Popliteal cysts and subgastrocnemius bursitis are associated with knee symptoms and structural abnormalities in older adults: a cross-sectional study

Yuelong Cao1,2*, Graeme Jones1, Weiyu Han1, Benny Antony1, Xia Wang1, Flavia Cicuttini1 and Changhai Ding1,3*

Abstract

Introduction: The role of popliteal cysts and subgastrocnemius bursitis in knee joint homeostasis is uncertain. The aim of this study is to describe cross-sectional associations between popliteal cysts, subgastrocnemius bursitis, knee symptoms and structural abnormalities in older adults.

Methods: A cross-sectional sample of 900 randomly-selected subjects (mean age 63 years, 48% female) were studied. Knee pain, stiffness and dysfunction were assessed by self-administered Western Ontario McMaster Osteoarthritis Index (WOMAC) questionnaire. Radiographic knee osteophyte and joint space narrowing (JSN) were recorded. Magnetic resonance imaging (MRI) was utilized to assess popliteal cysts, subgastrocnemius bursitis, cartilage defects and bone marrow lesions (BMLs).

Results: Popliteal cysts were present in 11.7% and subgastrocnemius bursitis in 12.7% of subjects. Subgastrocnemius bursitis was more common in those with popliteal cyst (36.2% versus 9.7%, P <0.01). In multivariable analyses, popliteal cysts were significantly associated with increased osteophytes in both medial and lateral tibiofemoral compartments while subgastrocnemius bursitis was associated with increased osteophytes and JSN in the medial tibiofemoral compartment. Both were significantly associated with cartilage defects in all compartments, and with BMLs in the medial tibiofemoral compartment. Furthermore, both popliteal cysts and subgastrocnemius bursitis were significantly associated with increased weight-bearing knee pain but these associations became non-significant after adjustment for cartilage defects and BMLs.

Conclusions: Popliteal cysts and subgastrocnemius bursitis are associated with increased symptoms as well as radiographic and MRI-detected joint structural abnormalities. Longitudinal data will help resolve if they are a consequence or a cause of knee joint abnormalities.

Introduction

Osteoarthritis (OA) is a slowly progressive and multifactorial disease characterised by gradual loss of articular cartilage and other structural changes in the whole joint [1]. Among these structural changes, cystic lesions in the knee joint are a common feature seen on magnetic resonance imaging (MRI) or ultrasound assessment but may not be detected by physical examination [2].

Most cystic lesions around the knee joint represent encapsulated fluid collections. Although bursae are synovium-lined structures and normally quiescent, bursitis can be detected when bursae become inflamed and/or filled with fluid [3]. Among those cystic lesions, popliteal cyst (Baker cyst [4]) and subgastrocnemius bursitis are most commonly detected on MRI with the prevalence of about 38% for popliteal cyst [5] and 15% [6] for subgastrocnemius bursitis. Popliteal cysts are usually formed...
from extrusion of joint fluid into the gastrocnemius- semimembranosus bursa [7] and can extend deep to gastrocnemius muscles, resulting in fluid in the subgastrocnemius bursa. A narrow neck connecting the bursa to the knee joint is usually identified on axial MRI images and is commonly found just below the proximal attachment site of the medial head of gastrocnemius [6]. Therefore, subgastrocnemius bursae can communicate with the knee joint and popliteal cyst if co-existing.

In adults, almost all popliteal cysts are secondary [8] since a communication exists between the cysts and the knee joint, allowing articular fluid to enter the cyst. It is therefore reasonable to postulate that popliteal cyst and subgastrocnemius bursitis may be associated with intra-articular derangement of the knee joint; however, this has only been examined in a limited number of studies. Although popliteal cyst has been associated with the presence of a meniscal lesion, [5,9] it is unclear if popliteal cyst and subgastrocnemius bursitis play roles in radiographic OA, and MRI-detected structural abnormalities. The aim of our study was, therefore, to describe the prevalence of popliteal cyst and subgastrocnemius bursitis in a cohort of older adults and to examine their associations with radiographic and MRI-detected structural abnormalities, as well as knee symptoms.

Methods
Study design and subjects
The study was based on the Tasmanian Older Adult Cohort (TASOAC) study, a prospective epidemiological study of persons aged 50 to 79 years, with a goal of identifying the environmental, genetic and biochemical factors associated with the development and progression of OA and osteoporosis. Participants were selected randomly using computer-generated random numbers from the electoral roll in Southern Tasmania (population 229,000), a comprehensive population listing, with an equal number of men and women. Institutionalised persons were excluded. The study was approved by the Southern Tasmanian Health and Medical Human Research Ethics Committee, and written informed consent was obtained from all participants. Self-reports of disease status such as rheumatoid arthritis (RA), asthma, cardiovascular disease and diabetes were recorded by questionnaire. Baseline measurements were conducted from April 2002 to September 2004, and the overall response rate was 57%.

Anthropometrics
Height was measured to the nearest 0.1 cm by using a stadiometer, with shoes, socks and headgear removed. Weight was measured to the nearest 0.1 kg (with shoes, socks and bulky clothing removed) using a single pair of electronic scales (Seca Delta Model 707, Bradford, MA, USA) that were calibrated using a known weight at the beginning of each clinic. Body mass index (BMI) (weight (kg)/height² (m²)) was calculated.

