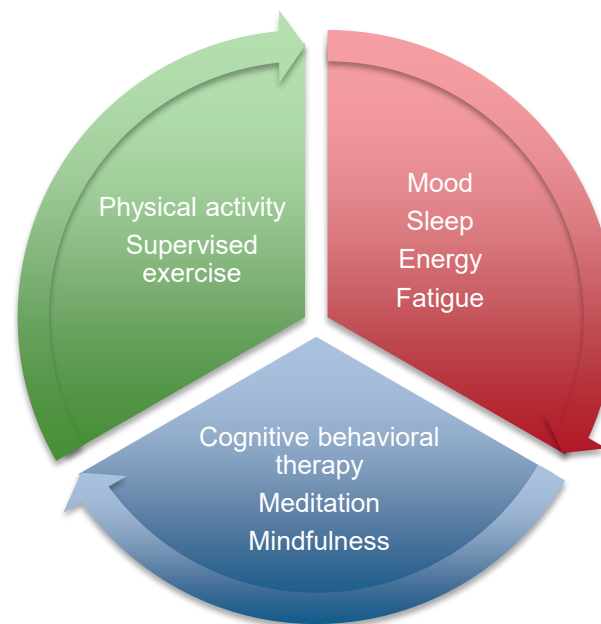
Multimodal Treatment

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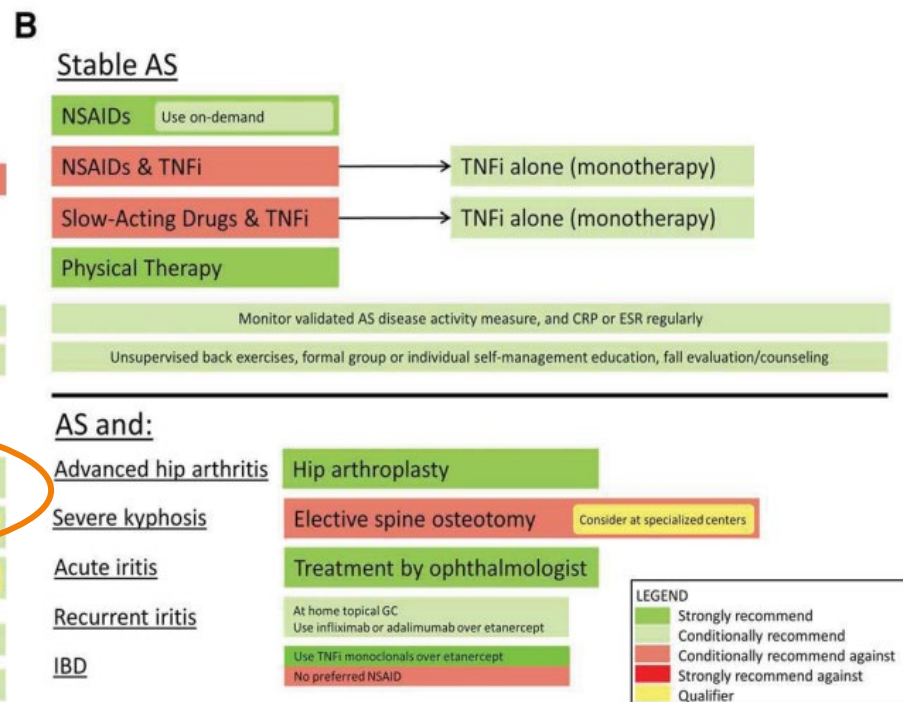
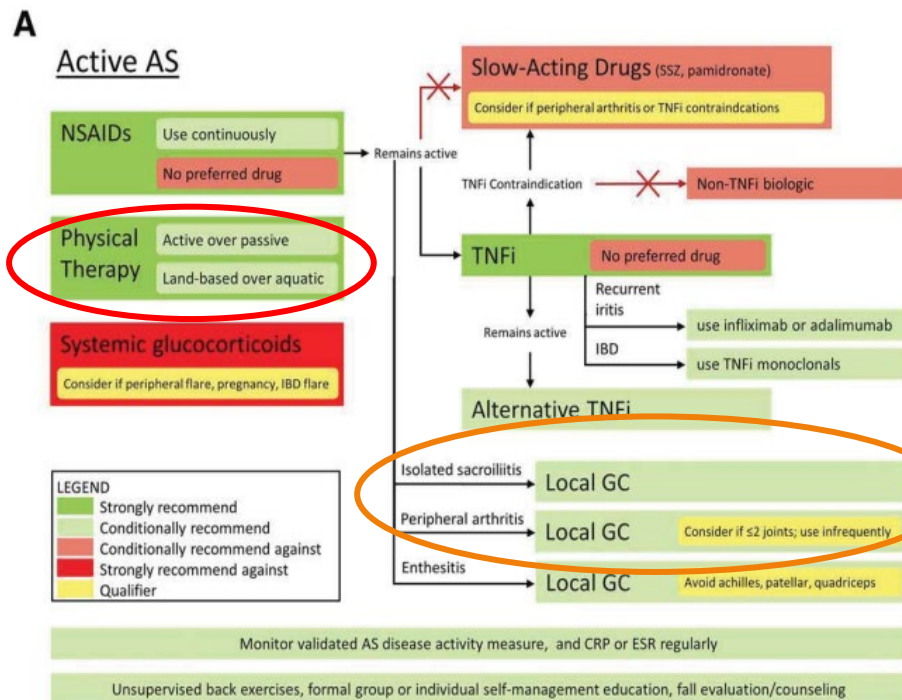
Goal of **multimodal** treatment is to improve **function**.

