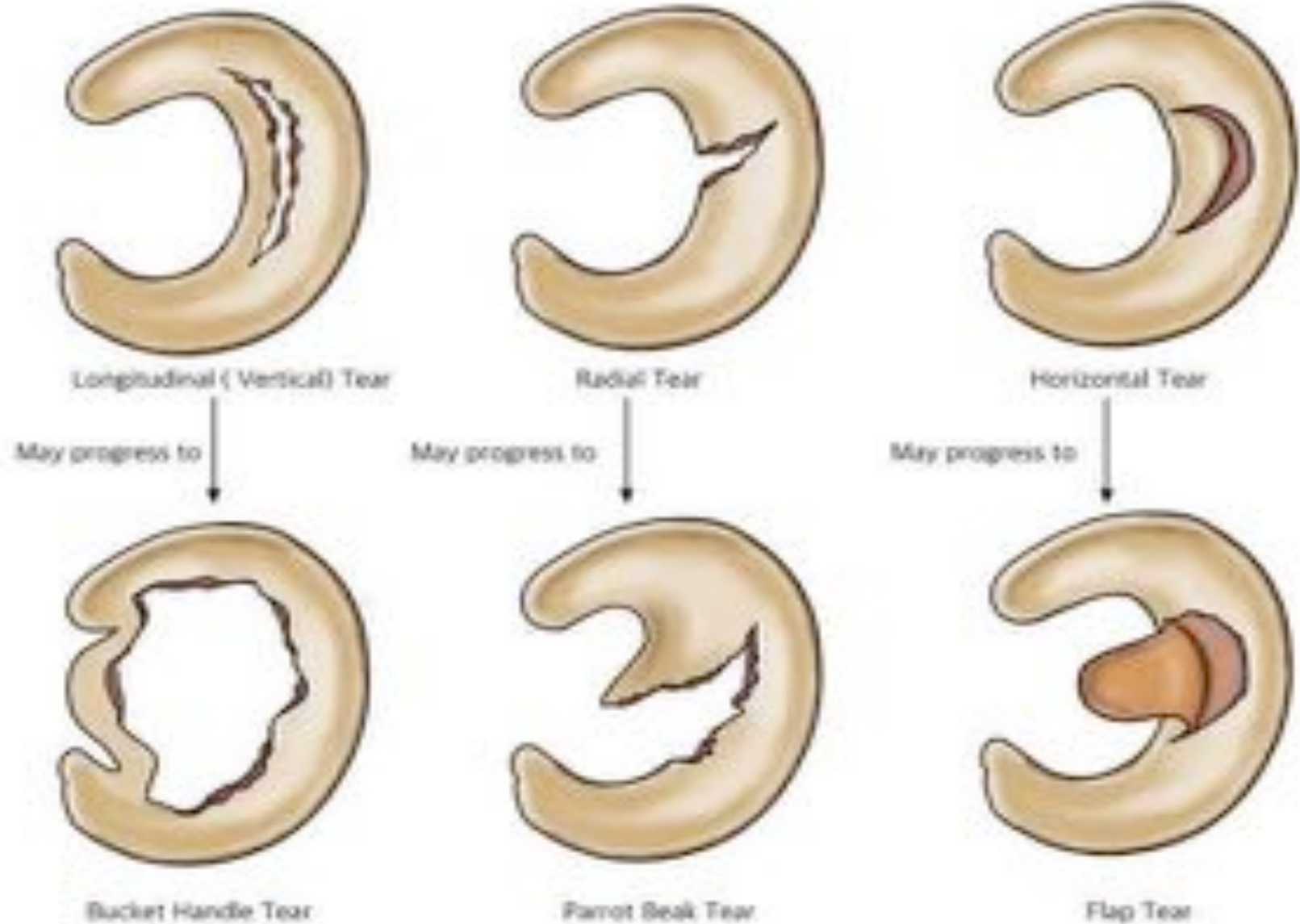
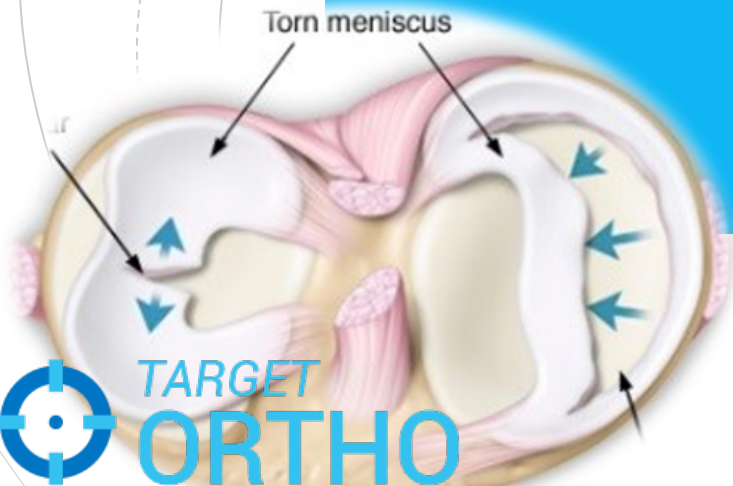
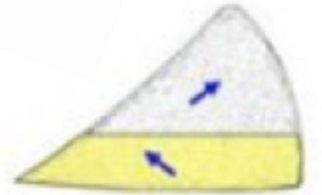
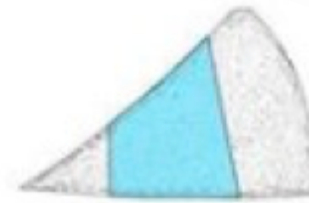
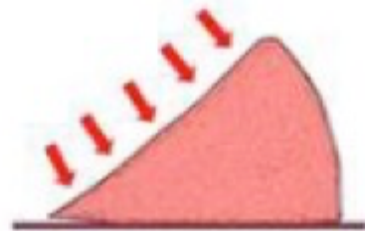