MRI
Using a commercial transmit/receive extremity coil, MRI of the right knee was performed with a 1.5 T whole-body magnetic resonance unit (Picker, Cleveland, OH, USA). The following sequence and parameters were used: a T1-weighted fat-suppressed three-dimensional gradient-recalled acquisition in the steady state, flip angle 30°, repetition time 31 ms, echo time 6.71 ms, field of view 16 cm, 60 partitions, 512 × 512-pixel matrix, acquisition time 5 minutes 58 seconds, 1 acquisition, sagittal images obtained at a partition thickness of 1.5 mm without a between-slice gap; and a T2-weighted fat-saturated two-dimensional fast spin echo, flip angle 90°, repetition time 3,067 ms, echo time 112 ms, field of view 16 cm/15 partitions, 228 × 256-pixel matrix; sagittal images were obtained at a partition thickness of 4 mm with a between-slices gap of 0.5 to 1.0 mm. Assessments were performed on a standard digital imaging and communications in medicine (DICOM) viewer (Osiris 4, University of Geneva, Geneva, Switzerland).

Popliteal cyst and subgastrocnemius bursitis assessments
Assessments were performed on T2-weighted sagittal and axial views (Figure 1). Following to a previously published method, [10,11] popliteal cysts and subgastrocnemius bursitis were scored using a semiquantitative scale: grade 0 for absent, grade 1 for small, grade 2 for moderate, grade 3 for large. Interobserver reproducibility was assessed between two readings in 82 images of randomly selected subjects, with interclass correlation coefficients of 0.87 for subgastrocnemius bursitis and 0.86 for popliteal cysts. A further random subset of these images was re-read for intraclass correlation coefficients (ICCs) with 0.93 and 0.94 for subgastrocnemius bursitis and popliteal cysts, respectively.

Cartilage defects and bone marrow lesion (BML) assessments
Cartilage defects were assessed at the medial tibial, medial femoral, lateral tibial, and lateral femoral sites as previously reported [12]: grade 0 = normal cartilage; grade 1 = focal blistering and intracartilaginous low-signal intensity area with an intact surface and bottom; grade 2 = irregularities on the surface or bottom and loss of thickness <50%; grade 3 = deep ulceration with loss of thickness >50%; grade 4 = full-thickness chondral wear with exposure of subchondral bone. A cartilage defect also had to be present in at least two consecutive slices. ICCs were 0.89 to 0.94, and interclass correlation coefficients were 0.85 to 0.93 [13].
BMLs were assessed on the T2-weighted images and defined as discrete areas of increased signal adjacent to the subcortical bone at the medial tibial, medial femoral, lateral tibial, and lateral femoral sites as previously described [14]. Each BML was scored on the basis of lesion size: grade 0 = no BML; grade 1 = only present on one slice; grade 2 = on two consecutive slices; grade 3 = on more than three consecutive slices. The highest score was used if more than one lesion was present on the same site. Intraobserver repeatability was assessed in 50 subjects with at least a one-week interval between two readings, with ICCs of 0.89, 0.96, 0.94, and 1.00 for the BML scores at lateral tibia, lateral femur, medial tibia, and medial femur, respectively.

Knee effusion was graded from 0 to 3 in terms of the estimated maximal distension according to the whole-organ magnetic resonance imaging score (WORMs) method [11].

Western Ontario McMaster osteoarthritis index (WOMAC) and knee pain assessment
Each component of knee pain, stiffness and dysfunction was assessed by self-administered WOMAC questionnaire [15] with a 10-point scale from 0 (no symptom) to 9 (most severe symptom). Total knee pain was the sum of all components to create a total pain (0 to 45) score. The five WOMAC pain questions related to knee OA were clinically constructed in three weight-bearing pain questions (pain on climbing stairs, on walking, and on standing) and two non-weight-bearing pain questions (pain in bed or when sitting/lying down) as suggested by a recent study [16].

Radiographic OA assessment
This was scored according to the Osteoarthritis Research Society International (OARSI) atlas [17] as previously reported [13]. A standing semi-flexed view of the right knee with 15° of fixed knee flexion was performed in all subjects and scored individually for joint space narrowing (JSN) and osteophytes on a scale of 0 to 3 (0 = normal and 3 = severe). Medial or lateral JSN was scored separately, and osteophytes were scored at each site of the medial tibia, medial femur, lateral tibia and lateral femur.

Data analysis
The Student t-test or χ² test was used to compare means or proportions, respectively, between those with and without popliteal cyst. Ordinal logistic regression was
results, we examined the associations between popliteal cysts, subgastrocnemius bursitis (dependent variables) and the anthropometric/other risk factors (independent variables). Univariable and multivariable ordinal logistic regression analysis was used to examine the associations between dependent variables (score of knee cartilage defects, BMLs, osteophytes and JSN in the medial or lateral tibiofemoral site) and independent variables (each score for popliteal cyst and subgastrocnemius bursitis) before and after adjustment for age, sex, BMI, smoking status, radiographic features, and/or disease status (RA, cardiovascular disease, asthma and diabetes). Binary logistic regression models were used to examine the associations between the cystic lesion score (independent variables) and presence of knee symptoms, including weight-bearing pain, non-weight-bearing pain, total knee pain, stiffness and dysfunction (dependent variables). The associations between cystic lesions and knee symptoms were further adjusted for effusion or cartilage defects/BMLs to see if they were mediated by these structural abnormalities.