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American College of Rheumatology Guidelines by Rheumatologic Disease

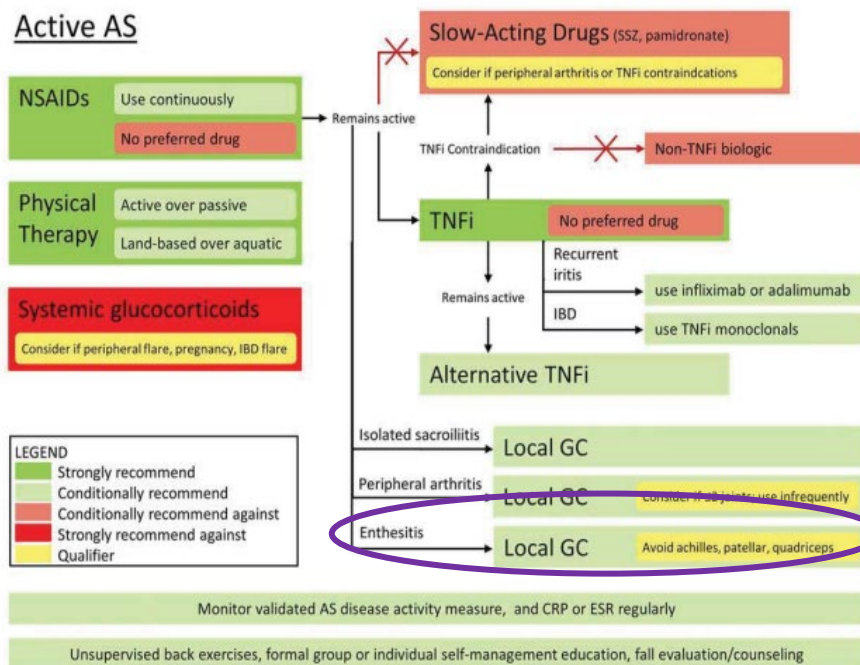
Ankylosing Spondylitis



Ankylosing Spondylitis

A

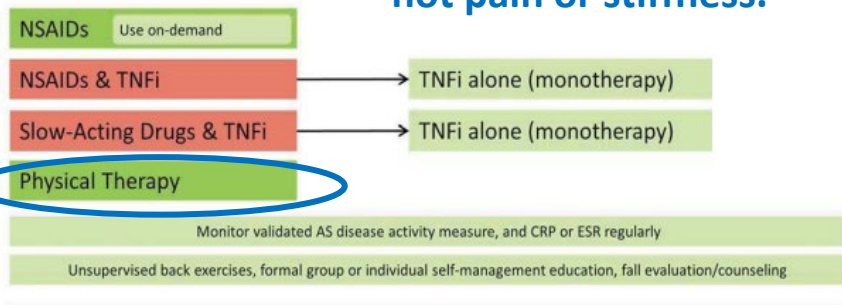
Active AS



B

Stable AS

PT improved disease activity/function
– not pain or stiffness.



AS and:

Advanced hip arthritis

Hip arthroplasty

Severe kyphosis

Elective spine osteotomy
Consider at specialized centers

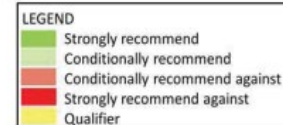
Acute iritis

Treatment by ophthalmologist

Recurrent iritis

At home topical GC
Use infliximab or adalimumab over etanercept
Use TNFi monoclonals over etanercept
No preferred NSAID

IBD



greater trochanter, pelvic rim, and plantar fascia?