TEAR PATTERNS







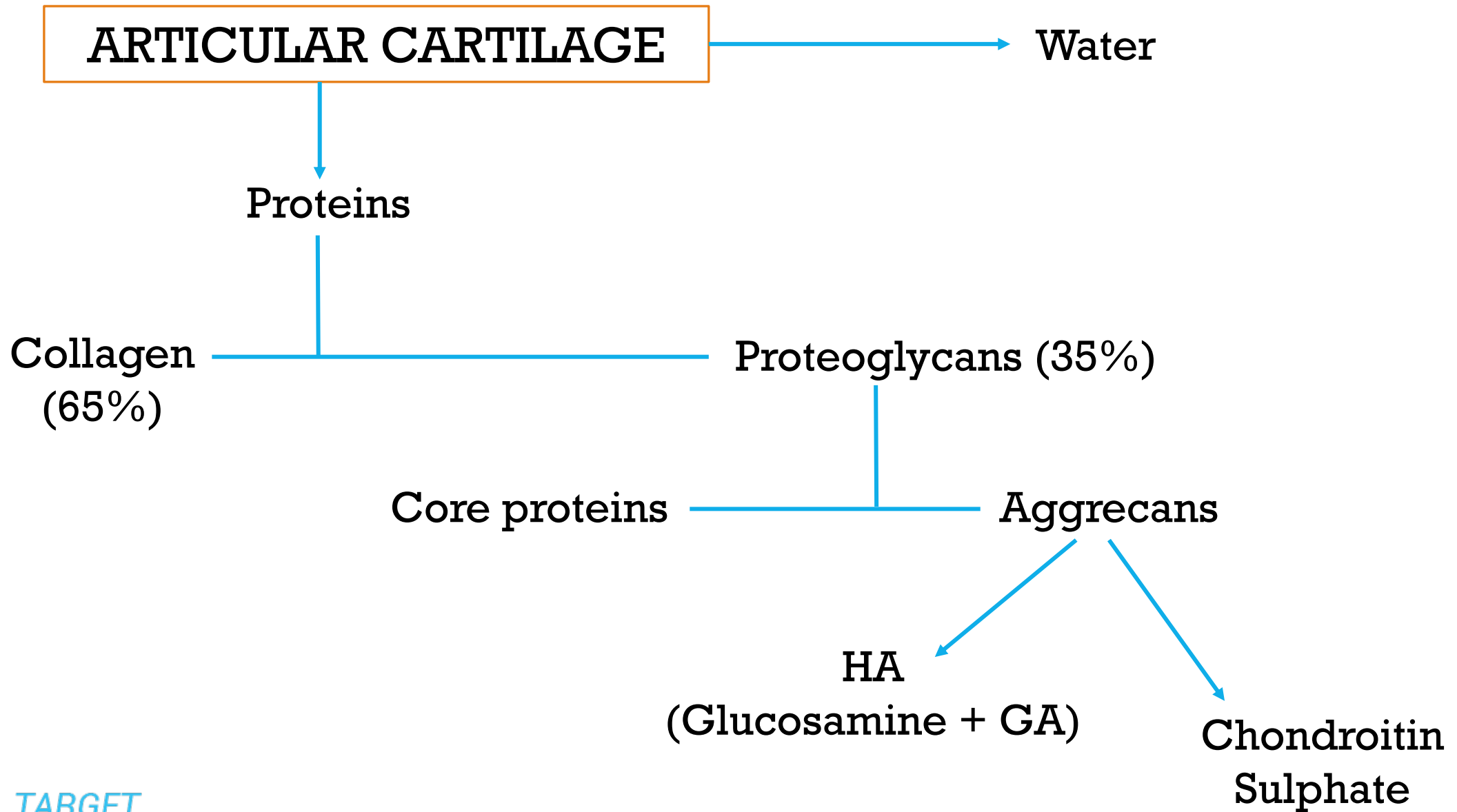
FOCAL ARTICULAR CARTILAGE LESIONS

of Knee

**MUKUL
MOHINDRA**



M.S [Ortho], DNB, MNAMS
Dip. SICOT [Belgium]
FNB [Sports Medicine]
Fellowship in MIA, Athens [SICOT]



Q. The predominant collagen type present in Articular cartilage?

- A. Type I
- B. Type II
- C. Type III
- D. Type IV

Q. The predominant collagen type present in Meniscus?

- A. Type I
- B. Type II
- C. Type III
- D. Type IV

PRE-DISPOSING FACTORS FOR CARTILAGE LESIONS

The primary factor in the development of a cartilage lesion is undoubtedly the relationship between the size of the lesion and the load surface, being adversely affected by:

- Obesity
- Age
- Axial misalignment
- Family history of osteoarthritis
- Overload activities



Obesity



Aging



Mechanical Stress



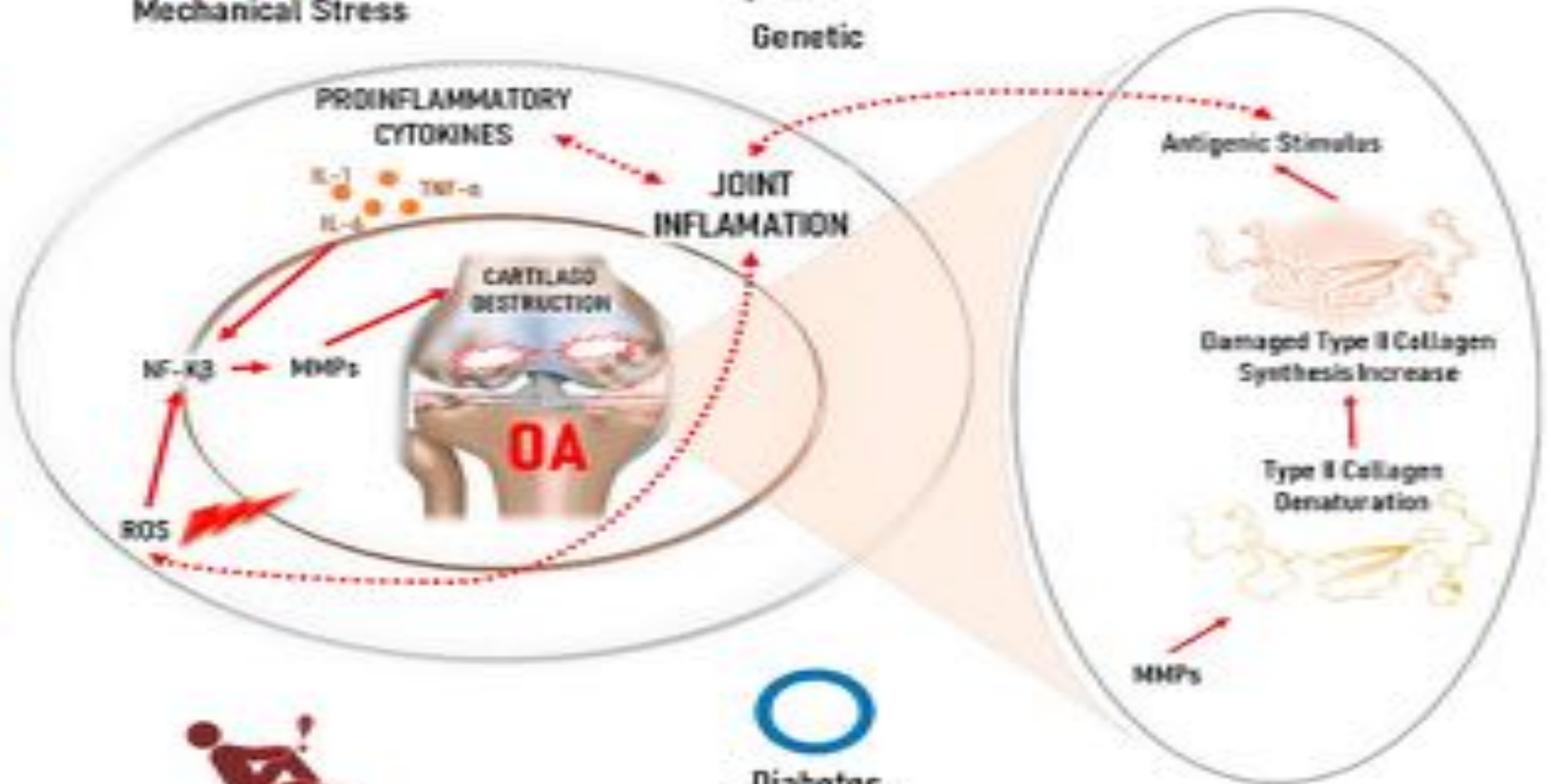
Genetic



Injury



Diabetes



CLASSIFICATION OF ARTICULAR CARTILAGE DEFECTS

Several classifications : Outerbridge, ICRS etc

Outerbridge classification:

Grade 0 - Normal cartilage

Grade 1 - Cartilage softened and swollen

Grade 2 - Cracking not reaching the subchondral bone; less than 1.5 cm

Grade 3 - Cracking reaching the subchondral bone without exposure; greater than 1.5 cm

Grade 4 - subchondral bone exposure of any diameter

ICRS Classification:

Normal: grade 0

Almost normal:

Grade 1a- superficial lesions / softening

Grade 1b - 1a and / or fissures or surface cracks

Abnormal:

Grade 2 - length < 50% thickness

Serious injury:

Grade 3 a - extension > 50%

Grade 3 b - to the calcified layer

Grade 3 c - to the surface of the subchondral bone (without entering)