Standard diagnostic checks of model fit and residuals were routinely made, and data points with large residuals and/or high influence were investigated for data errors. Interactions between sex and cystic lesions on OA measures were investigated by testing the statistical significance of interactions between sex and cystic lesions on OA measures. In univariable analysis both were significantly and positively associated with knee cartilage defects at all sites (all P <0.05) and more osteophytes and cartilage defects at all sites (all P <0.05). Subgastrocnemius bursitis (prevalence 12.7%) was more common in those with popliteal cyst (36.2% versus 9.7%, P <0.01). Thus, 38 subjects had both popliteal cysts and subgastrocnemius bursitis, 67 had popliteal cyst and 77 had subgastrocnemius bursitis only (Table 1).

Age, sex, BMI, and smoking were not associated with cystic lesions (data not shown); however, there was a significant association between history of knee surgery and the popliteal cyst score (knee surgery versus no knee surgery: β = 0.70, P = 0.01).

### Results

Of 900 subjects (49.7% female) included in the analysis, the average age at baseline was 62.3 years. The mean BMI was 27.7 kg/m². Popliteal cysts were present in 105 (11.7%) subjects, with 47 in grade 1, 40 in grade 2, and 18 in grade 3. Subjects with or without popliteal cyst were similar in terms of age, gender, BMI, non-weight-bearing pain, total knee pain, prevalence of BMLs, osteophytes and cartilage defects at all sites (all P >0.05). Subgastrocnemius bursitis (prevalence 12.7%) was more common in those with popliteal cyst (36.2% versus 9.7%, P <0.01). Thus, 38 subjects had both popliteal cysts and subgastrocnemius bursitis, 67 had popliteal cyst and 77 had subgastrocnemius bursitis only (Table 1).

### Table 1 Baseline characteristic of participants

| Characteristic | Without popliteal cyst (n = 795) | With popliteal cyst (n = 105) | P       |
|---------------|----------------------------------|------------------------------|---------|
| Age, years    | 62.2 (7.3)                       | 63.2 (8.0)                   | 0.18*   |
| Female gender,%| 49.9                            | 47.6                         | 0.65    |
| Body mass index, kg/m² | 27.6 (4.5)   | 27.9 (4.9)                   | 0.53*   |
| WOMAC knee pain | Weight-bearing, 0 to 27 mm       | 4.7 (4.9)                    | 5.8 (5.0) | 0.03*   |
|               | Non-weight-bearing, 0 to 18 mm    | 3.2 (4.7)                    | 3.1 (4.7) | 0.95*   |
|               | Total pain, 0 to 45 mm            | 5.1 (5.0)                    | 6.0 (4.9) | 0.07*   |
|               | Prevalence of bone marrow lesions, % | 34.0                        | 40.0 | 0.23    |
|               | Prevalence of osteophytes, %      | 8.2                          | 16.2    | 0.01    |
|               | Medial tibiofemoral JSN, %        | 52.6                         | 54.5 | 0.30    |
|               | Lateral tibiofemoral JSN, %       | 23.4                         | 27.3 | 0.34    |
|               | Cartilage defect, ≥2, %           | 19.7                         | 31.4 | <0.001  |
|               | Medial tibial                     | 20.4                         | 80.4 | <0.001  |
|               | Lateral tibial                    | 15.3                         | 28.6 | 0.01    |
|               | Medial femoral                    | 19.7                         | 31.4 | <0.001  |
|               | Lateral femoral                   | 9.6                          | 15.2 | 0.01    |
|               | Patellar                          | 38.9                         | 49.5 | 0.01    |
|               | Effusion, WOMAC ≥1, %             | 25.9                         | 30.5 | 0.56    |
|               | Subgastrocnemius bursitis, grade ≥1, % | 9.7                          | 36.2 | <0.001  |

Results presented as mean (SD) except for percentages. Data in bold denote statistically significant results. JSN, joint space narrowing; WOMAC, Western Ontario McMaster Universities osteoarthritis index; WOMS, whole-organ magnetic resonance imaging score; *t-tests; others were χ² tests.

### Popliteal cyst, subgastrocnemius bursitis and radiographic features

In multivariable analysis, whereas popliteal cysts were significantly associated with increased medial and lateral tibiofemoral osteophytes, subgastrocnemius bursitis was significantly associated with increased medial tibiofemoral JSN and osteophytes (Table 2, Figure 2). Popliteal cyst and subgastrocnemius bursitis were not significantly associated with lateral tibiofemoral JSN (Table 2).

