



The Essential Role of Primary Care in the Management of Knee Osteoarthritis

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Objectives

- Review knee OA diagnosis
- Utilize appropriate imaging
- Review non-operative management
- Understand how to access and apply AAOS CPG and AUC
- Identify timing and management for surgical referral

Important Note

- Evidence Based strategies from AAOS CPG and AUC
- CPG / AUC offers summaries – not mandates
- Individual patient context matters
- Clinical judgment required

What Is Knee Osteoarthritis?

- Degenerative joint disease--Pain and dysfunction
- Knee is most common joint involved
- Common but variable, 1/3 people > age 75 have it
- Primary care handles greatest burden

Why This Matters in Primary Care

- Very common condition
- Long-term management
- Information often misleading to patients
- Expectations shape outcomes

Diagnosis of Knee OA

- Clinical + radiographic
- OA often coexists with other conditions
- Imaging \neq pain
- Context matters, not all pain is OA

Clinical Presentation

- Pain and stiffness
- Swelling
- ROM limitation
- Decreased function

Differential Diagnosis

- Spine / radiculopathy
- Hip pathology, other bony issues
- Inflammatory arthritis, soft tissue injury
- Infection or neoplasm
- Vascular claudication

Diagnostic Caution

- Do not inject blindly
- Image first
- Rule out red flags
- Safety first

Imaging Approach

- X-ray almost always sufficient for AO Dx
- Weight-bearing views
- MRI rarely needed– adds anxiety
- Imaging ≠ symptoms, must correlate

Kellgren–Lawrence Grading

- Describes radiographic severity
- Used in guidelines, for imaging reliability
- Descriptor, Not a pain scale,
- Does not dictate treatment

Kellgren- Lawrence	OA Grading Scale
Grade 0–1	No or doubtful OA, possible osteophytes
Grade 2	Definite osteophytes, possible joint space narrowing
Grade 3	Joint space narrowing, sclerosis , osteophytes
Grade 4	Severe OA with deformity, joint space narrowing, osteophytes

K-L Grade 4 on R; Grade 3 on L**Non-Operative Treatment Goals**

- Reduce pain
- Improve function, preserve motion and activity
- Slow progression of disease and morbidity
- Manage expectations

CPG STRONG Recommendations

- Topical NSAIDs
- Supervised exercise
- Neuromuscular training
- Self-management programs

CPG MODERATE Recommendations

- Weight loss
- Assistive devices
- Bracing
- Corticosteroid injections

CPG Limited Evidence Treatments (Select Patients)

- Manual therapy
- Massage
- Acupuncture
- Laser therapy

Adjunctive Modalities (Limited Evidence)

- Electrical stimulation (TENS / PENS)
- Extracorporeal shockwave therapy
- Percutaneous denervation
- Dry needling

**CPG Limited Evidence:
Orthobiologics and Supplements**

- Platelet-rich plasma (PRP)
- Other orthobiologics
- Dietary supplements
- Select patient benefit

Limited Evidence – Clinical Framing

- Low to moderate evidence
- May be early in development/ need further study
- Generally low harm
- May benefit select patients
- Shared decision-making

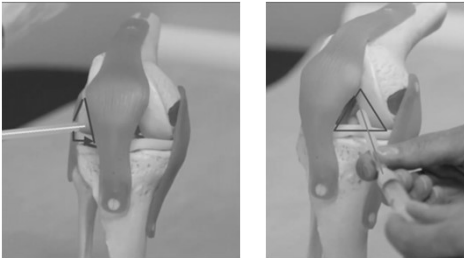
CPG Generally Avoid

- Chronic narcotics incl tramadol
- Arthroscopic debridement
- Routine HA injections

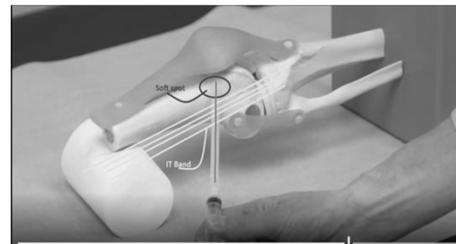
Knee Injection Principles

- Several techniques work
- Use what you know
- Sterility critical
- Patient comfort matters

Lateral and Medial injection sites



Superolateral (Pouch) injection



**Video - Knee injection, Superolateral
(suprapatellar pouch access) Approach**



When to Refer for TKA

- Pain limits daily life
- OA explains symptoms
- Non-op care failed
- Patient engaged and open to surgery

