



Definition

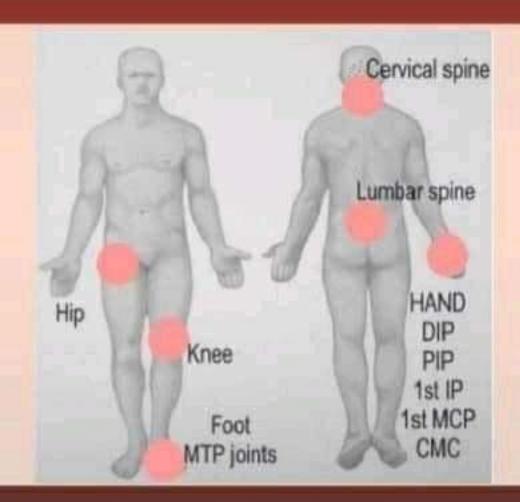
- Osteoarthritis (OA), which is also known as osteoarthrosis or degenerative joint disease (DJD):
- ➤ is a progressive disorder of the joints caused by gradual loss of cartilage and resulting in the development of bony spurs and cysts at the margins of the joints.



Epidemiology

- ➤ OA is the most common form of arthritis and the most common joint disease
- Characterized by degeneration of articular cartilage
- Leads to fibrillation, fissures, gross ulceration and finally disappearance of the full thickness of articular cartilage







Joint Protective Mechanisms

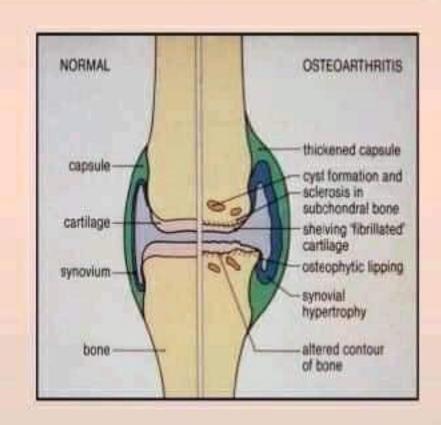
- √ Joint Capsule
- √ Ligaments
- ✓ Muscle
- √ Sensory afferents
- ✓ Underlying bone