### Popliteal cyst, subgastrocnemius bursitis and MRI-detected structural features

Popliteal cyst or subgastrocnemius bursitis was significantly and positively associated with knee cartilage defects at medial and lateral tibiofemoral sites before and after adjustment for age, sex, BMI, disease status and knee radiographic features. In univariable analysis both were also significantly associated with a patellar cartilage defect and remained significant, except for popliteal cyst, after adjustment for the covariates as mentioned above (Table 3). Moreover, in multivariable analysis both were...
significantly associated with medial tibiofemoral BMLs, but not with lateral tibiofemoral BMLs (Table 4).

**Popliteal cyst, subgastrocnemius bursitis and knee symptoms**

Both subgastrocnemius bursitis and popliteal cyst were associated significantly with the presence of weight-bearing knee pain, but not with the presence of non-weight-bearing pain or total knee pain, before and after adjustment for age, sex, BMI, disease status, and knee radiographic features. Moreover, subgastrocnemius bursitis was significantly associated with prevalent WOMAC stiffness and popliteal cyst was significantly associated with prevalent WOMAC dysfunction (Table 5). All these significant

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|                | Univariable | Multivariable* |
|----------------|-------------|----------------|
|                | β   | 95% CI | P      | β   | 95% CI | P      |
| Medial tibiofemoral JSN |               |               |        |               |               |        |
| Subgastrocnemius bursitis | 0.32 | (0.08, 0.56) | 0.01  | 0.29 | (0.05, 0.54) | 0.01  |
| Popliteal cyst | 0.04 | (−0.17, 0.25) | 0.68  | −0.03 | (−0.25, 0.19) | 0.77  |
| Lateral tibiofemoral JSN |               |               |        |               |               |        |
| Subgastrocnemius bursitis | −0.11 | (−0.42, 0.19) | 0.47  | −0.17 | (−0.49, 0.14) | 0.28  |
| Popliteal cyst | 0.01 | (−0.24, 0.26) | 0.92  | −0.02 | (−0.28, 0.24) | 0.87  |
| Medial tibiofemoral osteophyte |               |               |        |               |               |        |
| Subgastrocnemius bursitis | 0.63 | (0.30, 0.95) | <0.001 | 0.61 | (0.24, 0.97) | 0.01  |
| Popliteal cyst | 0.48 | (0.17, 0.78) | 0.01  | 0.60 | (0.25, 0.95) | 0.01  |
| Lateral tibiofemoral osteophyte |               |               |        |               |               |        |
| Subgastrocnemius bursitis | −0.02 | (−0.67, 0.63) | 0.95  | −0.09 | (−0.78, 0.58) | 0.78  |
| Popliteal cyst | 0.43 | (0.03, 0.82) | 0.03  | 0.46 | (0.02, 0.91) | 0.03  |

Dependent variable: joint space narrowing (JSN) or osteophyte score. Independent variable: subgastrocnemius bursitis or popliteal cyst, per grade. *Adjusted for age, sex, body mass index, disease status, medial or lateral tibiofemoral osteophytes for JSN, or medial or lateral tibiofemoral JSN for osteophytes. Data in bold denote statistically significant results.

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![Figure 2](http://arthritis-research.com/content/16/2/R59)

**Figure 2** Associations between popliteal cysts, subgastrocnemius bursitis, and knee radiographic features. (A) Subjects with popliteal cysts had significantly higher medial tibiofemoral osteophyte score than those without popliteal cysts. (B) Subjects with subgastrocnemius bursitis had a significantly increased score for medial tibiofemoral joint space narrowing (JSN) and osteophytes compared with those without subgastrocnemius bursitis.
associations remained unchanged after further adjustment for effusion, but decreased in magnitude and became non-significant after further adjustment for cartilage defects and BMLs (Table 5).

There were no significant interactions between sex and cystic lesions on OA measures so male and female subjects were combined for analysis (data not shown). The results remained unchanged when subjects with RA were excluded from analysis (data not shown).

**Discussion**

To the best of our knowledge this is the first study to comprehensively examine the relationship between popliteal cyst, subgastrocnemius bursitis and imaging abnormalities of knee OA as well as symptoms in older adults.

We documented that popliteal cyst or subgastrocnemius bursitis was not only associated with radiographic features (osteophytes and JSN), cartilage defects and BMLs, but also with knee OA symptoms, including weight-bearing pain, stiffness and dysfunction. The associations were independent of confounders including age, sex, BMI, radiographic OA and/or disease status, suggesting popliteal cyst and subgastrocnemius bursitis may play roles in the pathological process of OA.

In adults, the aetiology of popliteal cyst may be related to an inflammatory process, meniscal tears, or mechanical intra-articular derangements of the knee joint. In our study, knee surgical history was associated with popliteal cyst. We did not find that age, sex, BMI and smoking, the OA risk factors, were associated with popliteal cyst and subgastrocnemius bursitis in older adults.

Both JSN and osteophytes are the key elements of radiographic OA. We documented that popliteal cyst was associated with osteophytes both in the medial and lateral tibiofemoral compartments whereas subgastrocnemius bursitis was associated with osteophytes and JSN in the medial tibiofemoral compartment, suggesting that popliteal cyst and subgastrocnemius bursitis can co-exist with radiographic OA.