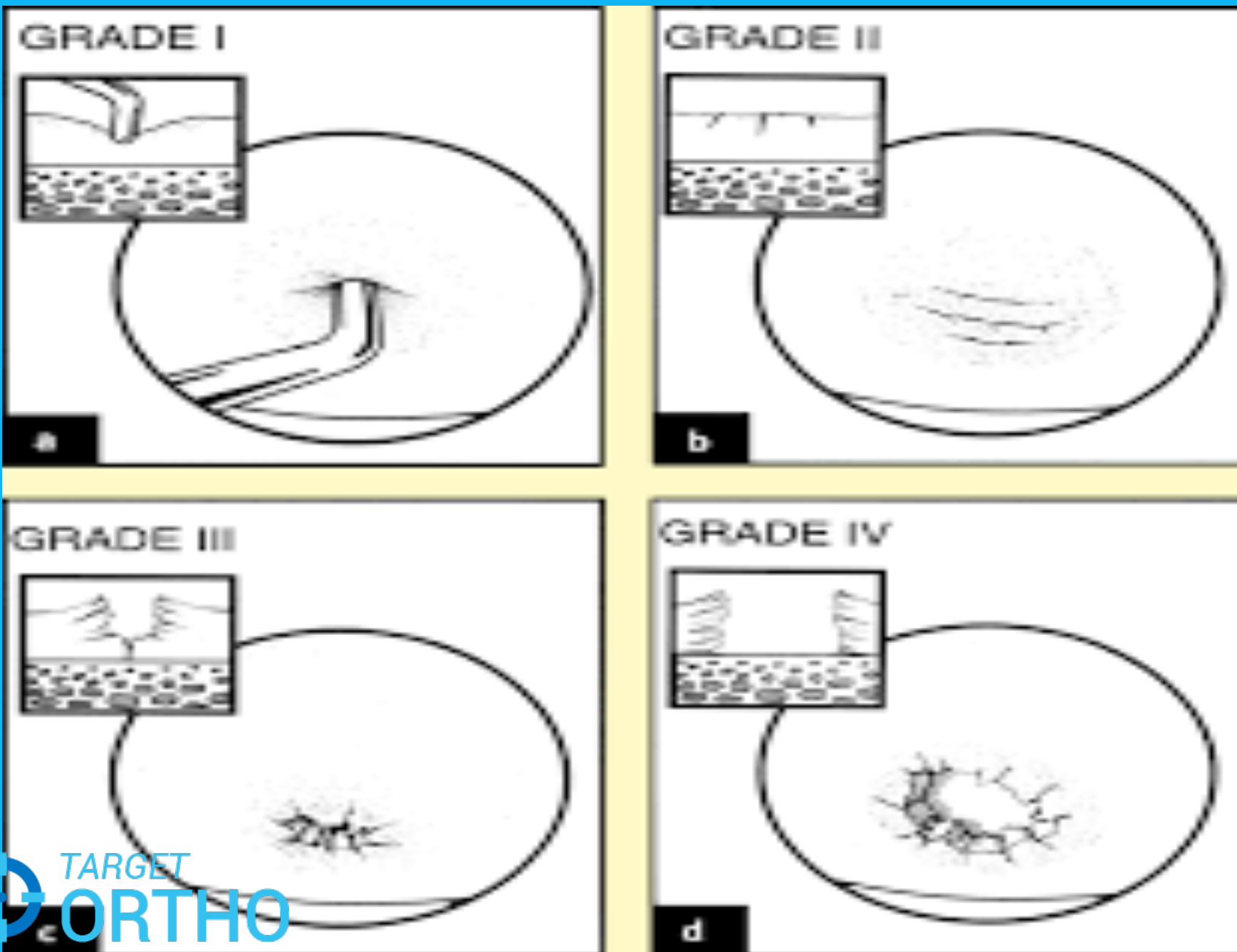
Grade 3 d - includes blisters

Very serious injury:

Grade 4 a - penetration of the subchondral bone but not the full diameter of the defect

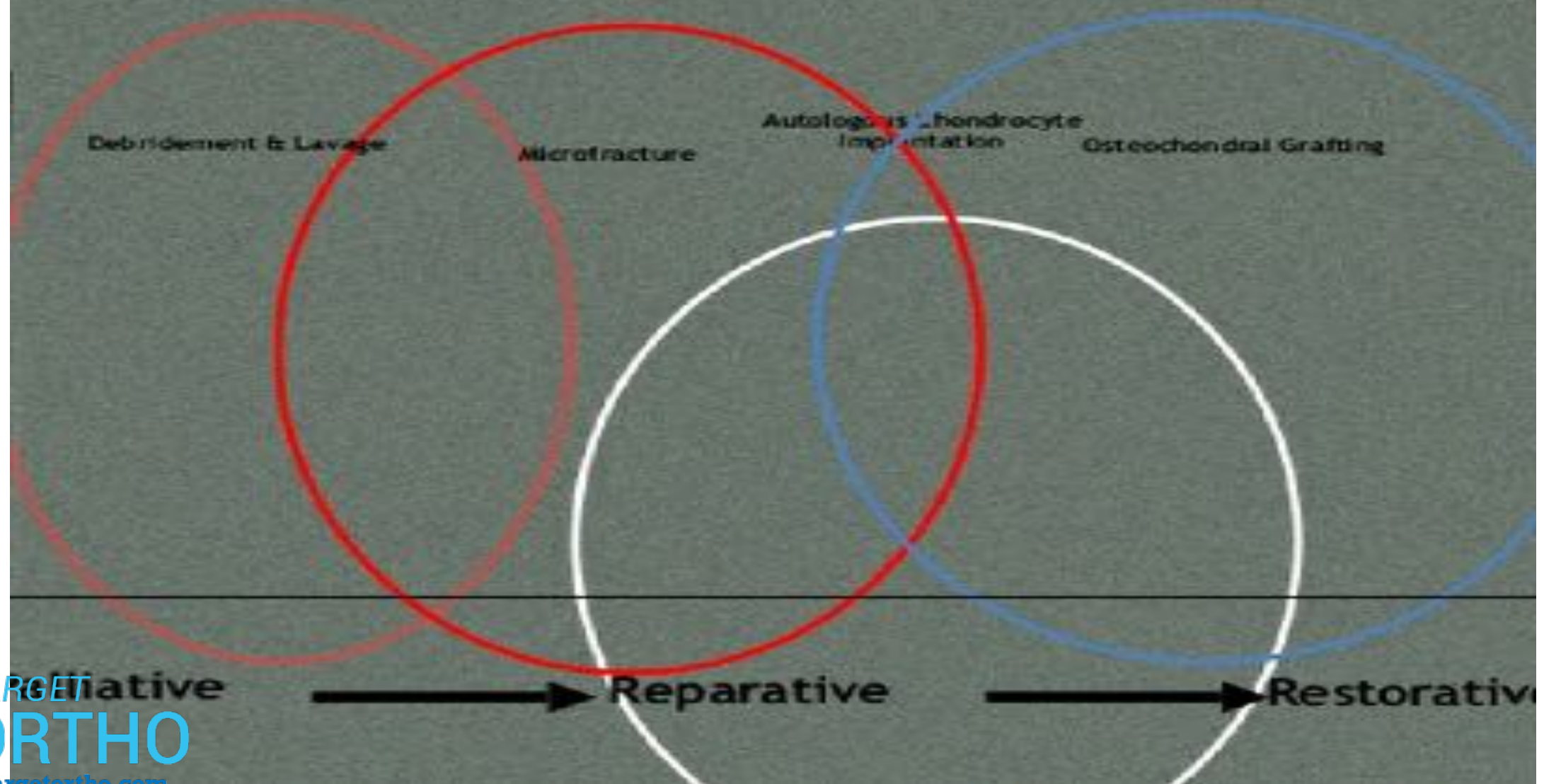
Grade 4b - penetration across the diameter of the defect

