



# Morphological assessment of non-human primate models of osteoarthritis using HR-MRI and $\mu$ CT arthrography

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## Introduction

- Small animal models of osteoarthritis (OA) do not mimic perfectly the complex conditions occurring in human OA.
- OA that closely resembles the human condition occurs naturally in primate. Non-human primates (NHP) could be a useful model for human OA.
- Non-invasive techniques such as 3D HR-MRI have been validated to directly assess the cartilage thickness on guinea pigs (1) and different cartilage compartment volumes on rat models of OA (2-3).
- Nonetheless, spatial resolution is limited compared to µCT scanner that however needs contrast agent injected in the joint to depict cartilage limits.

## Objectives

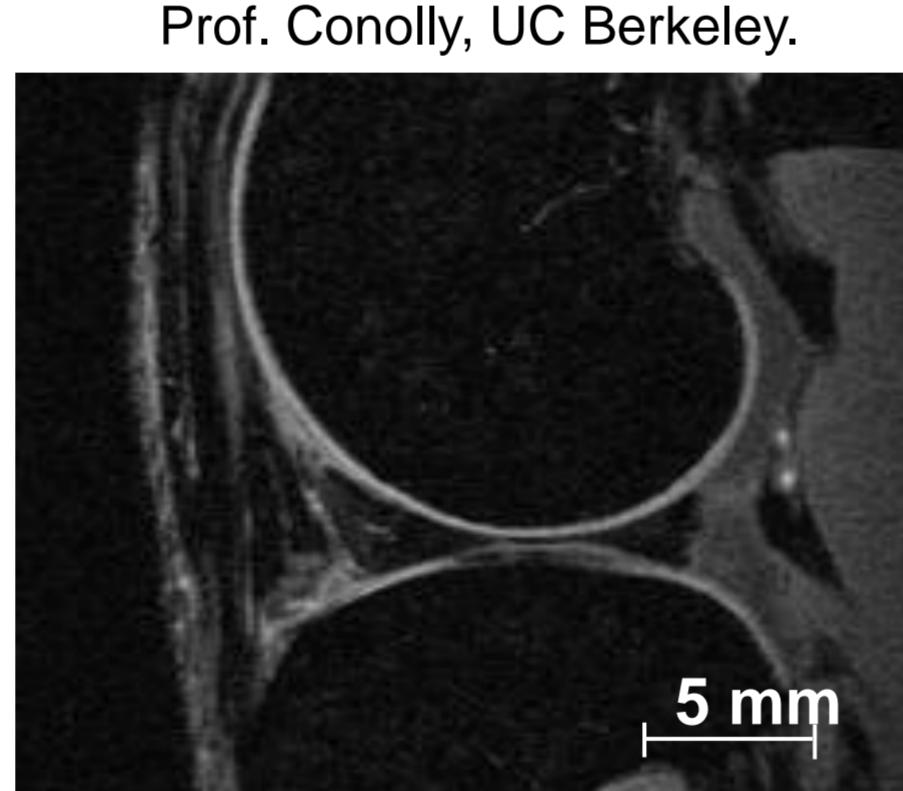
- The aim of this work, based on morphological parameters assessed on MRI and µCT arthrography (µCTA) acquisitions, was:
- To assess the potential µCTA protocol impact on the model follow-up
- To compare quantifications results based on both imaging modalities
- To characterize an induced model of OA by transection of the anterior cruciate ligament (ACL).

## Material and method

- The ethical guidelines for animal experimental investigations were followed and the experimental protocol was approved by the Animal Ethics Committee from Ecole nationale vétérinaire de Lyon (VetAgro Sup), Marcy l'Etoile.
- Group 1&3 (n=3+3): control animals – only the right knee was injected with HexabrixTM for µCTA imaging. MR imaging of both knees.
- Group 2 (n=6): ACL transection of the right knee only. Multi-modal imaging of both knees.
- Longitudinal follow-up using HR-MRI and µCTA of 4 year old female cynomolgus at baseline, D15 (µCTA only), D30, D60, D90, D180.