Nevertheless, radiographic JSN is a surrogate method used to assess knee cartilage, [18] and has been regarded as insensitive to change. Using arthroscopy, previous studies reported that 41% to 83% of joint disorders including meniscal tears and chondral lesions were associated with popliteal cysts [19-21]. MRI allows non-invasive, direct, accurate and reliable assessment of cartilage and other joint structures with their changes over time [22]. So far, there are no reports describing the associations between knee cyst lesions and MRI-based measurements such as

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**Table 3 Association between subgastrocnemius bursitis, popliteal cyst and cartilage defects**

|                      | Univariable | Multivariable* |
|----------------------|-------------|---------------|
|                      | β           | 95% CI        | P     | β           | 95% CI        | P     |
| Medial tibiofemoral defects |             |               |       |             |               |       |
| Subgastrocnemius bursitis | 0.41        | (0.18, 0.64)  | <0.001 | 0.40        | (0.16, 0.65)  | 0.01 |
| Popliteal cyst       | 0.39        | (0.18, 0.59)  | <0.001 | 0.32        | (0.10, 0.53)  | 0.01 |
| Lateral tibiofemoral defects |             |               |       |             |               |       |
| Subgastrocnemius bursitis | 0.39        | (0.16, 0.62)  | 0.01  | 0.38        | (0.13, 0.62)  | 0.01 |
| Popliteal cyst       | 0.47        | (0.27, 0.67)  | <0.001 | 0.41        | (0.20, 0.62)  | <0.001 |
| Patellar defects     |             |               |       |             |               |       |
| Subgastrocnemius bursitis | 0.37        | (0.14, 0.59)  | 0.00  | 0.34        | (0.10, 0.59)  | 0.01 |
| Popliteal cyst       | 0.25        | (0.04, 0.45)  | 0.01  | 0.18        | (−0.02, 0.39)  | 0.08 |

Dependent variable: cartilage defects. Independent variable: subgastrocnemius bursitis or popliteal cyst, per grade. *Adjusted for age, sex, body mass index, disease status, and radiographic osteoarthritis. Data in bold denote statistically significant results.

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**Table 4 Association between subgastrocnemius bursitis, popliteal cyst and BMLs**

|                      | Univariable | Multivariable* |
|----------------------|-------------|---------------|
|                      | β           | 95% CI        | P     | β           | 95% CI        | P     |
| Medial tibiofemoral BMLs |             |               |       |             |               |       |
| Subgastrocnemius bursitis | 0.20        | (−0.06, 0.47) | 0.13  | 0.29        | (0.01, 0.57)  | 0.03 |
| Popliteal cyst       | 0.26        | (0.03, 0.49)  | 0.02  | 0.26        | (0.02, 0.49)  | 0.03 |
| Lateral tibiofemoral BMLs |             |               |       |             |               |       |
| Subgastrocnemius bursitis | 0.20        | (−0.07, 0.48) | 0.15  | 0.18        | (−0.12, 0.48) | 0.24 |
| Popliteal cyst       | 0.08        | (−0.17, 0.34) | 0.52  | 0.13        | (−0.13, 0.39) | 0.32 |

Dependent variable: bone mass lesions (BMLs). Independent variable: subgastrocnemius bursitis or popliteal cyst, per grade. *Adjusted for age, sex, body mass index, disease status, and radiographic osteoarthritis. Data in bold denote statistically significant results.
cartilage defects and BMLs. Cartilage defects have been associated with tibiofemoral osteophytes, increased tibial bone area, decreased knee cartilage volume and type II collagen breakdown [13]. Knee cartilage defects are predictive of knee cartilage loss over 2 years, [23] suggesting a potential target for intervention in OA [24]. BMLs are commonly detected in knee OA as increased signal intensity within the bone marrow on MRI. BMLs are strongly associated with knee pain and have appeared to be a promising target for OA [25,26]. Using MRI we have reported that in the current study, popliteal cyst and subgastocnemius bursitis were positively associated with cartilage defects in all compartments. These associations were independent of covariates including female sex, age, BMI and radiographic features, all which have been associated with change in knee cartilage defects [27]. Furthermore, we found that both popliteal cyst and subgastocnemius bursitis were associated with medial tibiofemoral BMLs in older adults.

The reasons underlying the association between popliteal cyst, subgastocnemius bursitis and knee structural abnormalities, particularly in the medial compartment, are unclear. The medial compartment of the knee bears the majority of the mechanical loads. Abnormal loading can lead to changes in the composition, structures, [28] and mechanical properties of articular cartilage [29]. It is highly possible that weight-bearing loads can be perceived by joint fluid triggering two-way flow between the cavity and bursae, and thus induces more cartilage loss, BMLs or osteophyte formation in the medial compartment.