OUTBRIDGE CLASSIFICATION

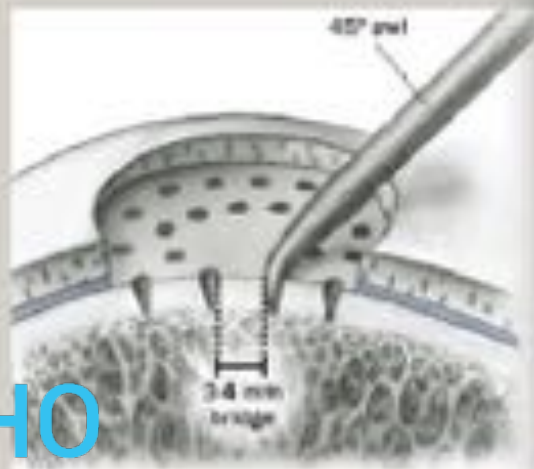
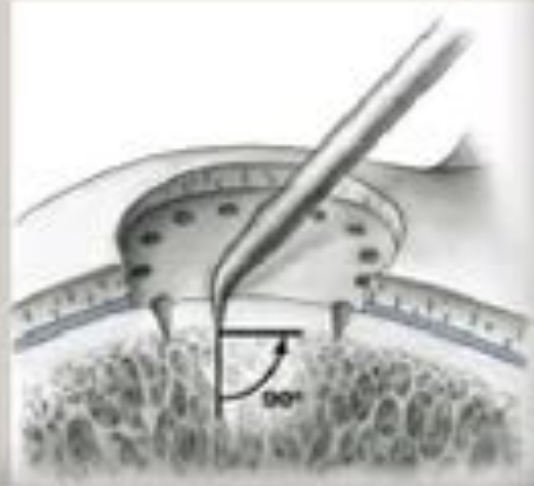
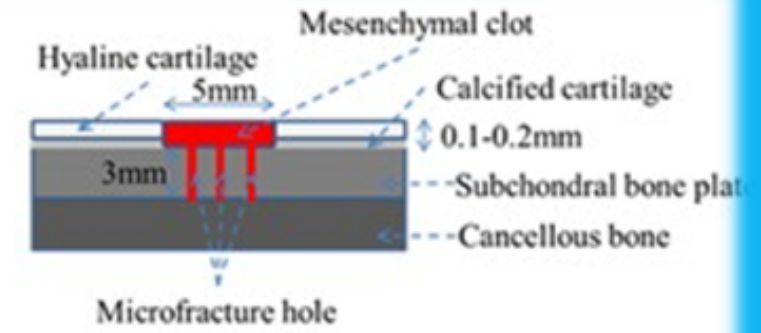


TREATMENT OPTIONS

Treatment Options for the Cartilage Bio-surgeon in 2009



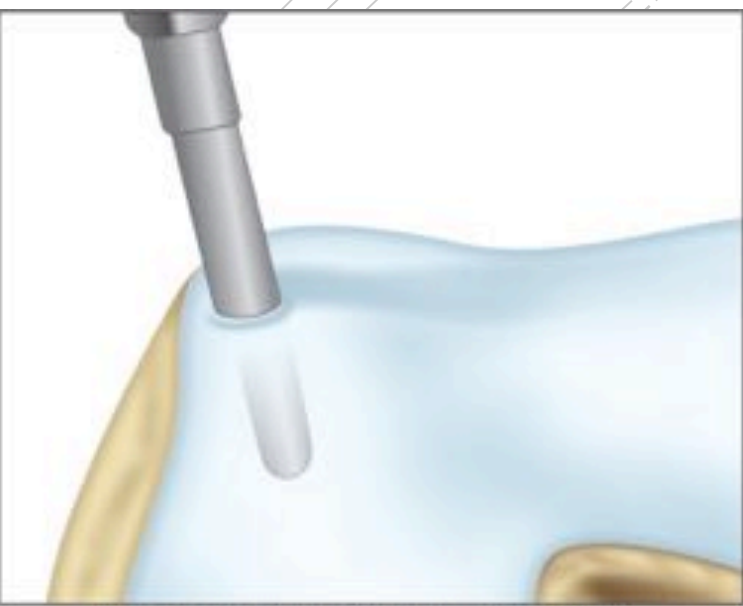
MICROFRACTURE



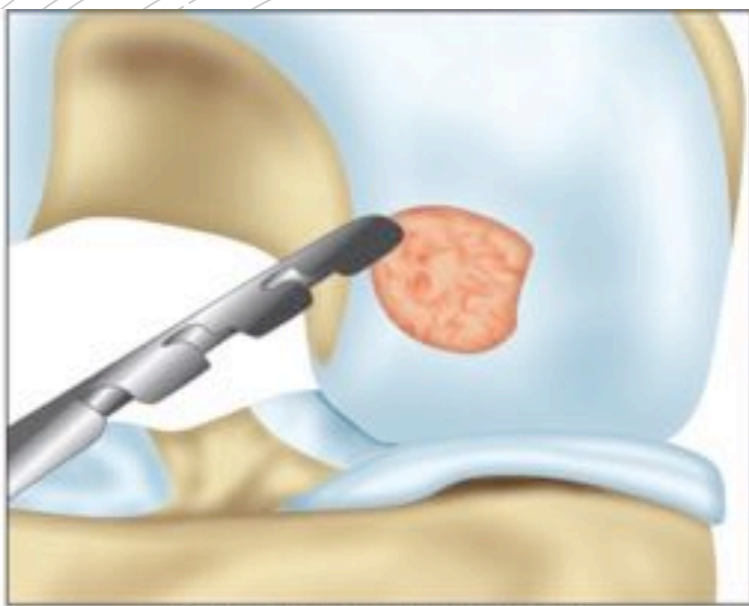


OATS

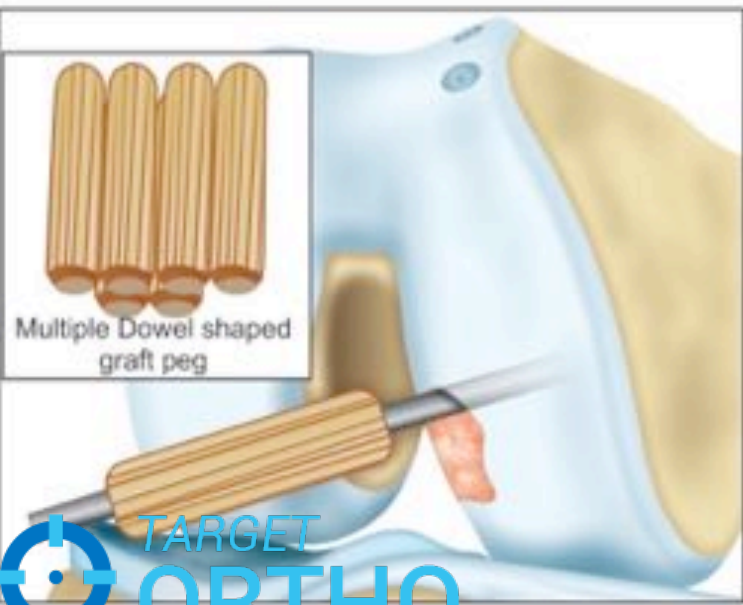
Osteochondral Autologous
Chondrocyte Transplantation
Surgery