Rheumatoid Arthritis

Table 4. Treatment modification*

2021 Trea	Recommendations	Certainty of evidence	Based on the evidence report(s) of the following PICO(s)	Evidence table(s), in Supp. App. 2
Liana Fr	A TTT approach is strongly recommended over usual care for patients who have not been previously treated with bDMARDs or tsDMARDs.	Low	PICO 12.a	p. 191
Kristine	A TTT approach is conditionally recommended over usual care for patients who have had an inadequate response to bDMARDs or tsDMARDs.	Very low	PICO 12.b	p. 199
Mary C.	A minimal initial treatment goal of low disease activity is conditionally recommended over a goal of remission.	Low	PICO 13	p. 201
Shilpa Ve	Addition of a bDMARD or tsDMARD is conditionally recommended over triple therapy for patients taking maximally tolerated doses of methotrexate who are not at target.	Very low	PICO 19.C2–C6†	p. 240–1
Jennifer	Switching to a bDMARD or tsDMARD of a different class is conditionally recommended over switching to a bDMARD or tsDMARD belonging to the same class for patients taking a bDMARD or tsDMARD who are not at target.	Very low	PICO 24–27†	p. 293–338
Lara Ka	Addition of/switching to DMARDs is conditionally recommended over continuation of glucocorticoids for patients taking glucocorticoids to remain at target.	Very low	PICO 23	p. 292
Namrata	Addition of/switching to DMARDs (with or without IA glucocorticoids) is conditionally recommended over the use of IA glucocorticoids alone for patients taking DMARDs who are not at target.	Very low	PICO 28.C1–C2	p. 339–40

* PICO = population, intervention, comparator, and outcomes; Supp. App. 2 = Supplementary Appendix 2, available on the *Arthritis Care & Research* website at <http://onlinelibrary.wiley.com/doi/10.1002/acr.24596/abstract>; TTT = treat-to-target; bDMARDs = biologic disease-modifying antirheumatic drugs; tsDMARDs = targeted synthetic DMARDs; IA = intraarticular.

**DMARDs should be adjusted to reduce disease activity,
irrespective of treatment with IA glucocorticoids.**

Integrative Medicine: RA

Arthritis Care & Research

Vol. 75, No. 8, August 2023, pp 1603–1615
 DOI 10.1002/acr.25117
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2022 American College of Rheumatology Guideline for Exercise, Rehabilitation, Diet, and Additional Integrative Interventions for Rheumatoid Arthritis

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Regular, tailored exercise results in improved pain + function – whether aerobic, resistance, aquatic, mind-body.

Table 1. Recommendations on integrative interventions for the management of rheumatoid arthritis (RA)

Exercise	Rehabilitation	Diet	Additional
Consistent engagement in exercise (++)	Comprehensive occupational therapy (+)	Mediterranean-style diet (+)	Standardized self-management program (+)
Aerobic exercise (+)	Comprehensive physical therapy (+)	Against formally defined diet other than Mediterranean-style (-)	Cognitive behavioral therapy and/or mind-body approaches (+)
Aquatic exercise (+)	Hand therapy exercises (+)	Against dietary supplements (-)	Acupuncture (+)
Resistance exercise (+)	Splinting, orthoses, compression, bracing, and/or taping (+)		Massage therapy (+)
Mind-body exercise (+)	Joint protection techniques (+)		Thermal modalities (+)
	Activity pacing, activity modification, energy conservation, and/or fatigue management (+)		Against electrotherapy (-)
	Assistive devices, adaptive equipment, and/or environmental adaptations (+)		Against chiropractic therapy (-)
	Vocational rehabilitation, work site evaluations and/or modifications (+)	Strong recommendation FOR Conditional recommendation FOR Conditional recommendation AGAINST	

Exercise in RA

Journal of Pain Research

Dovepress
Taylor & Francis Group

 Open Access Full Text Article

REVIEW

Effect of Exercise Interventions for Rheumatoid Arthritis: A Systematic Review and Network Meta-Analysis of Randomised Controlled Trials

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“Exercise interventions are effective supplements for RA treatment, with specific exercises offering distinct benefits: Pilates for pain, aerobic exercise + resistance exercise for morning stiffness, traditional Chinese exercise for disease activity.”