It is also possible that there are elevated cytokines (for example, IL-6, TNF-α) or growth factors (for example, transforming growth factor (TGF)-β) in the cysts that can induce increased JSN, cartilage loss [30] and osteophytes [31]. Hill and colleagues reported 43.2% prevalence of popliteal cysts in knees with moderate or larger effusions, compared with 22.7% in those with little or no effusion, and the presence of popliteal cysts was not associated with knee pain [10]. Chatzopoulos et al. reported that popliteal cysts are a common finding in the knee among subjects with chronic OA pain [32]. These studies did not distinguish weight-bearing from non-weight-bearing pain, which are two distinct constructs of WOMAC pain [16]. By utilizing these classifications of pain, we have been able to detect that weight-bearing pain rather than non-weight-bearing pain was related to popliteal cyst or subgastocnemius bursitis. This finding is consistent with common belief, as weight-bearing pressure can cause compression of the adjacent cyst structures, and a relatively high proportion of weight-bearing pressure inside the cyst can induce knee pain. Increased pressure can promote extravasation of joint fluid through the posteromedial joint capsule into the bursae [33] and cause knee pain; however, it may not be related to joint cavity effusion observed simultaneously, because the significant association remained unchanged after further adjustment for effusion. The significant associations between knee pain and popliteal cyst and subgastocnemius bursitis may be mediated partly by cartilage

| Table 5 Association between subgastrocnemius bursitis, popliteal cysts and WOMAC measures |
|---------------------------------|-----------------|-----------------|-----------------|
|                                | Odds ratio | 95% CI       | P       | Odds ratio | 95% CI       | P       | Odds ratio | 95% CI       | P       |
| Total pain                     |            |               |         |            |               |         |            |               |         |
| Subgastrocnemius bursitis      | 1.31       | (0.99, 1.73)  | 0.05    | 1.31       | (0.99, 1.73)  | 0.05    | 1.20       | (0.91, 1.61)  | 0.19    |
| Popliteal cyst                 | 1.24       | (0.98, 1.56)  | 0.07    | 1.24       | (0.98, 1.56)  | 0.07    | 1.19       | (0.94, 1.51)  | 0.15    |
| Weight-bearing pain            |            |               |         |            |               |         |            |               |         |
| Subgastrocnemius bursitis      | 1.34       | (1.02, 1.77)  | 0.03    | 1.34       | (1.02, 1.77)  | 0.04    | 1.24       | (0.94, 1.64)  | 0.14    |
| Popliteal cyst                 | 1.27       | (1.01, 1.60)  | 0.04    | 1.27       | (1.01, 1.60)  | 0.04    | 1.23       | (0.97, 1.56)  | 0.09    |
| Non-weight bearing pain        |            |               |         |            |               |         |            |               |         |
| Subgastrocnemius bursitis      | 0.85       | (0.63, 1.13)  | 0.27    | 0.85       | (0.64, 1.14)  | 0.28    | 0.85       | (0.64, 1.14)  | 0.28    |
| Popliteal cyst                 | 0.97       | (0.76, 1.24)  | 0.82    | 0.97       | (0.76, 1.24)  | 0.82    | 0.97       | (0.76, 1.24)  | 0.81    |
| Stiffness                      |            |               |         |            |               |         |            |               |         |
| Subgastrocnemius bursitis      | 1.35       | (1.03, 1.79)  | 0.03    | 1.36       | (1.03, 1.80)  | 0.03    | 1.29       | (0.98, 1.72)  | 0.07    |
| Popliteal cyst                 | 1.20       | (0.95, 1.52)  | 0.12    | 1.20       | (0.94, 1.52)  | 0.12    | 1.18       | (0.93, 1.49)  | 0.18    |
| Dysfunction                    |            |               |         |            |               |         |            |               |         |
| Subgastrocnemius bursitis      | 1.18       | (0.9, 1.57)   | 0.23    | 1.18       | (0.9, 1.57)   | 0.23    | 1.14       | (0.85, 1.53)  | 0.37    |
| Popliteal cyst                 | 1.29       | (1.00, 1.67)  | 0.04    | 1.29       | (1.00, 1.67)  | 0.04    | 1.24       | (0.96, 1.61)  | 0.09    |

Dependent variable: presence of pain, stiffness or dysfunction (yes versus no). Independent variable: subgastrocnemius bursitis or popliteal cyst, per grade.

*Adjusted for age, sex, body mass index, disease status, and radiographic osteoarthritis; #further adjusted for effusion; †further adjusted for bone mass lesions and cartilage defects. Data in bold denote statistically significant results. WOMAC, Western Ontario McMaster Universities osteoarthritis index.
defects and BMLs, because the associations decreased in magnitude and became non-significant after adjustment for these structural abnormalities. Although arthroscopic excision is commonly performed in symptomatic patients with popliteal cyst, our study suggests that such cysts and bursitis may not be the primary etiology for symptoms of OA, although this needs to be confirmed by future longitudinal studies.