Graft are taken from a nonweight bearing part of the cartilage

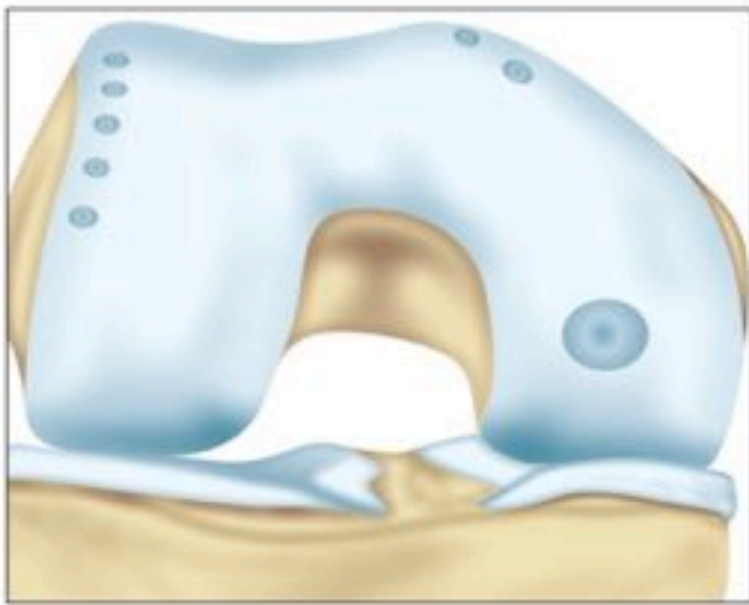


Holes drilled at defect site



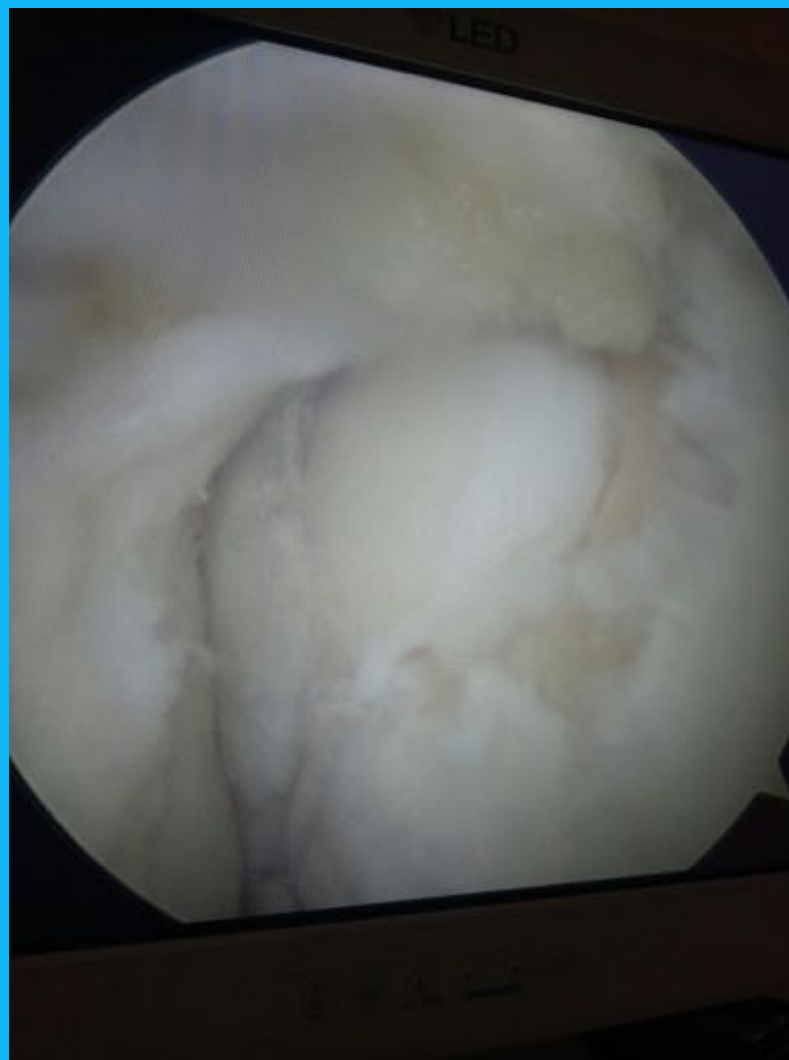
Multiple Dowel shaped graft peg

Dowel-shaped graft peg inserted at hole at chondral lesion site

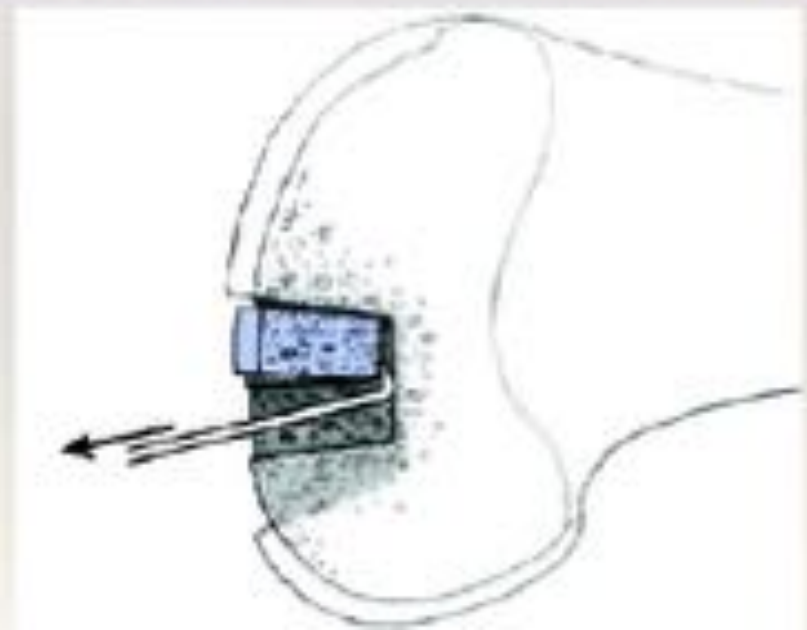


Lesion and donor sites





- Cylinder perpendicular to articular surface
- Avoid the cylinder sinking; 1 - 2 mm above the adjacent surface
- No load for 3 - 5 weeks



Osteochondral allograft

Restrictions

- Obtainment difficulties
- Rejection
- Difficulty in graft incorporation
- Diseases transmission
- Potential high cost
- Technical difficulty in graft conditioning



de Oliveira
ondonal do Joel
/ F. Medicina d

ACI

AUTOLOGOUS CHONDROCYTE IMPLANTATION

GENERATIONS OF ACI

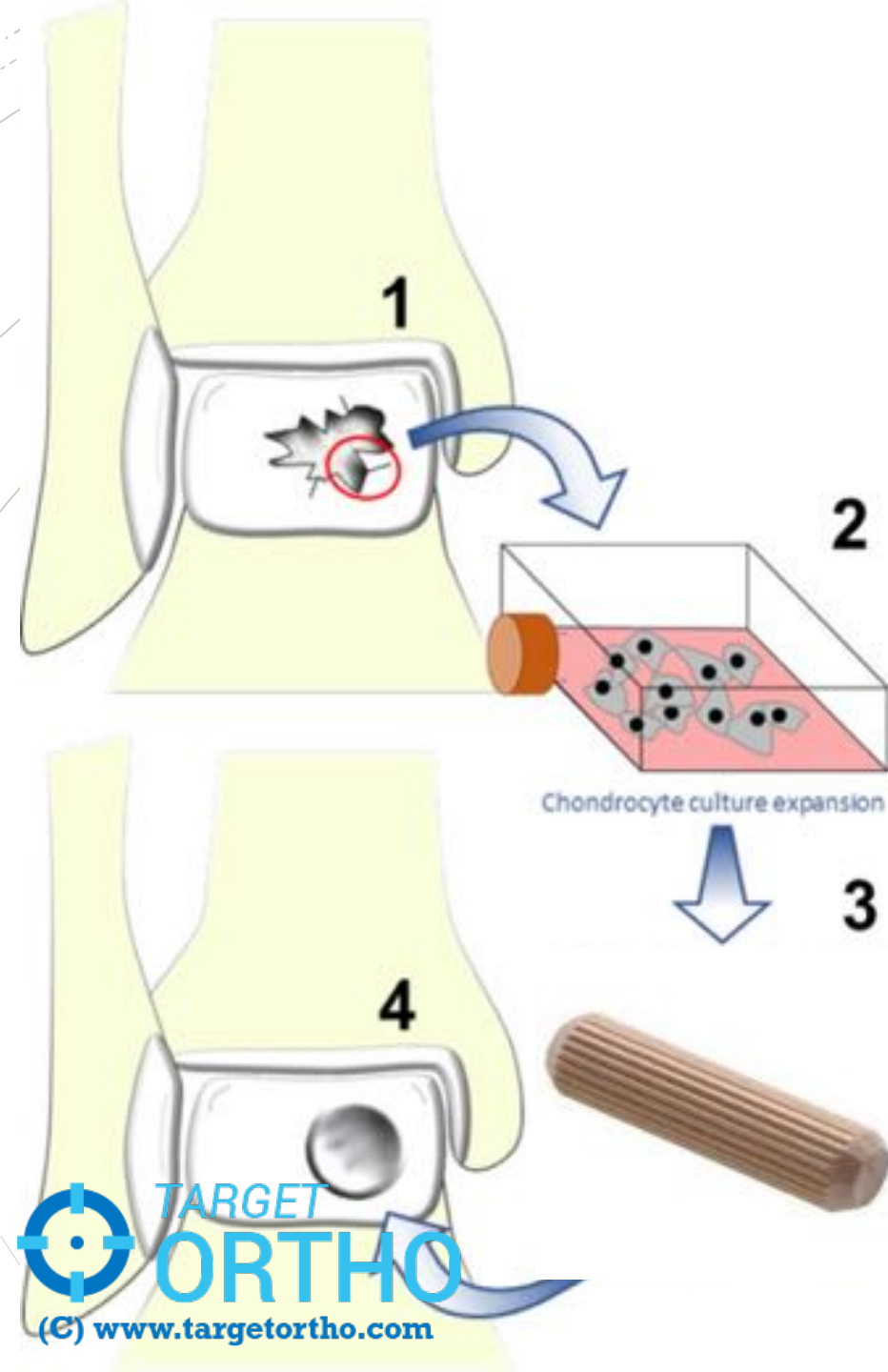
First generation ACI

Autologous chondrocytes were grown in culture, made into paste which was then injected into the defect after 3-6 weeks beneath a “periosteal flap” or a “collagen membrane” e.g. MACI