Prerequisites for Considering TKA

- Mental and emotional readiness
- Diagnosis is correct and explains symptoms
- Modifiable Risks identified and mitigated
- Expectations aligned

Medical Optimization Before Referral

- Optimize chronic disease management
- Strongly Encourage smoking cessation
- Address nutrition and weight
- Coordinate care early

Relative Contraindications to TKA

- Minimal pain or dysfunction—Little benefit, poor satisfaction
- Incorrect diagnosis
- Poor engagement- patient not ready
- Unsafe social situation

- Not necessarily permanent contraindications

Modifiable Medical Risk Factors

- Poor diabetes control (Hgb A1c>7)
- Tobacco or nicotine exposure
- Morbid obesity (BMI>40) malnutrition
- Active infection
- Other medical co- morbidities: CAD, PVD, HTN

Largely Unmodifiable Contraindications

- End-stage renal disease on dialysis
- Permanent non-ambulatory state
- Life expectancy less than 6 months

Higher-Risk but Possible Situations

- Long-term immunosuppression
- Active or recent cancer treatment
- Sickle cell disease
- Cognitive or mental health differences
- Prior joint infection

Role of Primary Care

- The Heavy Lifting
- Longitudinal OA management
- Risk factor modification
- Expectation-setting
- Appropriate referral

Transitional Q & A



Non-Arthroplasty Treatments for Knee Osteoarthritis

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The Importance of Joint Preservation

- Why should we care about joint preservation?
 - The Burden of Osteoarthritis (OA)
 - Patient Apprehension
 - The Potential for Regeneration

The Importance of Joint Preservation

- Why should we care about joint preservation?
 - The Burden of Osteoarthritis (OA)
 - High expenditures on patient care, time lost from work
 - Worsens co-morbid conditions and increases risk of heart disease
 - 55% increase in all-cause mortality**

The Burden Of Osteoarthritis: A Serious Disease

242 Million people worldwide have symptomatic and activity-limiting OA of the hip and/or the knee. OA pain affects sleep quality, mood, and participating in everyday life.

The Cost Of Osteoarthritis

- Prevalence Costs: \$10.3 billion
- Total Costs: \$136 billion
- Average Direct Costs: \$11,000 person/year

Indirect costs are \$17 billion (i.e. lost earnings). Direct costs are \$65 billion (i.e. medical expenditures). 37 most rapidly rising condition associated with disability, just behind diabetes and dementia.

A third of people with OA have 5 or more chronic conditions. OA significantly limits a person's ability to self-manage other conditions, such as diabetes and hypertension. OA increases the risk of developing heart disease by 50%.

55% Increase in all cause mortality as a result of reduced levels of physical activity, comorbid conditions, and adverse effects of medications.

The Importance of Joint Preservation

- Why should we care about joint preservation?
 - Patient Apprehension
 - Fear of infection, VTE, poor outcome
 - The Potential for Regeneration
 - OA has traditionally been understood as progressive, with no disease-modifying treatments
 - New(er) treatments show disease-modifying potential that may further prolong the lifespan of the native joint

OA Pathophysiology and Therapeutic Targets

- OA is a multi-factorial degenerative condition with hallmark pathophysiologic features
 - Loss of synovial fluid viscoelasticity
 - Cartilage degradation (apoptosis)
 - Subchondral bone resorption (manifested as bone marrow edema and insufficiency fractures)
 - Synovitis
 - Mechanical axis deviations (malalignment and flexion contractures)

OA Pathophysiology and Therapeutic Targets

Osteoarthritis: Pathophysiologic Hallmarks & Solutions	
Pathophysiologic Hallmarks	Therapeutic Solutions
Loss of Viscoelasticity	Viscosupplementation with Hyaluronate
Loss of Cartilage	PRP: Reduces Chondrocyte Apoptosis & Promotes Chondrocyte Proliferation
Synovitis	PRP: Reduces Synovitis & Inflammation
Bone Marrow Edema	PRP & Shockwave Therapy
Malalignment	Implantable Mechanical Supports

Viscosupplement

- Approved by the FDA in 1997, this is either avian-derived or fermented (from streptococcus metabolism) and then purified for patient application
- Although widely used, it came under scrutiny when the AAOS recommended against it in its 2013 clinical practice guideline
- That posed a major issue for patients
- This left steroid as the only other injectable therapy
 - Known to be chondrotoxic and is associated with increased rate of knee arthroplasty
- This created a significant need for knee OA injectables