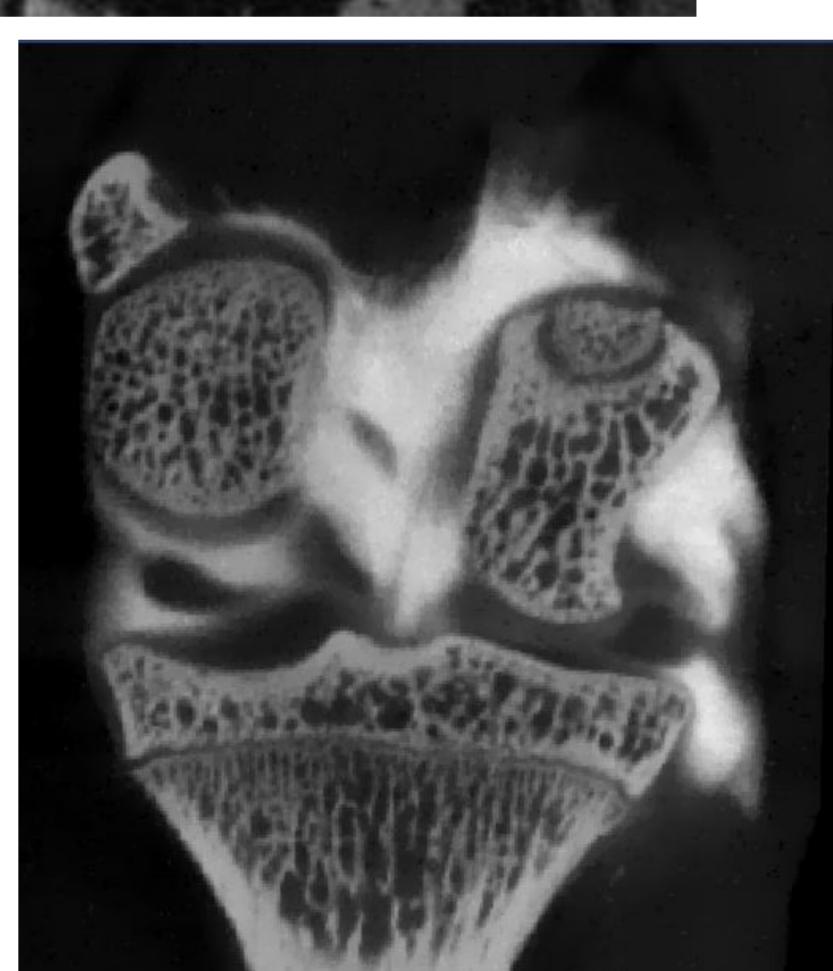
### MRI acquisition protocol

- 1.5T Siemens Sonata system
- 3D water excitation FLASH sequence: 25° flip angle, 27 ms TR, 11.7 ms TE, 70 Hz/Pixel receiver bandwidth
- A pair of homemade two-channel array coil
- In-plane pixel: 112x131 µm<sup>2</sup>, partition thickness: 220 µm
- scan time/knee : 20 min
- Total examination time: 90 min