Second generation ACI

Grown chondrocytes were mounted into a **3D SCAFFOLD** and the scaffold was **hammered press fit** into the defect e.g. hyalograft C, bioseed- C



Third generation ACI

The matrix with chondrocytes is subjected to mechanical stimulation, with hydrostatic pressure to chondrocytes for a minimum of seven days which will increase the production of collagen type II, aggrecan and other normal components of hyaline cartilage

Easy application and adaptation to different sizes and shapes of defects

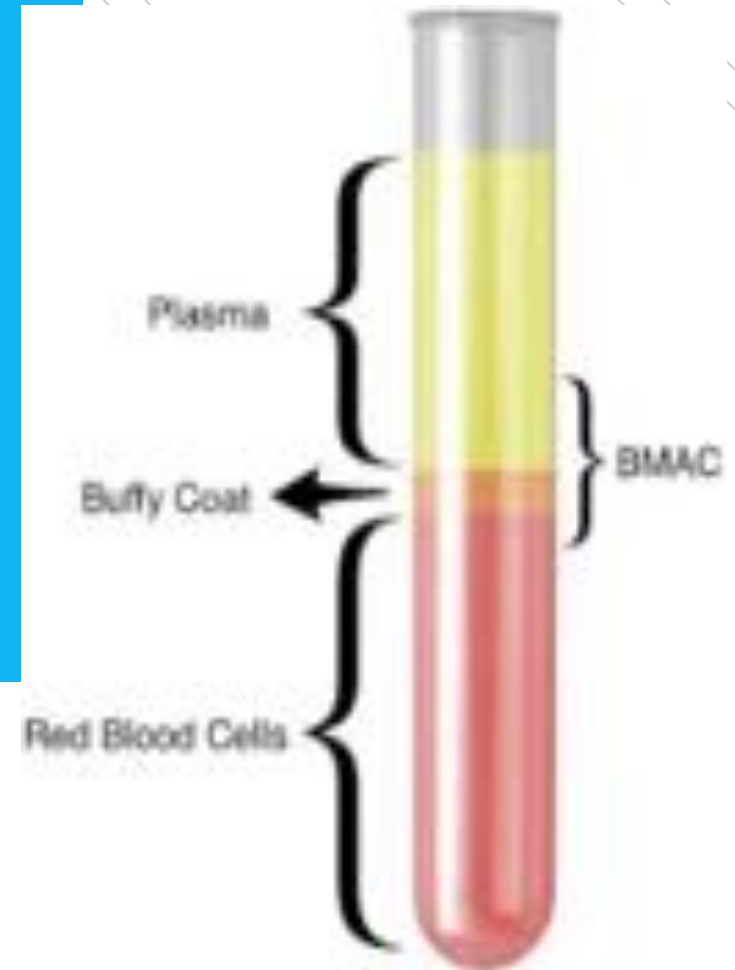
MODIFIED

Third generation ACI

- Latest modification in 3rd generation ACI is **CARTILAGE AUTOGRAFT IMPLANTATION SYSTEM (CAIS)**.
- Articular cartilage after harvest is **MINCED** into small fragments encapsulated directly **IMPLANTED** into scaffold (w/o culture expansion).
- Chondrocytes from fragments migrate into scaffold (made of polycaprolactone+PGA +PDS) and produce ECM.
- A minimal amount of chondrocytes produce much of tissue for a relatively larger defect.