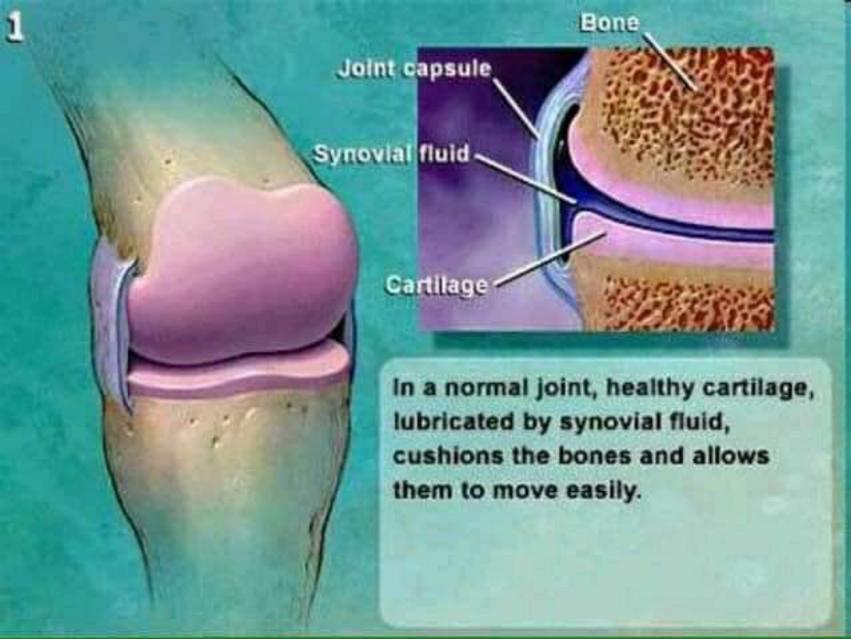


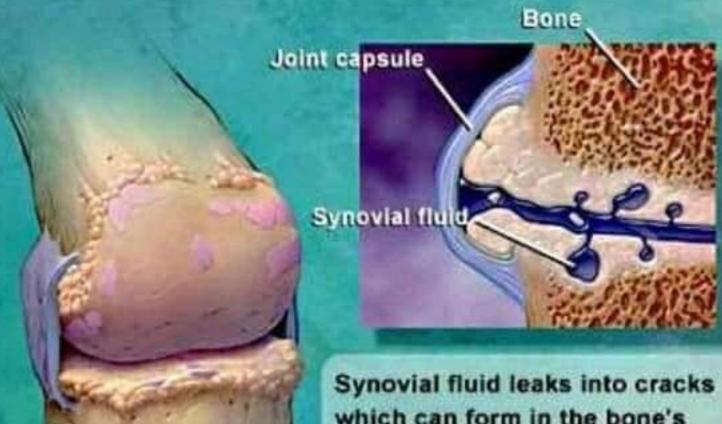
OA is primarily disease of cartilage.

II-1 is a potent proinflammatory cytokine, which capable of inducing chondrocytes and synovial cells to synthesize MMPs. MMPs is responsible of degradation of articular cartilage.

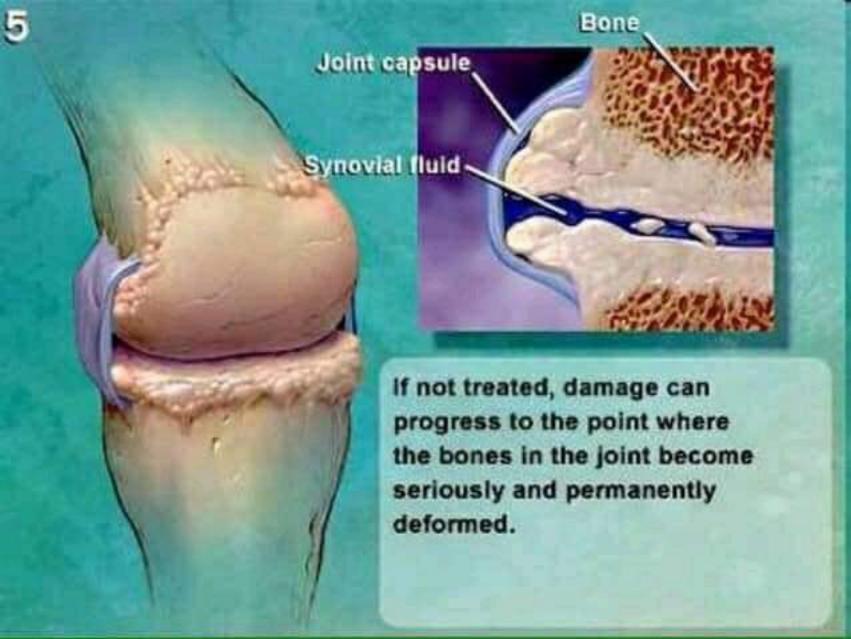
In addition, IL-1 inhibits synthesis of collagen II, proteoglycans and growth factor B stimulated chondrocyte proliferation







Synovial fluid leaks into cracks which can form in the bone's surface when this replacement cartilage wears away. This causes further damage and in some cases can lead to cysts in the bone or other deformities.





Age

- Age is the most potent risk factor for OA
- OA is rare in under age 40
- OA occurs in more than 50% of persons over age 70





Female gender

In general, arthritis occurs more frequently in women than in men. Before age 45, OA occurs more frequently in men; after age 45, OA is more common in women. OA of the hand is particularly common among women.





Joint's Abnormalities

People with joints that move or fit together incorrectly, such as bow legs, a dislocated hip, or doublejointedness, are more likely to develop OA in those joints.