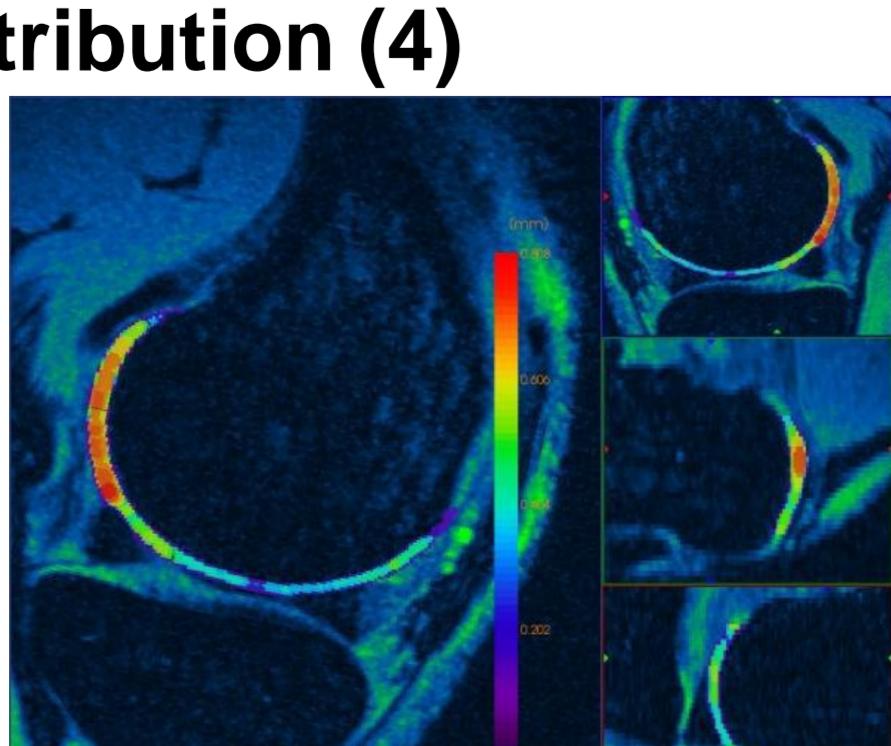
### µCTA acquisition protocol

- 2mL HexabrixTM (320mg/mL) with 40/60 dilution in PBS was injected in the synovial capsule with 23G needle.
- GE Locus µ-CT (standard voltage and amperage)
- Isotropic voxel of 90 µm
- Scan time for both knees : 15 min
- Total examination time: 30 min



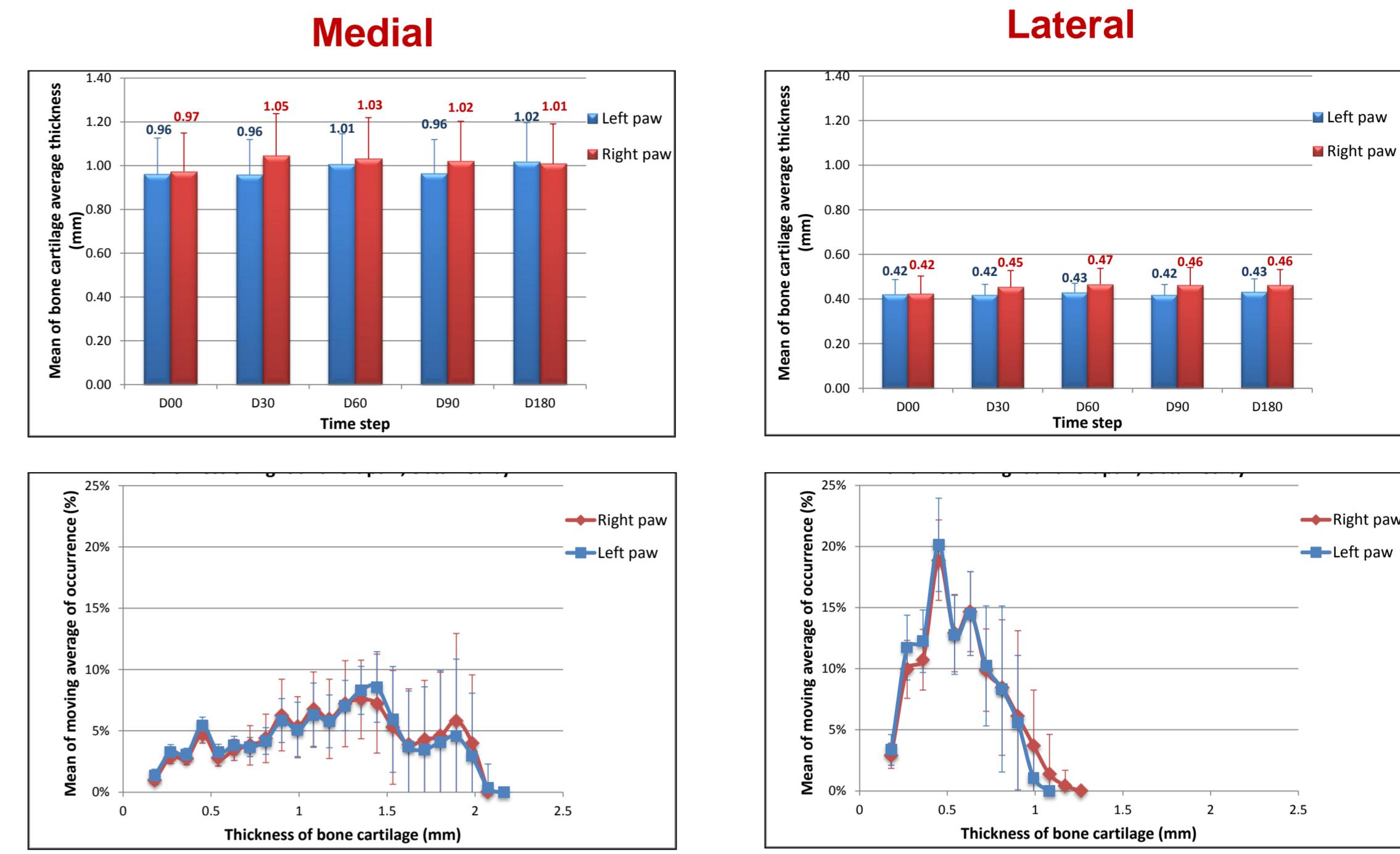
### Image processing protocol and cartilage quantification parameters