Rheumatoid Arthritis

Cox et al. BMC Medicine (2025) 23:54
https://doi.org/10.1186/s12916-025-03870-0

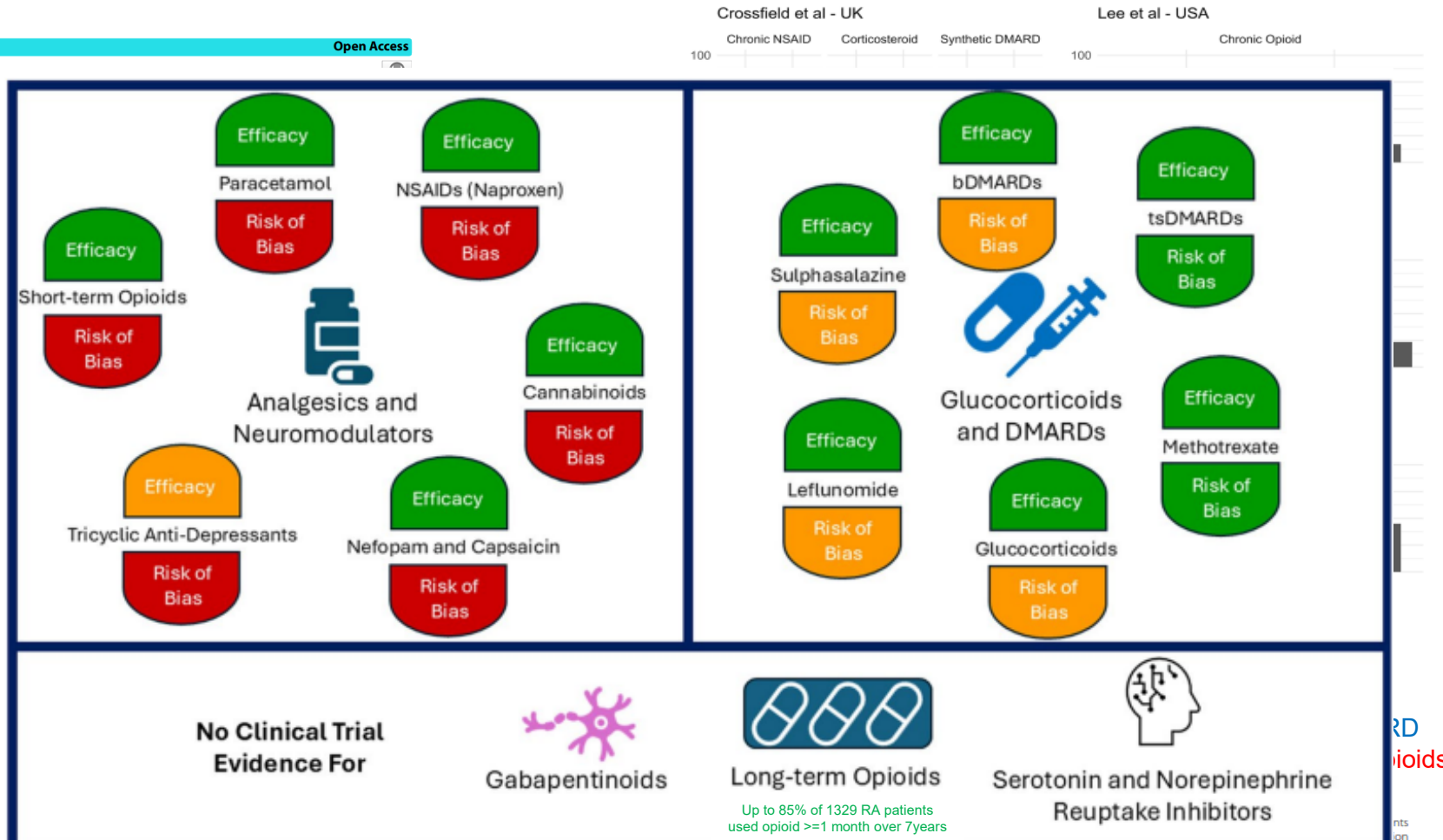
BMC Medicine

REVIEW

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Pharmac
in patient
literature