BMAC

Bone marrow aspirate
stem cell concentrate



OUTERBRIDGE GRADE

Grade I and II

*Debridement, Lavage,
Chondroplasty*

Grade III/ IV

Activity level
LOW

Microfracture

Activity level GOOD

Size of the Defect

< 2.5 cms

*OATS
(autograft)*

2.5 -5 cms

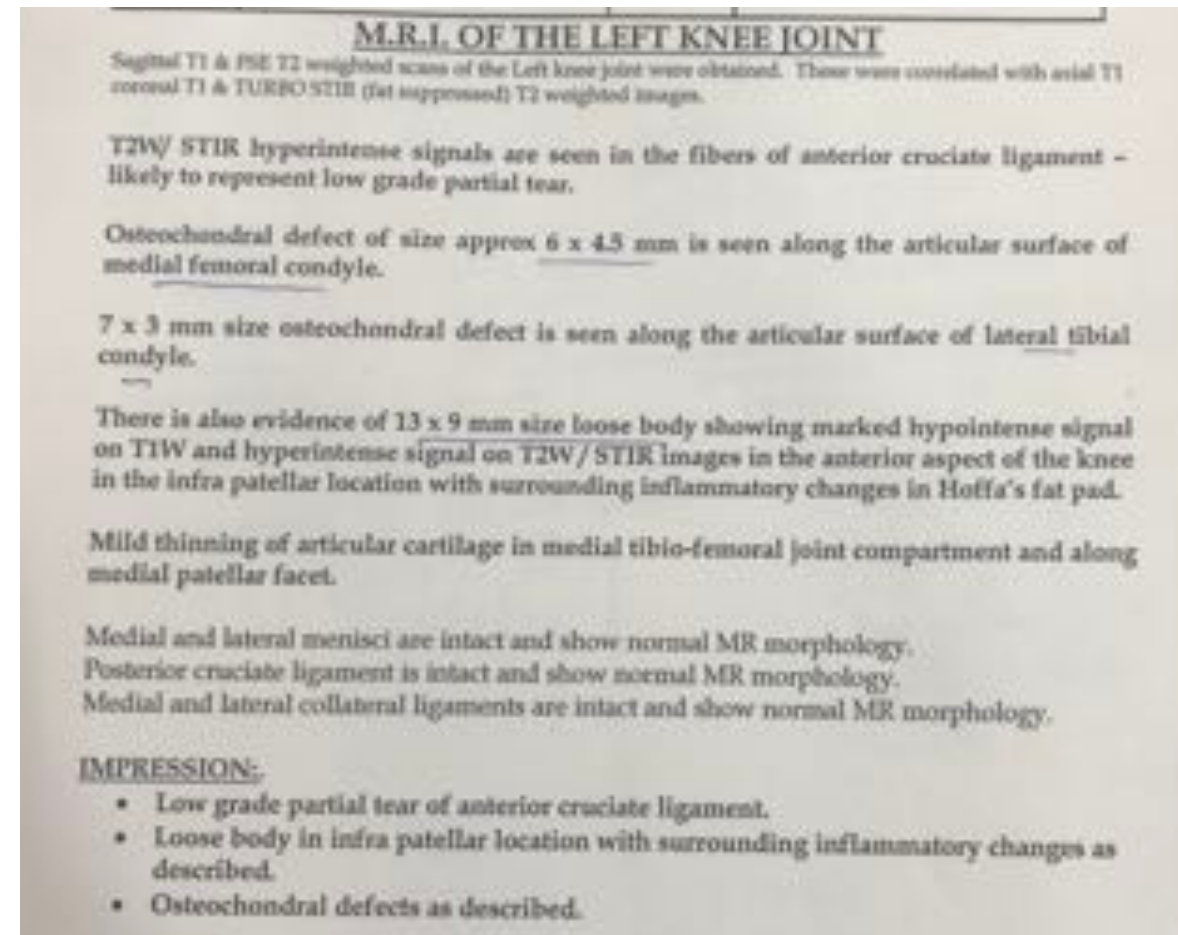
*OATS
(allograft)*

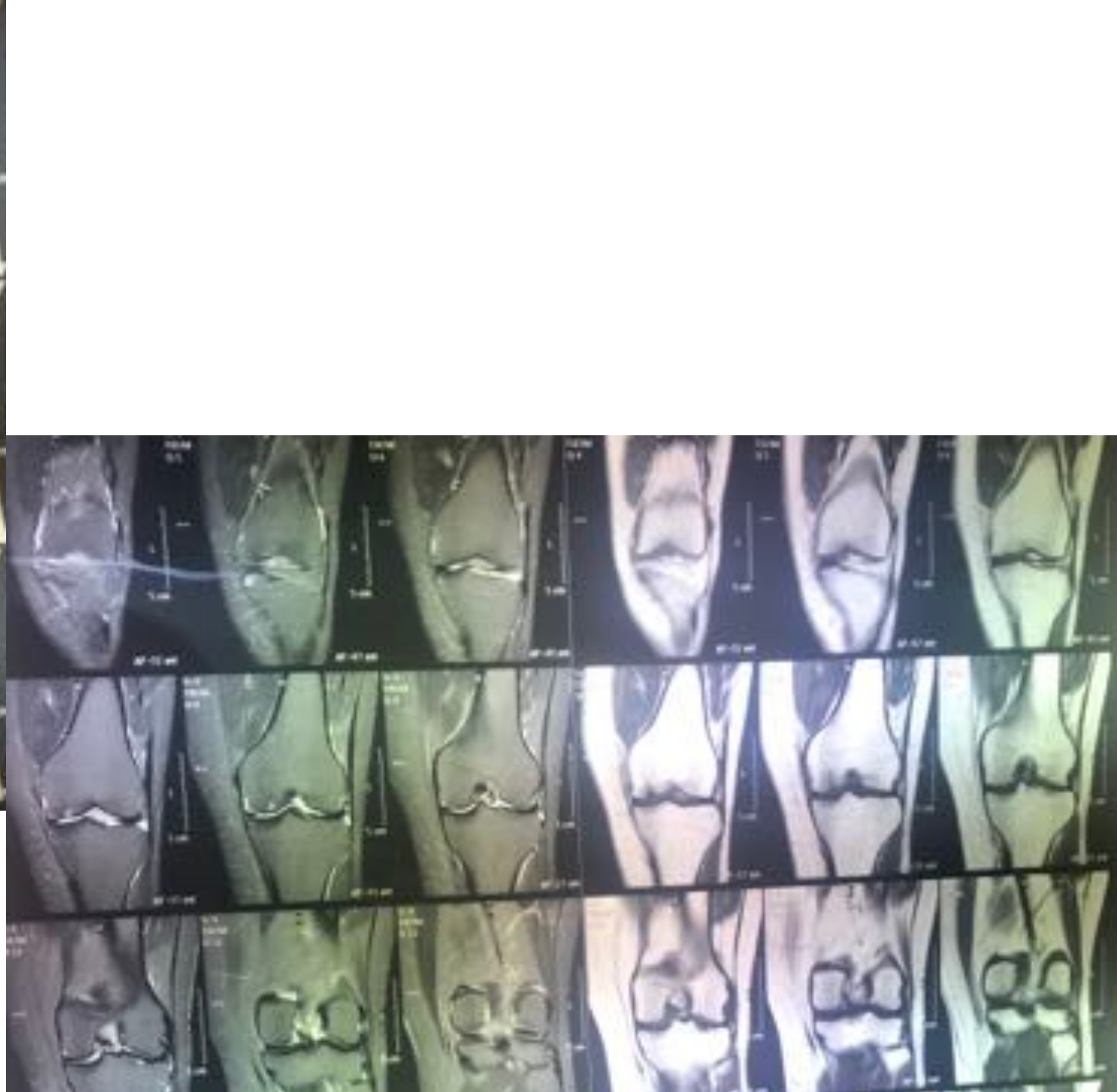
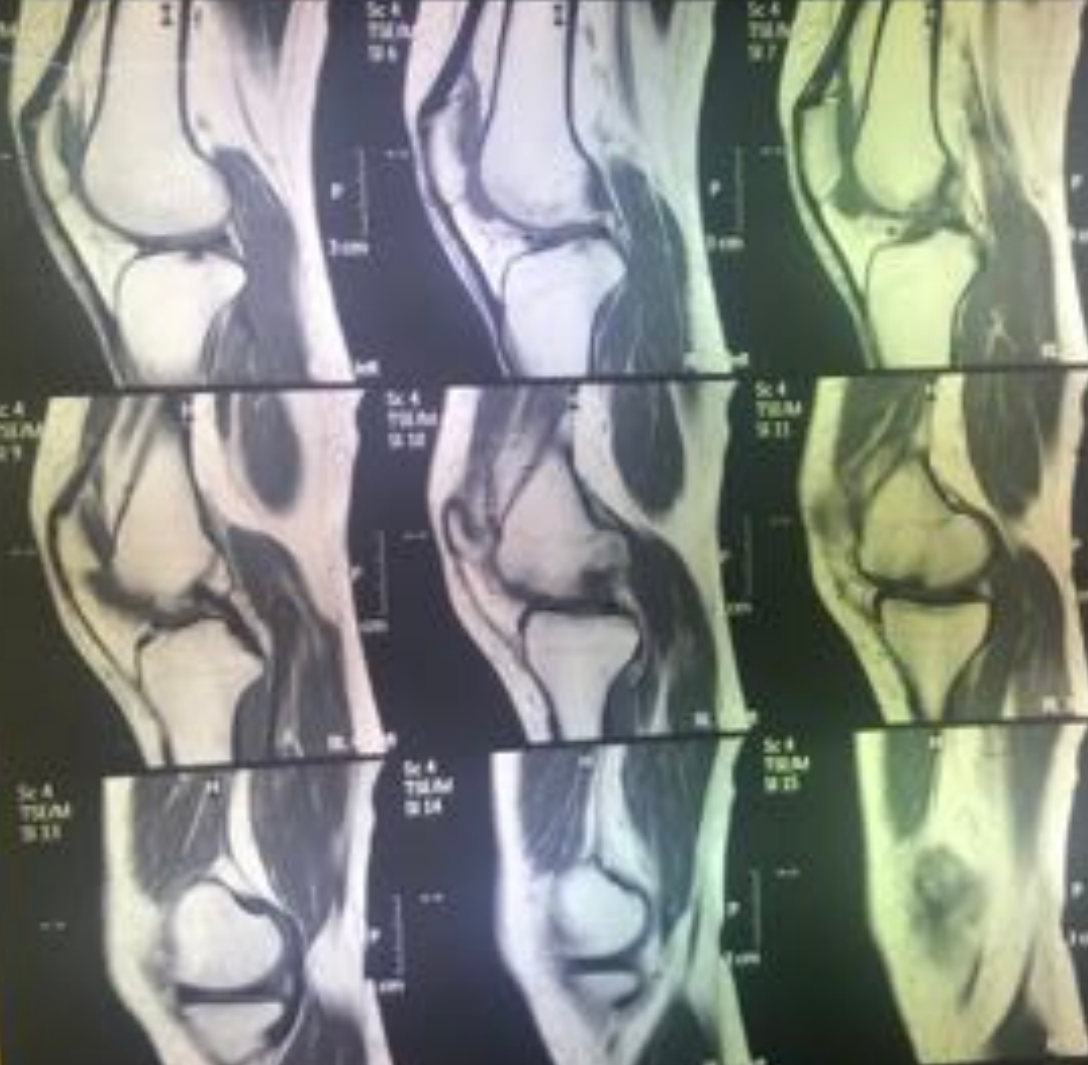
2.5-10 cms

ACI/ BMAC

Q. A 35 years female landed up with twisting injury to knee that was managed conservatively. 3 months post injury she started having multiple episodes of locking. MRI films and reports are shown. Surgical PLAN?