An Opportunistic Window

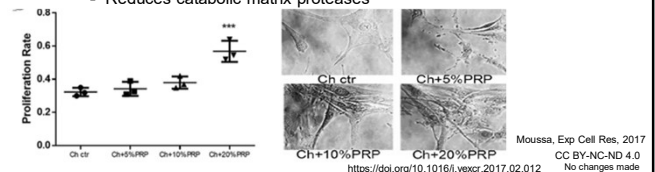
- Given the increased scrutiny on viscosupplement products, this created a new gap in treatment options
- At the same time, cohort studies were being published from Italy on positive patient outcomes after treatment with platelet-rich plasma (PRP)
- Originally used to aid in sternal wound closures and gum grafting, PRP was applied to OA
 - Rationale: OA is a non-healing condition, so a boost of growth factors may improve symptoms and / or disease progression

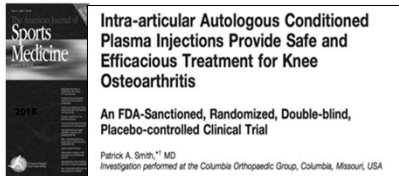
PRP: Mechanistic Insight

- Complex interplay of tissue healing, anti-inflammatory, and anti-catabolic effects
- When platelets contact damaged collagen, they degranulate and release their anabolic 'cargo'
 - Growth factors (TGF, IGF, FGF, VEGF, etc)
 - Exosomes
- These GF and proteins then exert a paracrine effect and incorporate into cells (like synoviocytes)
 - Alters phenotypic expression to one more resilient in the face of chronic disease

Mechanisms of Tissue Healing

- PRP results in dose-dependent effects
 - Anabolic
 - Promotes chondrocyte proliferation
 - Increases vacuole formation as a marker of autophagy, promoting balanced cellular turnover
 - Anti-catabolic
 - Reduces arthritic chondrocyte apoptosis
 - Reduces catabolic matrix proteases



ACP

Confirmed safety and efficacy of ACP with statistical and clinical significance up to 1 year

Mechanisms of Tissue Healing

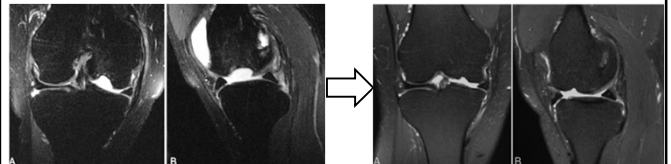
- These cellular level changes have potential to cause macroscopic improvement
- Yoshioka et al completed a RCT comparing 3 LP-PRP vs saline
- MRI improvement seen by 24 weeks

Yoshioka, AJSM, 2024

Mechanisms of Tissue Healing

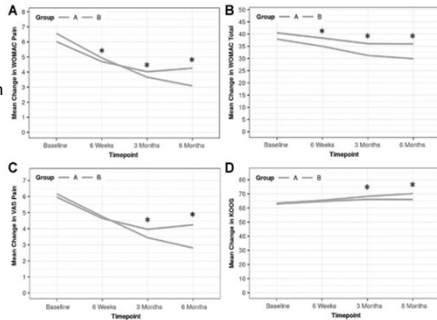
- Chu et al randomized over 600 knee OA patients to either a series of 3 LP-PRP or saline
- After only 1 treatment session (series of 3 weekly injections), the LP-PRP group had less cartilage loss compared to placebo at 5 years
- Take home point: A single treatment changed the disease course over 5 years

Chu, KSSTA, 2022

Clinical Observations of Structural Healing

Dosing Primer

- Patel et al performed a dose comparison study
 - 5.6 billion vs 2.8 billion platelets (LP-PRP)
 - PRP concentrated approximately 3.5x over baseline
 - 5.6 billion superior as an initial dose



Patel et al. OJSM, 2024 doi: 10.1177/23259671241227863 CC BY-NC-ND 4.0 No changes made

Dosing Primer

- Gobbi et al demonstrated that a proactive 'booster' cycle at 1 year promotes further symptom relief compared to a single series without repeat administration