Natasha Cox^{1,2}, Chri



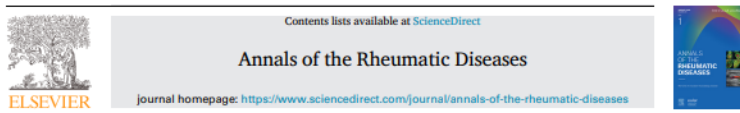
includes patients with RA, psoriatic arthritis, and axial spondyloarthritis (although RA was the commonest diagnosis). Crossfield et al.: DMARD and corticosteroid prescriptions are long-term. Navarro-Millan et al.: the population comprises beneficiaries of Social Security Disability Insurance (no longer working because they are considered disabled) and <65 years old. Studies included in the figure are those reporting the prevalence of chronic NSAID, chronic opioid, DMARD, and corticosteroid prescriptions/use in > one calendar year that contain extractable data

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Rheumatoid Arthritis: Injections

Ann Rheum Dis 84 (2025) 937–948



Rheumatoid arthritis

Treatment with methotrexate plus oral prednisolone versus triple therapy (methotrexate/sulfasalazine/hydroxychloroquine) plus intra-articular glucocorticoids in early rheumatoid arthritis: a prespecified nonrandomised subgroup analysis of clinical and radiographic data at 48 weeks from the NORD-STAR trial's conventional treatment arm

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Early RA: triple therapy + mandatory IA GCI had numerically better clinical outcomes, fewer withdrawals, fewer adverse events, and lower cumulative dose of glucocorticoids, but slightly worse radiographic outcomes than treatment with methotrexate and oral prednisolone.

Gout

ACR GUIDELINE FOR MANAGEMENT OF GOUT

Table 6. Gout flare management*

M	Recommendation	PICO question	Certainty of evidence
Joh An Ca Jan Pu Tai An	For patients experiencing a gout flare, we strongly recommend using oral colchicine, NSAIDs, or glucocorticoids (oral, intraarticular, or intramuscular) as appropriate first-line therapy for gout flares over IL-1 inhibitors or ACTH (the choice of colchicine, NSAIDs, or glucocorticoids should be made based on patient factors and preferences). When colchicine is the chosen agent, we strongly recommend low-dose colchicine over high-dose colchicine given its similar efficacy and fewer adverse effects.	32	High
	For patients experiencing a gout flare for whom other antiinflammatory therapies are poorly tolerated or contraindicated, we conditionally recommend using IL-1 inhibition over no therapy (beyond supportive/analgesic treatment).	33	Moderate
	For patients who may receive NPO, we strongly recommend glucocorticoids (intramuscular, intravenous, or intraarticular) over IL-1 inhibitors or ACTH.	32	High
	For patients experiencing a gout flare, we conditionally recommend using topical ice as an adjuvant treatment over no adjuvant treatment.	31	Low
	Strongly recommend Conditionally recommend Strongly recommend against Conditionally recommend against		

IA steroid OR oral NSAID treatment are strongly recommended in managing acute gout flares.

Systemic Lupus

2025 American College of Rheumatology (ACR) Guideline for the Treatment of Systemic Lupus Erythematosus (SLE)

Guideline Summary

Recommendations and Good Practice Statements	Strength	Level of Evidence
Organ-specific manifestations		
Musculoskeletal		
GPS: <i>Initial therapy for acute or recurrent episodes of inflammatory arthritis in people with SLE may include a course of NSAID or a limited course of oral glucocorticoid while waiting for recommended long-term therapies to take effect.</i>		
Arthritis: For persistent or recurrent active SLE arthritis on HCQ, regardless of prior/current NSAIDs or short-term glucocorticoid therapy: ...We conditionally recommend initial therapy with MTX, MPAA, or AZA, with a low threshold to add or substitute with belimumab or anifrolumab for inadequate response over initial biologic therapy.	Conditional	Very Low to Low

Recommend **limited use** of **oral NSAID** or **steroid** for **acute SLE arthritis** – goal is to start OR escalate long-term therapy.

Psoriatic Arthritis

Arthritis & Rheumatology

Vol. 71, No. 1, January 2019, pp 5–32

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SPECIAL ARTICLE

2018 American Foundation

Jasvinder A. Singh,
 Maureen Dubreuil,
 Paula Marchetti,
 Bernadette Siat,
 Nancy Sullivan,
 Nowell,²⁴ Ana-Maria
 Jessica A. Walsh

Non-pharmacologic therapies	physical therapy, occupational therapy, smoking cessation, weight loss, massage therapy, exercise
Symptomatic treatments	nonsteroidal anti-inflammatory drugs, glucocorticoids, local glucocorticoid injections
OSM	methotrexate, sulfasalazine, cyclosporine, leflunomide, apremilast
TNFi	etanercept, infliximab, adalimumab, golimumab, certolizumab pegol
IL12/23i	ustekinumab
IL17i	secukinumab, ixekizumab, brodalumab
CTLA4-Ig	abatacept
JAK inhibitor	tofacitinib