- Binarization performed in two steps:
  - Manual segmentation
  - Global thresholding
- Thickness distribution (4)



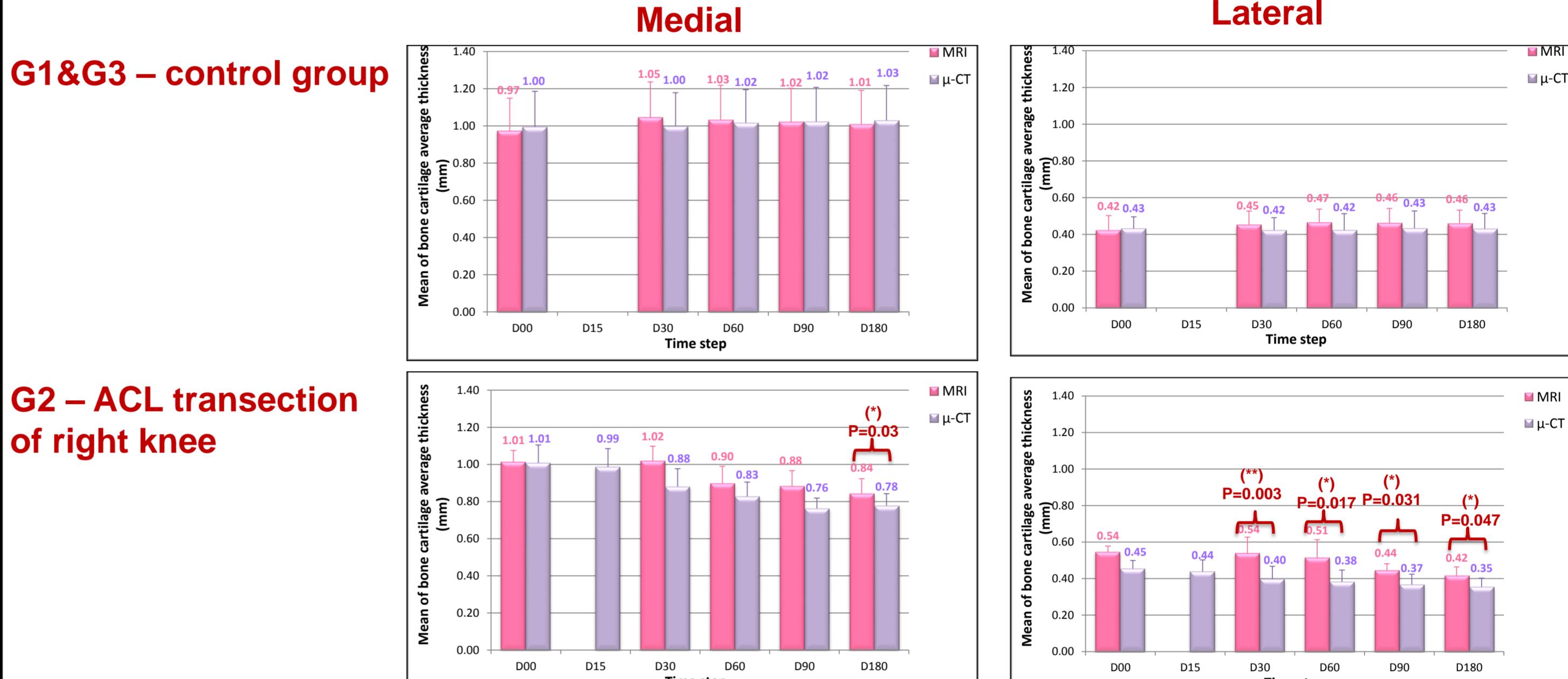
## Results

### G1 & G3 - MRI-based quantification of left and right tibial plateaus



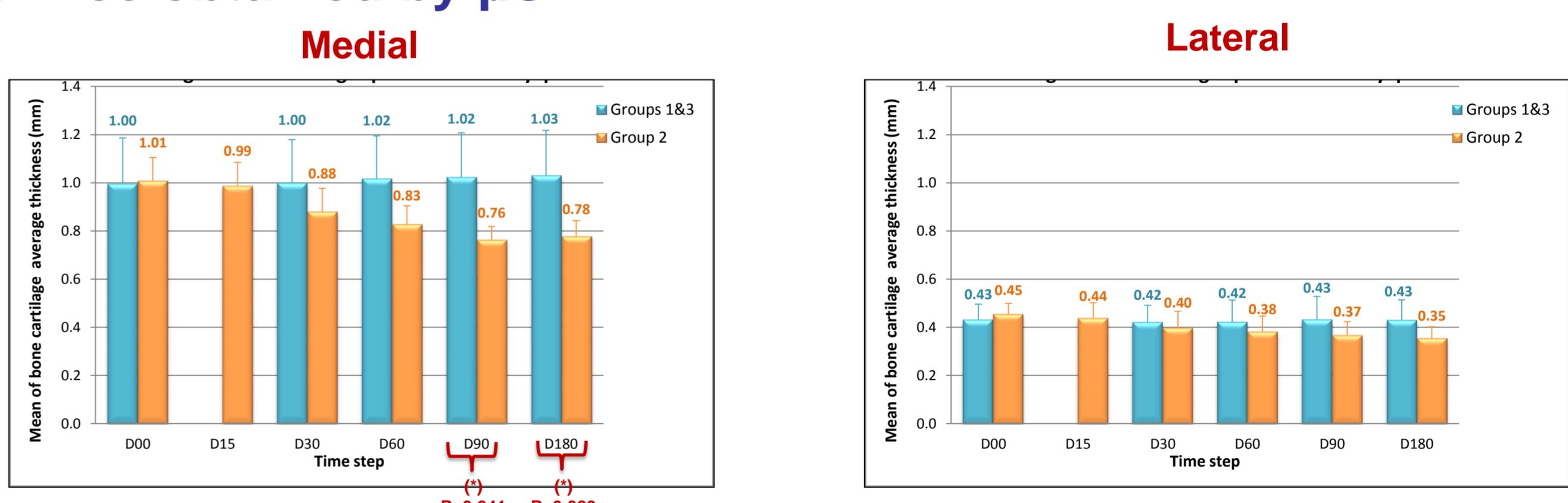
Repeated injections for µCTA have no impact on cartilage quantification

### G1&G3 vs G2 - Mean of tibial cartilage average thicknesses of right knee obtained by µCTA and MRI



Similar trend between µCTA and MRI with systematic thickness overestimation by MR

### G1&G3 vs G2 - Mean of tibial cartilage average thicknesses of right knee obtained by µCTA



Significative decrease of cartilage thickness starting at D90 and D180 on NPH model of OA

## Conclusion

- No differences were found with MRI examination between non injected and injected knees (required for the µCTA protocol) over the time.
- No difference was found between µCTA- and MRI-based cartilage thickness methods on control groups (G1&G3) since no residual measurement between both methods were found above the resolution of these techniques. Differences shown with G2 have to be further investigated.
- NHP model of OA was characterized by both imaging methods showing a monotone progression of the cartilage thinning up to  $-24.6 \pm 5.7\%$  on D90 and  $-27.2 \pm 5.2\%$  on D180.

## Perspectives

- MRI and µCTA modalities are valuable to measure cartilage morphology.
- Additional information can be obtained:
  - Indirectly about cartilage structure (T2, T1rho...) with MRI
  - Subchondral bone density with µCTA

### References:

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