Obesity

Being overweight during midlife or the later years is among the strongest risk factors for OA of the knee





Joint overuse or injury

Traumatic injury (ex. Ligament or meniscal tears) to the knee or hip increases your risk for developing OA in these joints. Joints that are used repeatedly in certain jobs may be more likely to develop OA because of injury or overuse





Sources of pain

- Increase hemostatic pressure within bone rise in OA
- ➢ Bone marrow edema
- ➤ Synovial inflammation
- ➤ Joint effusions
- ➤ Osteophytes

Symptoms & Signs

- ✓ Pain increases with activity and worse at night.
- ✓ Morning Stiffness less than 30 mins
- √ Swelling
- ✓ Deformities
- √ Joint instability
- √ Loss of function
- ✓ Neurological signs if spine Involved.
- ✓ Crepitus
- ✓ Muscle wasting



Diagnosis – Blood Tests

No Blood tests are routinely indicated for workup of patients with OA unless symptoms & signs suggest inflammatory arthritis.



Diagnosis – Synovial Fluid

Examination of the synovial fluid is more helpful diagnostically than an X-Ray.

❖ Synovial Fluid in OA:

>WBC<1000/mm3

➤ Clear color

▶ High Viscosity



Diagnosis – Radiography

X-Rays are indicated to evaluate chronic hand pain and hip pain thought to be due to OA

For knee pain, X-Ray should be obtained if symptoms or signs are not typical of OA or Pain persists after inauguration of effective treatment



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Radiographic Features

- > Erosion
- ➤ Bone Cyst
- ➤ Subchondral Sclerosis
- ➤ Osteophytes
- ➤ Narrow Joint Space



Management of OA

Treatment of OA:

- ▶ Pharmacotherapy
- ➤ Non Pharmacotherapy
- Patients with mild and intermittent symptoms may need only Non Pharmacotherapy
- Patients with ongoing, disabling pain are likely to need both Non Pharmacotherapy and Pharmacotherapy



Management of OA

Treatment of OA:

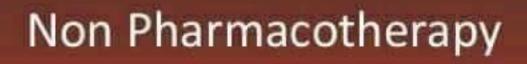
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Pharmacotherapy

Treatment	Dosage	Comments
Acetaminophen	Up to 1 g tid	Prolongs half-life of warfarin. Make sure patient is not taking other treatments containing acetaminophen to avoid hepatic toxicity.
Oral NSAIDs and COK-2 inhibitors Naproxen Salsalate Ibuprofen	375-500 mg bid 1500 mg bid 600-800 mg 3-4 times a day	Take with food, increased risk of myocardial infanction and stroke for some NSAIDs and especially COX-2 inhibitors. High rates of gastrointestinal side effects, including ulcess and bleeding, occur. Patients at high risk for gastrointestinal side effects should also take either a proton pump inhibitor or misoprostol." There is an increase in gastrointestinal side effects or bleeding when taken with acetylsalicylic acid. Can also cause edema and renal insufficiency.
Topical NSAIDs Dictoferrac Na 1% get	4 g and (for knees, hands)	Rub onto joint. Few systemic side effects. Skin irritation common.
Opiates	Various	Common side effects include dizziness, sedation, nausea or vomiting, dry mouth, constipa- tion, urinary retention, and praritus. Respiratory and central nervous system depression can occur.
Capsaidn	0.025-0.075% cream 3-4 times a day	Can irritate mucous membranes.
Intraarticular injections		
Steroids		
Hyaluronans	Varies from 3–5 weekly injections depending on preparation	Mild to moderate pain at injection site. Controversy exists regarding efficacy.

Patients at high risk include those with previous gentromeratinal events, persons 260 years, and persons taking glucocorrocads. Trials have shown the efficacy of proton pump whitefors and misoprostol in the prevention of silvers and bleeding. Misoprostol is essociated with a high rate of diamnes and cramping therefore, proton pump inhibitors are more widely used to reduce NSAID related gestro-mentional symptoms.



Avoid unnecessary stress Increase muscle strength Reduce Load on the Non Pharmacotherapy joint Use of walking stick Exercise Weight loss Correction of Malalignment



Thank you