Figure 1. Pharmacologic, nonpharmacologic, and symptomatic therapies for psoriatic arthritis. Pharmacologic therapies are displayed in the blue boxes and include oral small molecules (OSMs), tumor necrosis factor inhibitor (TNFi) biologics, interleukin-17 inhibitor (IL-17i) biologics, an IL-12/23i biologic, CTLA4-immunoglobulin, and a JAK inhibitor. While there are numerous nonpharmacologic therapies available, 6 of these are addressed in this guideline. Symptomatic therapies include nonsteroidal antiinflammatory drugs, systemic glucocorticoids, and local glucocorticoid injections. **Systemic glucocorticoids or local injections are not addressed in this guideline.**

Psoriatic Arthritis

Table 8. Recommendations for treatment of patients with active psoriatic arthritis with nonpharmacologic interventions (PICOs 1–8)*

	Level of evidence (evidence [refs.] reviewed)†
In adult patients with active PsA,	
1. Recommend exercise over no exercise (PICO 1) Conditional recommendation based on low-quality evidence; may consider no exercise in patients with existing muscle/tendon injury or multiple inflamed symptomatic joints with worsening pain with exercise.	Low (128)
2. Recommend low-impact exercise (e.g., tai chi, yoga, swimming) over high-impact exercise (e.g., running) (PICO 2) Conditional recommendation based on very-low-quality evidence; may consider high-impact exercise due to patient preference.	Very low
3. Recommend physical therapy over no physical therapy (PICO 3) Conditional recommendation based on very-low-quality evidence; may consider no physical therapy due to patient preference, out-of-pocket cost, distance to physical therapy site, or lack of transportation.	Very low
4. Recommend occupational therapy over no occupational therapy (PICO 4) Conditional recommendation based on low-quality evidence; may consider no occupational therapy due to patient preference, out-of-pocket cost, distance to occupational therapy site, or lack of transportation.	Low (129, 130)
5. Recommend weight loss over no weight loss for patients who are overweight/obese (PICO 5) Conditional recommendation based on low-quality evidence; may consider no weight loss due to additional patient burden involved with weight-loss program.	Low (131–133)
6. Recommend massage therapy over no massage therapy (PICO 7) Conditional recommendation based on very-low-quality evidence; may consider no massage therapy due to associated costs.	Very low (134)
7. Recommend acupuncture over no acupuncture (PICO 8) Conditional recommendation based on very-low-quality evidence; may consider no acupuncture due to associated costs.	Very low (135)
8. Recommend smoking cessation over no smoking cessation (PICO 6) Strong recommendation supported by moderate-quality evidence, rated down for indirectness.	Moderate (136, 137)

Patients with **active PsA**
 should use some form or
 combination of **exercise,**
physical therapy,
occupational therapy,
massage therapy, and
acupuncture.

Low-impact exercise is
recommended over high-
impact exercise.

* Active psoriatic arthritis (PsA) is defined as disease causing symptoms at an unacceptably bothersome level as reported by the patient, and judged by the examining clinician to be *due to PsA* based on ≥1 of the following: swollen joints, tender joints, dactylitis, enthesitis, axial disease, active skin and/or nail involvement, and extraarticular inflammatory manifestations such as uveitis or inflammatory bowel disease.

Psoriatic Arthritis: Injections

Clinical Rheumatology (2020)
<https://doi.org/10.1007/s1006>

ORIGINAL ARTICLE

*Intra-articular Treatment Options in the
Management of Joint Disorders*

Special Collection

 *Therapeutic Advances in Musculoskeletal Disease*

Review

Effectiveness o
from a multice

Nicolò Girolimetto^{1,2} •
Federica Martinis³ • Al
Carlo Salvarani^{2,5} • Ral

Corticosteroid injection treatment for dactylitis in psoriatic arthritis

Antonio Carriero^{id}, Ennio Lubrano^{id}, Valentina Picerno, Angela Anna Padula
and Salvatore D'Angelo^{id}

Ther Adv Musculoskel Dis

2021, Vol. 13: 1–12

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1759720X211041864

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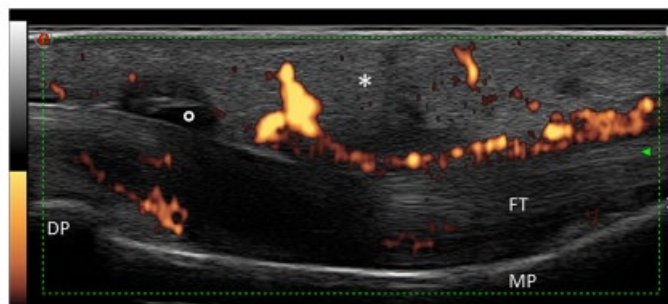