Gobbi, KSSTA, 2014

Autologous "Stem Cell" Therapies

- Both bone marrow and adipose are known sources of mesenchymal stem cells (MSC)
 - Relative concentration of MSCs in both cell populations is quite low
 - Stem cells are unlikely to be the primary driver of effect
- Despite the popularity due to stem cell content, there are no studies showing either are superior to either placebo or PRP
- More invasive
- More costly

Off-the-Shelf "Stem Cell" Therapies

- Usually derived from birth tissue
 - Umbilical cord
 - Placenta
 - Amniotic fluid
- Studies have shown these products have no stem cell content
 - Mainly composed of hyaluronate
- None are FDA approved
- Substantial harm has occurred
 - Sepsis / septic arthritis
 - Reactive arthritis in HLA-B27+ patients
- Not indicated for any clinical use
 - Off-label use doesn't apply since this is allogeneic

Shockwave Therapy

- High-energy acoustic waves
- In-office administration
- Therapeutic effects
 - Promotes angiogenesis
 - Stimulates tissue regeneration
 - Reduces pain signaling
- Many RCTs demonstrate superiority for knee OA over control therapies (including physiotherapy and alendronate)



On The Horizon

- Several biologics are in the developmental pipeline and worth monitoring
- Autologous
 - Stromal vascular fraction (SVF)
 - Adipose is harvested in a procedure room similar to a low-volume liposuction
 - Collagenase is mixed with the tissue to release MSCs from the collagen matrix
 - Tissue is centrifuged to concentrate a higher amount of true stem cells

On The Horizon

- SVF
 - Passed Phase 2 FDA trials
 - Successfully complete phase 3 FDA trials
 - Awaiting regulatory review

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Article - Knee

Clinical Efficacy of Intra-articular Mesenchymal Stromal Cells for the Treatment of Knee Osteoarthritis: A Double-Blinded Prospective Randomized Controlled Clinical Trial

Jaime R. Garza, MD¹, Richard E. Campbell, BS¹, Fotios P. Tjonnakeris, MD¹, Kevin B. Freedman, MD¹, Lawrence S. Miller, MD², Daniel Santa Maria, MD¹, and Bradford S. Tucker, MD^{1,3}

On The Horizon

- Platelet-derived exosomes
 - Exosomes are small extracellular vesicles (sEV) found in blood cells and plasma
 - Consist of proteins, lipids, and nucleic acids (i.e. miRNA) that deliver cell signaling information
 - Promotes tissue restoration and homeostasis

On The Horizon

- Exosomes; Two primary applications
 - Exosomes as the therapy
 - When derived from platelets, carry the same therapeutic information as PRP
 - Purified
 - Consistent dosing
 - Off-the-shelf (stable shelf life for years, eliminates need to draw / spin blood in the clinic)
 - Exosomes as the delivery vehicle
 - Exosomes can be loaded with another therapeutic drug / biologic
 - Lipid bilayer protects the therapeutic content
 - Improves targeted delivery

On The Horizon

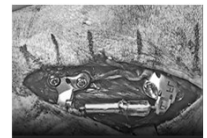
- Platelet-derived exosomes
 - Currently in phase I studies for
 - Knee OA
 - Radiation and fistulizing wounds
 - Myocardial infarction
 - In phase 2 for
 - Diabetic foot ulcers

Surgical Innovations for Knee OA

- Bracing intuitively helps patients
- Address malalignment
- Improves muscle contractility
- However, patients cannot brace 24/7
- Two surgical, non-arthroplasty treatments are worth considering

Surgical Innovations for Knee OA

- Medial Implantable Shock Absorber (MISHA)
- For medial knee OA
- Reduces impact by 30%
- In the Phantom Trial, 96% of patients had an improvement of 20% or more on the WOMAC scale
 - 4% removal rate
- Native joint preserved, OA injectables still an option



Surgical Innovations for Knee OA

- Osteotomy
 - High tibial osteotomy for medial pain / varus knees
 - Distal femoral osteotomy for lateral pain / valgus knees
 - Corrects malalignment with no implants and no replacement
 - Native joint is preserved
 - OA injectables can still be performed



Summary

- PRP has excellent safety and efficacy track record for over 10 years
- New biologic injectables may offer additional injection based care
 - Autologous and off-the-shelf options in FDA pathway
- Joint sparing surgical techniques (MISHA / HTO) may offer additional benefit when rehabilitation and injectable options have yielded suboptimal results