

Relationship Between Patient-Reported
Outcome Measures (Proms) And MRI
Abnormalities In Early Knee Osteoarthritis

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Faculty Disclosure Information

There is nothing to disclosure





MRI is not recommended as an aid to identify or define early OA

- 1. KOOS ≤85% in at least 2 out of these 4 categories
 - pain(9items)
 stiffness and other symptoms (7items)
 - function and daily living (4items)quality of life(4items)

2. Clinical examination:

joint line tenderness or crepitus

3. Knee radiographs

Kellgren & Lawrence (KL) grade of 0 or 1





Early diagnosis and treatment of KOA are required to reduce the health care burden²⁾



It is important to understand the pathogenesis of early KOA using MRI

Purpose

To elucidate the association between MRI abnormalities and patientreported outcomes (KOOS) in patients with early KOA



Medial knee joint pain

K–L grade ≤ 1

Nonrandomized, prospective, multicentor clinical trial



Exclusion

history of ipsilateral lower extremity surgery

Age (years)	Sex (knees)	
59.1 (±12.1)	male 63	
	female 94	

MRI abnormalities*



The following items evaluated in WORMS³⁾ were investigated

- 1 cartilage abnormalities
- 2 bone marrow lesions (BMLs)
- ③ subchondral bone cysts (BMC)
- 4 subchondral bone attrition (SBA)
- ⑤ osteophytes
- 6 joint effusion**
- 7 medial meniscus tears * * *
- ® medial collateral ligament (MCL) bursitis

*The MRI images were interpreted twice by the speaker, with a two-week interval.

* * Medial meniscus injuries included horizontal, transverse, flap, and dorsal root tears.

Horizontal tears were defined as Mink classification 3 or higher, while those classified as 2 or lower were considered normal.

* * * Joint effusion was considered abnormal if graded as 1 or higher.

Statistics



Mann-Whitney U test was used to compare the mean values of each of the five KOOS subscales between two groups, categorized by the presence or absence of each MRI lesion.



We performed multiple regression analysis with the five KOOS subscales as outcome variables and age, BMI, and MRI lesions (significant in the Mann-Whitney U test) as explanatory variables. Statistical significance was set at p < 0.05.



High prevalence of cartilage abnormalities, osteophytes, and MCL bursitis

	Presence (knees)	Absence (knees)	Prevalence (%)
cartilage abnormalities	148	9	94.2
medial meniscus tears	131	26	83.4
joint effusion	122	35	77.7
osteophytes	118	39	75.2
MCL bursitis	115	42	73.3
MME>2mm	108	49	68.8
SBA	93	69	59.2
BMLs	67	97	42.7
BMC	29	140	18.5

Significant in the Mann-Whitney U test





BMLs→ significantly lower KOOS Pain/ADL Meniscal tear → significantly lower 5 KOOS subscales

BML	KOOS Symptom	KOOS Pain	KOOS ADL	KOOS Sport	KOOS QOL
presence	61.40	53.86	70.01	41.27	38.64
absence	68.02	62.28	76.36	46.00	39.82
P-value	0.057	0.004	0.027	0.294	0.723

Meniscal tear	KOOS Symptom	KOOS Pain	KOOS ADL	KOOS Sport	KOOS QOL
presence	63.36	56.19	72.04	42.02	37.34
absence	74.44	71.26	81.79	53.85	49.30
P-value	0.007	<0.001	0.003	0.043	0.004

Significant in the Mann-Whitney U test





Joint effusion → significantly lower KOOS Symptom/ADL MME>2mm → significantly lower KOOS Symptom/Pain

Joint effusion	KOOS Symptom	KOOS Pain	KOOS ADL	KOOS Sport	KOOS QOL
presence	63.09	57.40	72.10	42.30	38.60
absence	72.54	63.17	79.07	49.86	41.86
P-value	0.017	0.07	0.032	0.196	0.572

MME >2mm	KOOS Symptom	KOOS Pain	KOOS ADL	KOOS Sport	KOOS QOL
presence	62.80	56.45	72.04	41.39	38.05
absence	70.48	63.60	77.19	49.69	42.11
P-value	0.022	0.048	0.0788	0.064	0.304

Significant in the Multiple regression analysis

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KOOS Symptom	Regression coefficient (β)	95%CI	P-value
Age	0.344	(0.068-0.621)	0.009
Joint effusion	-8.061	(-14.77-0.532)	0.068
KOOS Pain	Regression coefficient (β)	95%CI	P-value
Meniscus tear	-12.701	(-20.7504.652)	0.002
BMLs	-7.362	(-13.0281.695)	0.011
KOOS Sport	Regression coefficient (β)	95%CI	P-value
Age	0.3660	(0.009-0.722)	0.045
KOOS Symptom	Regression coefficient (β)	95%CI	P-value
Age	0.303	(0.017-0.590)	0.038
Meniscus tear	-11.968	(-21.0962.841)	0.011

Discussion





Cartilage abnormalities were present in almost all cases (94%)

· Hada, et al. reported that all patients with early KOA had cartilage lesions⁴⁾



We propose that the pathogenesis of early KOA involves cartilage abnormalities



Joint effusion, medial meniscus tear, MME>2mm and BMLs were associated with knee symptoms

- Torres, et al. noted that joint effusion correlated best with knee pain⁵⁾
- There have been reports that meniscal lesions, BMLs, and joint effusion are associated with each other⁶⁾⁷⁾



Medial meniscus tears, joint effusion, MME and BML were associated with KOOS

Early treatment targeting these lesions might be key in preventing the progression of KOA

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