Figure 2. Tenosynovitis of the III right flexor tendon characterized by marked synovial proliferation, a moderate increase of the tendon thickness and a thickened and hypoechoic peritendineal tissue. Synovial sheath widening (circle) associated with soft-tissue edema (asterisk). Power Doppler function revealed diffuse and severe vascular signal inside and around the tendon sheath (Grade 3). DP, distal phalanx; FT, flexor tendon; MP, medial phalanx.

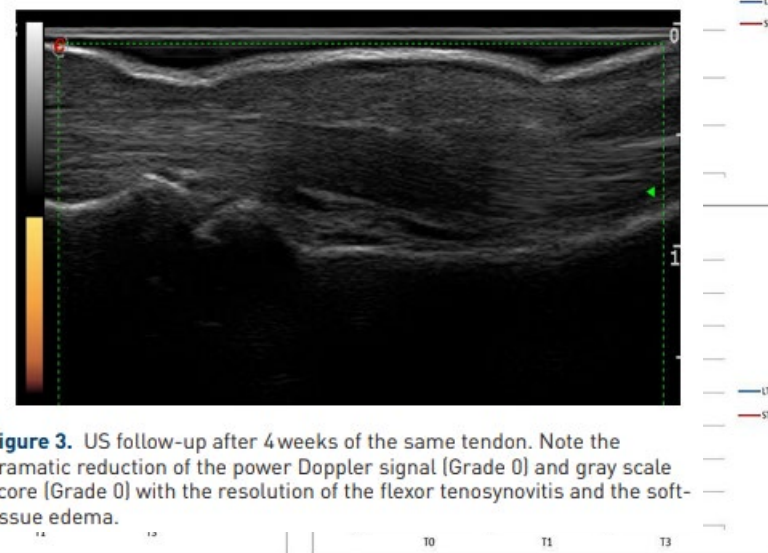
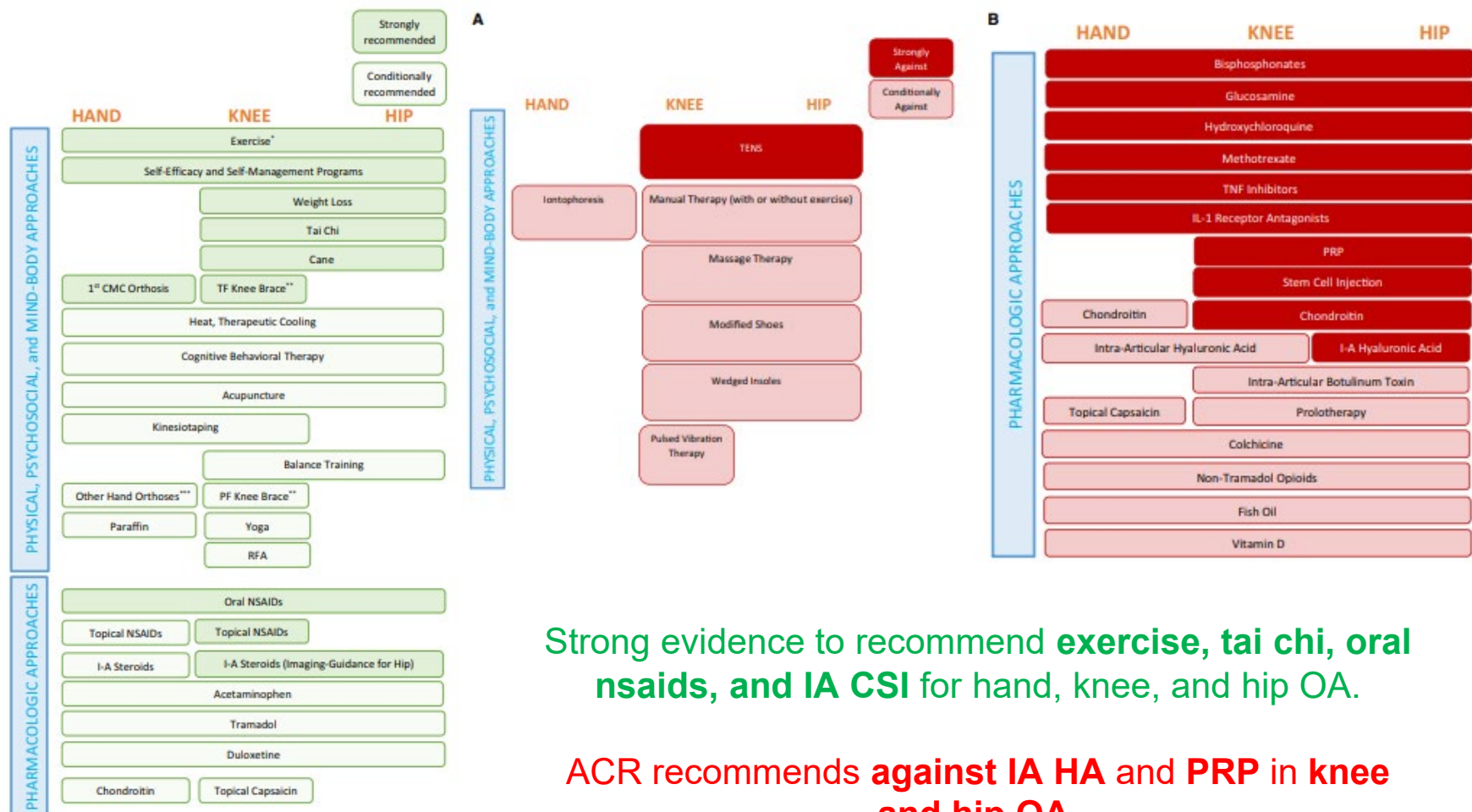