- A. Loose body removal and VS
- B. Microfracture and VS
- C. OATS
- D. ACI





Mosaicplasty

Bobic (1996)

Mosaicplasty in 12 patients with ACL injury: with good or excellent results in 10/12 cases

Wang (2002)

Mosaicplasty in 16 patients with 2 to 4 years of follow-up: 80% good or excellent results

Sharp et al (2005)

Combined chondrocyte transplantation and mosaicplasty with follow up of 3 years in 13 patients with large lesions
excellent results in 10/13 patients

10 years of follow-up
In conclusion : Osteochondral autograft transfer is recommended for smaller lesions, lesions in high-demand athletes, and lesions with associated bone loss, while microfracture is suited for medium-size defects with little or no bone loss in lower-demand patients and they should therefore be reserved for revision situations.

Conclusions: Impact insertion of osteochondral grafts generates damaging loads that cause chondrocyte death, particularly in the superficial zone, mainly as a result of apoptosis mediated the activation of caspases.

Clinical Relevance: Chondrocyte death that occurs during impact insertion of osteochondral grafts may lead to compromised function. Understanding the mechanisms and consequences of such impact loading may provide insights into potential therapeutic interventions, or lead to changes in the insertion technique, to decrease the cell injury associated with impact loading.

CHONDRO- PROTECTIVE AGENTS

- **COLLAGEN TYPE II SUPPLEMENTS**
- **GLUCOSAMINE**
- **CHONDROITIN SULPHATE**
- **S-ADENOSYL METHIONINE**
- **DIACERIN**

VISCOSUPPLEMENTATION



Obesity



Aging



Mechanical Stress



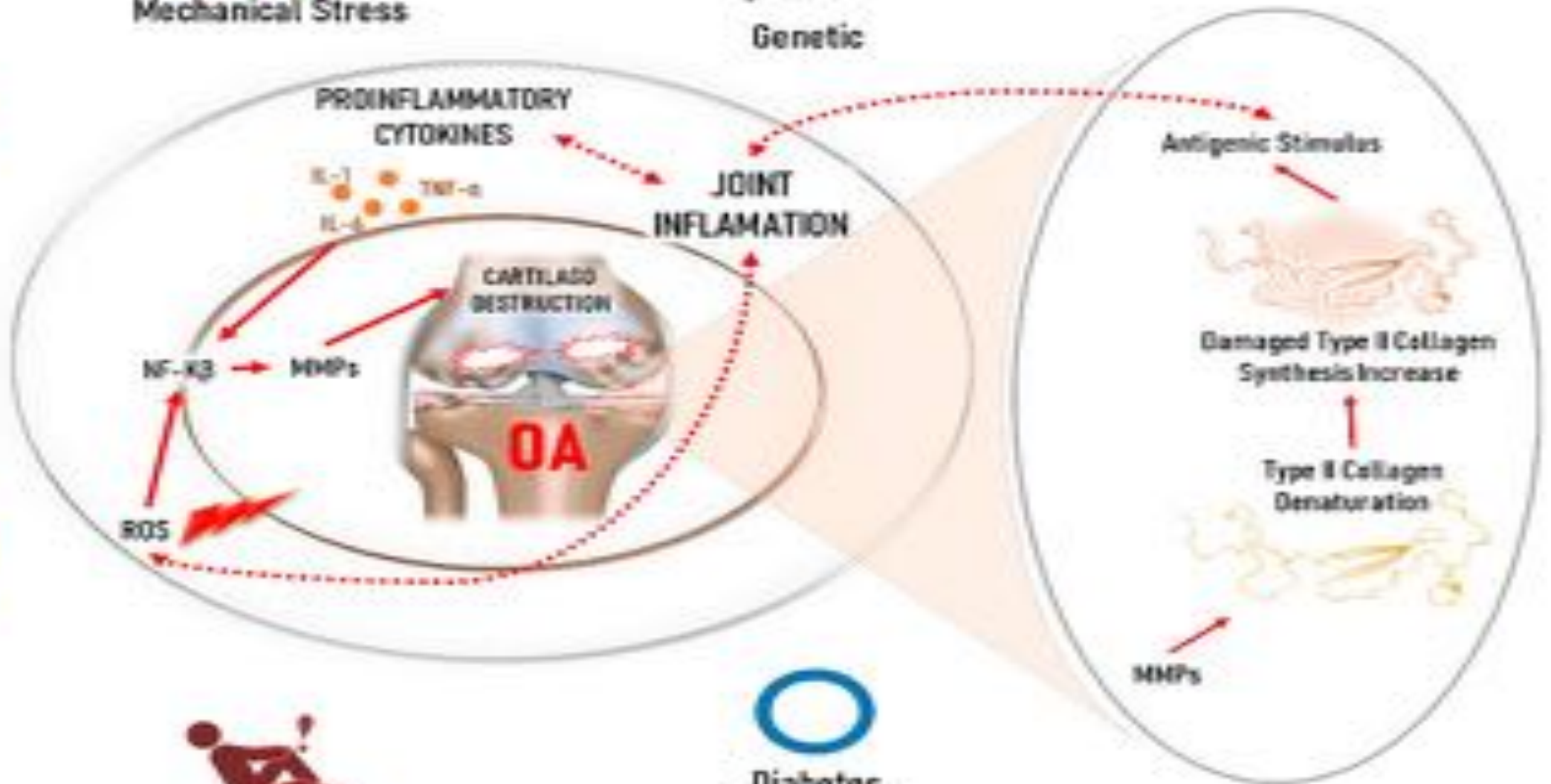
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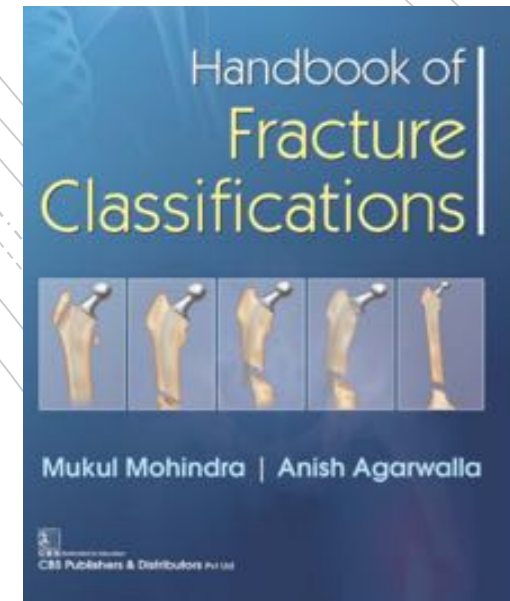
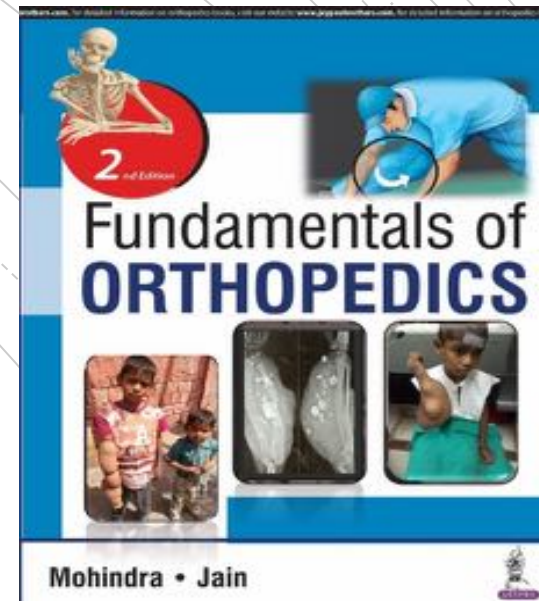
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THANK YOU