The strengths of the present study lie in a large sample with knee radiographic and MRI measurements. There are a number of potential limitations in this study. First, this is a cross-sectional study, precluding our ability to draw conclusions about causal relationships between cysts or bursitis and the structural abnormalities. Nevertheless, the existence of the consistent relationships between cysts and many intra-articular pathologies of OA suggests that those cystic lesions may be involved in symptomatic and structural changes in OA. This is supported by a recent study, which confirmed that removal of the popliteal cyst and associated pathology would provide improvements in function and pain of the knee after surgery, and in patient satisfaction [34]. Second, we only assessed two cystic lesions in the current study. Other cystic lesions were not as prevalent as these two lesions [6] and low prevalence would disallow us to observe significant associations between other cystic lesions and outcome measures. Third, the response rate at baseline was 57%, possibly due to the demands on study participants that each visit took three hours. However, there were no significant differences in age and gender between those who responded and those who did not. Fourth, although we studied a well-characterized population of older adults with a high level of knee pain (46%), this randomly selected sample unavoidably included subjects with other diseases, which may have affected the associations. Nevertheless, the results were largely unchanged when the analysis was adjusted for disease status or the subjects with other diseases were excluded. Last, measurement error may influence results; however, given all measures (for example, cystic lesions, cartilage defect and BML) were highly reproducible, this is considered unlikely.

Conclusions
This cross-sectional study documents that popliteal cyst and subgastrocnemius bursitis are associated with radiographic and MRI-detected joint structural abnormalities as well as knee symptoms. Longitudinal data will help resolve if they are a consequence or a cause of knee joint abnormalities.

Abbreviations
BMI: body mass index; BML: bone marrow lesion; ICC: interclass correlation coefficient; IL: interleukin; JSM: joint space narrowing; MRI: magnetic resonance imaging; OA: osteoarthritis; OARSI: Osteoarthritis Research Society International; RA: rheumatoid arthritis; TASOAC: Tasmania Older Adult Cohort; TNF: tumour necrosis factor; WOMAC: Western Ontario McMaster Universities osteoarthritis index; WORMS: whole-organ magnetic resonance imaging score.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
CD and YC conceived the study. CD, GJ and FC participated in its design and coordination. YC, WH, WW and BA performed MRI measurements. YC, GJ and CD drafted the manuscript. All authors revised the manuscript and gave final approval of the version to be submitted.

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Author details
1Menzies Research Institute Tasmania, University of Tasmania, Private Bag 23, Hobart, Tasmania 7000, Australia. 2Research Institute of Orthopaedics, Shuguang Hospital Affiliated to Shanghai University of Traditional Chinese Medicine, 528 Zhangheng Road, Shanghai 2003, China. 3Department of Epidemiology and Preventive Medicine, Monash University, Melbourne 3004, Australia.

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References
1. Felson DT: Clinical practice. Osteoarthritis of the knee. N Engl J Med 2006, 354:841–848.
2. Handy Jr: Popliteal cysts in adults: a review. Semin Arthritis Rheum 2001, 31:108–118.
3. Beaman FD, Peterson JJ: MR imaging of cysts, ganglia, and bursae about the knee. Magn Reson Imaging Clin N Am 2007, 15:39–52.
4. Baker WM: On the formation of synovial cysts in the leg in connection with disease of the knee-joint. Clin Orthop Relat Res 1944, 187:2–10.
5. Marti-Bonmati L, Molla E, Dosda R, Casillas C, Ferrer P: MR imaging of Baker cysts – prevalence and relation to internal derangements of the knee. MAGMA 2000, 10:205–210.
6. Hayashi D, Roemer FW, Dhina Z, Kwoh CK, Hannon MJ, Moore C, Guermazi A: Longitudinal assessment of cyst-like lesions of the knee and their relation to radiographic osteoarthritis and MRI-detected effusion and synovitis in patients with knee pain. Arthritis Res Ther 2010, 12:R72.
7. Marra MO, Crema MD, Chung M, Roemer FW, Hunter DJ, Zaim S, Diaz L, Guermazi A: MRI features of cystic lesions around the knee. Knee 2008, 15:423–438.
8. Fritschy D, Fasel J, Imbert JC, Bianchi S, Verdonk R, Wirth CJ: The popliteal cyst. Knee Surg Sports Traumatol Arthrosc 2006, 14:623–628.
9. Miller TT, Saron RB, Kohlmeiser T, Levin TL, Feldman F: MR imaging of baker cysts: association with internal derangement, effusion, and degenerative arthropathy. Radiology 1996, 201:247–250.
10. Hill CI, Gale DG, Chaisson CE, Skinner K, Kazis L, Gale ME, Felson DT: Knee effusions, popliteal cysts, and synovial thickening: association with knee pain in osteoarthritis. J Rheumatol 2001, 28:1330–1337.
11. Peterfy CG, Guermazi A, Zaim S, Timran PF, Mlaux Y, White D, Kothari M, Lu Y, Fye K, Zhao S, Genant HK: Whole-organ magnetic resonance imaging score (WORMS) of the knee in osteoarthritis. Osteoarthritis Cartilage 2004, 12:177–190.
12. Ding C, Ciucurini F, Parameswaran V, Burgess J, Quinn S, Jones G: Serum levels of vitamin D, sunlight exposure, and knee cartilage loss in older...
adults: the Tasmanian older adult cohort study. *Arthritis Rheum* 2009, 60:1381–1389.

13. Ding C, Gamero P, Cicuttini F, Scott F, Cooley H, Jones G. Knee cartilage defects: association with early radiographic osteoarthritis, decreased cartilage volume, increased joint surface area and type II collagen breakdown. *Osteoarthritis Cartilage* 2005, 13:198–205.