Figure 3. US follow-up after 4 weeks of the same tendon. Note the dramatic reduction of the power Doppler signal (Grade 0) and gray scale score (Grade 0) with the resolution of the flexor tenosynovitis and the soft-tissue edema.

Fig. 2 Changes in clinical parameters within time in both groups. * $p < 0.001$; ** $p = 0.008$. T1, 1 month; T3, 3 months; LDI-b, Leeds Dactylitis Index basic; VAS, visual analogue scale

Osteoarthritis



Strong evidence to recommend **exercise, tai chi, oral ns aids, and IA CSI** for hand, knee, and hip OA.

ACR recommends **against IA HA and PRP** in knee and hip OA.

Limitations

- ☐ *Rheumatology Advances in Practice*, 2024, **8**(4), rkae128
<https://doi.org/10.1093/rap/rkae128>
Guideline



British Society for
Rheumatology

RHEUMATOLOGY
ADVANCES IN PRACTICE

OXFORD

Guideline

- ☐ **Pain management in people with inflammatory arthritis:
British Society for Rheumatology guideline scope**

12

- ☐ Ian C. Scott ^{1,2,*}, Opeyemi Babatunde ¹, Christopher Barker³, Rebecca Beesley⁴, Richard Beesley⁴, Hollie Birkinshaw⁵, Mel Brooke⁶, Hema Chaplin ^{7,8}, Lara Chapman ⁹, Coziana Ciurtin ¹⁰, James Dale¹¹, Dervil Dockrell ¹², Emma Dures¹³, Kathryn Harrison¹⁴, Meghna Jani ^{15,16}, Charlotte Lee¹⁷, Maura McCarron^{18,19}, Christian D. Mallen ¹, Assie O'Connor², Claire Pidgeon²⁰, Tamar Pincus¹⁷, Dee Pratt²¹, Yeliz Prior ²², Karim Raza^{23,24}, Zoe Rutter-Locher ²⁵, Seema Sharma ²⁶, Katie Shaw²⁷, Samantha Small²⁸, Tilli Smith¹, Lesley Tiffin²⁹, Jordan Tsigarides ^{30,31}, Mikalena Xenophontos^{32,33}, Nicholas G Shenker ^{34,35}

Coming 2026! Reconvene TPS 2026?

Summary

- ❑ Inflammatory arthritis affects 300+ million individuals worldwide and contributes to functional disability
- ❑ Pain in rheumatologic diseases is multi-factorial and neurogenic inflammation contributes to chronic pain
- ❑ Multimodal treatment is mainstay of treatment
- ❑ Interventions are recommended by ACR in:
 - ❑ Acute gout
 - ❑ PsA dactylitis
 - ❑ Isolated sacroiliitis or 1-2 peripheral joints affected in AS
 - ❑ RA: combine with triple therapy?
- ❑ Need for high-quality randomized placebo-controlled trials

Thank you for your attention!



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