14. Dore O, Quinn S, Ding C, Wisenbaker T, Jones G. Correlates of subchondral BMD: a cross-sectional study. *J Bone Miner Res* 2007–2015, 2009:24.

15. Bellamy N, Buchanan WW, Goldsmith CH, Campbell J, Stitt LW. Validation study of WOMAC: a health status instrument for measuring clinically important patient relevant outcomes to antirheumatic drug therapy in patients with osteoarthritis of the hip or knee. *J Rheumatol* 1833–1840, 1988:15.

16. Stratford PW, Kennedy DM, Woodhouse LJ, Spadoni GF. Measurement properties of the WOMAC LK 3.1 pain scale. *Osteoarthritis Cartilage* 2007, 15:266–272.

17. Altman RD, Gold GE. Atlas of individual radiographic features in osteoarthritis, revised. *Osteoarthritis Cartilage* 2007, 15:A1–56.

18. Reginster JY, Deray R, Rovati LC, Lee RI, Lejeune E, Bruyere O, Gaccioulli G, Henrotin Y, Dacce JE, Gossett C. Long-term effects of glucosamine sulphate on osteoarthritis progression: a randomised, placebo-controlled clinical trial. *Lancet* 2001, 357:251–256.

19. Bickel WH, Burleson RJ, Dahlin DC. Popliteal cyst: a clinicopathological survey. *J Bone Joint Surg Am* 1956, 38-A:1265–1274.

20. Sansonne V, De Ponti A. Arthroscopic treatment of popliteal cyst and associated intra-articular knee disorders in adults. *Arthroscopy* 1999, 15:368–372.

21. Rupp S, Seil R, Jochum P, Kohn D. Popliteal cysts in adults. Prevalence, associated intraarticular lesions, and results after arthroscopic treatment. *Am J Sports Med* 2002, 30:112–115.

22. Ding C, Cicuttini F, Jones G. Tibial subchondral bone size and knee cartilage defects: relevance to knee osteoarthritis. *Osteoarthritis Cartilage* 2007, 15:479–486.

23. Cicuttini F, Ding C, Wu A, Davis S, Ebeling PR, Jones G. Association of cartilage defects with loss of knee cartilage in healthy, middle-age adults: a prospective study. *Arthritis Rheum* 2003–2039, 2005:52.

24. Carnes J, Stannus O, Cicuttini F, Ding C, Jones G. Knee cartilage defects in a sample of older adults: natural history, clinical significance and factors influencing change over 2.9 years. *Osteoarthritis Cartilage* 2012, 20:1541–1547.

25. Davies-Tuck ML, Wluka AE, Wang Y, English DR, Giles GG, Cicuttini F. The natural history of bone marrow lesions in community-based adults with no clinical knee osteoarthritis. *Ann Rheum Dis* 2009, 68:904–908.

26. Felson DT, Chaisson CE, Hill CL, Torreman SM, Gale ME, Skinner KN, Kazis L, Gale DR. The association of bone marrow lesions with pain in knee osteoarthritis. *Ann Intern Med* 2001, 134:541–549.

27. Ding C, Cicuttini F, Scott F, Cooley H, Boon C, Jones G. Natural history of knee cartilage defects and factors affecting change. *Arch Intern Med* 2006, 166:651–658.

28. Amin S, LaValley MP, Guermazi A, Grigoryan M, Hunter DJ, Clancy M, Niu J, Gale DR, Felson DT. The relationship between cartilage loss on magnetic resonance imaging and radiographic progression in men and women with knee osteoarthritis. *Arthritis Rheum* 2005, 52:3152–3159.

29. Mundermann A, Dyrby CO, Andriacchi TP. Secondary gait changes in patients with medial compartment knee osteoarthritis: increased load at the ankle, knee, and hip during walking. *Arthritis Rheum* 2005, 52:2835–2844.

30. Stannus OP, Jones G, Blizzard L, Cicuttini FM, Ding C. Associations between serum levels of inflammatory markers and change in knee pain over 5 years in older adults: a prospective cohort study. *Ann Rheum Dis* 2013, 72:535–540.

31. Bakker AC, van de Loo FA, van Beuningen HM, Sime P, van Lent PL, van der Kraan PM, Richards CD, van den Berg WB. Overexpression of active TGF-beta 1 in the murine knee joint: evidence for synovial-layer-dependent chondro-osteocyte formation. *Osteoarthritis Cartilage* 2001, 9:128–136.

32. Chatzopoulos D, Moralidis E, Markou P, Makri V, Ardos G. Baker's cysts in knees with chronic osteoarthritic pain: a clinical, ultrasonographic, radiographic and scintigraphic evaluation. *Rheumatol Int* 2008, 29:141–146.

33. Torreggiani WC, Al-Ismael K, Munk PL, Roche C, Keogh C, Nicolaou S, Marchinkow LP. The imaging spectrum of Baker's (popliteal) cysts. *Clin Radial* 2002, 57:681–691.

34. Malinowski K, Synder M, Siibinski M. Selected cases of arthroscopic treatment of popliteal cyst with associated intra-articular knee disorders primary report. *Ortop Traumatol Rehabil* 2011, 13:573–582.