Patient characteristics associated with a poor response to non-surgical multidisciplinary management of knee osteoarthritis: a multisite prospective longitudinal study in an advanced practice physiotherapist-led tertiary service

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ABSTRACT

Objectives To explore patient characteristics recorded at the initial consultation associated with a poor response to non-surgical multidisciplinary management of knee osteoarthritis (KOA) in tertiary care.

Design Prospective multisite longitudinal study.

Setting Advanced practice physiotherapist-led multidisciplinary orthopaedic service within eight tertiary hospitals.

Participants 238 patients with KOA.

Primary and secondary outcome measures Standardised measures were recorded in all patients prior to them receiving non-surgical multidisciplinary management in a tertiary hospital service across multiple sites. These measures were examined for their relationship with a poor response to management 6 months after the initial consultation using a 15-point Global Rating of Change measure (poor response (scores −7 to +1)/positive response (scores +2 to +7)). Generalised linear models with binomial family and logit link were used to examine which patient characteristics yielded the strongest relationship with a poor response to management as estimated by the OR (95% CI).

Results Overall, 114 out of 238 (47.9%) participants recorded a poor response. The odds of a poor response increased with higher patient expectations of benefit (OR 0.74 (0.63 to 0.87) per 1/10 point score increase) and higher self-reported knee function (OR 0.67 (0.51 to 0.89) per 10/100 point score increase) (p<0.01). The odds of a poor response increased with a greater degree of varus frontal knee alignment (OR 1.35 (1.03 to 1.78) per 5° increase in varus angle) and a severe (compared with mild) radiological rating of medial compartment degenerative change (OR 3.11 (1.04 to 9.3)) (p<0.05).

Conclusions These characteristics may need to be considered in patients presenting for non-surgical multidisciplinary management of KOA in tertiary care. Measurement of these patient characteristics may potentially better inform patient-centred management and flag the need for judicious monitoring of outcome for some patients to avoid unproductive care.

INTRODUCTION

Knee osteoarthritis (KOA) is a leading cause of disability in the community. Patients with more advanced pain and functional limitations due to KOA may be referred to tertiary hospitals for specialist opinion regarding
management options, but for many of these patients, non-surgical multidisciplinary management is the first line of care. While evidence supports non-surgical interventions such as exercise and weight loss for the management of KOA, individual patient responses may vary. It would be clinically and economically advantageous to identify patients with KOA at risk of a poor response to non-surgical multidisciplinary management at the initial consultation. Potentially at-risk patients could receive additional care to address identified risk factors, and their treatment response closely monitored to facilitate a timely referral for surgical consultation if not responding. However, it is crucial that studies are specific to patient populations, service settings and intervention types. Potentially interpreting the meaning of risk factors may be misguided when models are applied to different patients in different health settings (eg, primary vs tertiary) where marked variation in patient demographics and interventions may exist.

Our previous retrospective audit of medical records explored patient characteristics associated with a poor response to non-surgical multidisciplinary management of KOA. This audit was conducted within a multisite advanced practice physiotherapist-led multidisciplinary orthopaedic service (herein referred to as the ‘service’) in tertiary hospitals. The service employs experienced musculoskeletal physiotherapists (service leader) to assess and determine management pathways for non-urgent patients within orthopaedic specialist outpatient departments. Where non-surgical management is appropriate, care provided by the service is patient centred with an emphasis on progression from supported to self-managed and is multidisciplinary (as required; physiotherapy, occupational therapy, dietetics and/or psychology) to pragmatically address the mix of biopsychosocial factors potentially underlying each patient’s KOA presentation.

The retrospective audit showed that patients who reported lower knee function and higher levels of anxiety at their initial appointment within the service were at greater odds of a poor response to non-surgical multidisciplinary management. There were marked limitations of this prior study due to its retrospective nature and limited patient characteristics recorded. In particular, the literature suggests that many other potentially relevant risk factors should be considered such as patient expectations and physical examination findings of the knee, which were previously not examined due to a lack of standardised recording within medical records.

To address limitations of our previous retrospective audit, this study prospectively explored the relationship between a broader selection of patient characteristics and a poor outcome to non-surgical multidisciplinary management of KOA in the service. Based on our previous findings and others, we hypothesised that a mix of patient-reported, physical and radiological patient characteristics may be associated with a poor outcome.

METHODS
Study design
A prospective longitudinal study was conducted across eight tertiary hospital service sites in Queensland, Australia. Service leader engagement ensured selected measures and equipment (eg, questionnaires, goniometers, radiological findings) were accessible across sites. This manuscript follows the Strengthening the Reporting of Observational Studies in Epidemiology reporting guidelines.

Patient and public involvement
No patient involved.

Participants
Participants were recruited within the service at participating hospitals between January 2014 and December 2016 (follow-up completed July 2017). Potential participants had been triaged from the specialist orthopaedic outpatient waiting list by the medical consultant and/or service leader to undergo non-surgical management within the service for their knee condition. At the initial service consultation, potential participants were further screened for study eligibility based on an accepted clinical and radiological diagnosis of KOA. Specifically, participants had to report persistent knee pain (pain on most days for at least a month or longer in the past year), limited morning stiffness (less than 30 min) and functional limitations (self-reported functional impairment, eg, knee muscle weakness), as well as demonstrate radiological evidence of KOA on imaging (determined from a radiological report or medical consultant notes within the medical records). Participants were excluded if they presented with a potentially serious medical condition (red flags), knee pain of lumbar spine or hip origin, an active inflammatory condition (eg, rheumatoid arthritis), severe symptoms likely to be aggravated by therapies, significant or unstable neurovascular involvement, or if the referring medical practitioner or patient requested specialist medical consultation specifically.

Criteria for a poor response to non-surgical multidisciplinary management (dependent variable)
A poor response to management for each patient case was determined with a 15-point Global Rating of Change (GROC). Patients rated their overall perception of change since beginning treatment on a scale ranging from −7 (a very great deal worse) to zero (about the same) to +7 (a very great deal better). Investigators dichotomised (binary outcome) responses between −7 (a very great deal worse) and +1 (almost the same, hardly any better at all) as a poor response, and responses between +2 (a little better) and +7 (a very great deal better) as a positive response. This decision was based on the generally high level of KOA presentation severity seen within this specialty tertiary service, where marginal improvements in baseline function may for some patients be considered a positive response. Therefore, scores between −7 and
+1 representing worsening or negligible change in the disorder were considered a poor response. This outcome was assessed at 6 months after entry to the service as this is a conservative timeframe for decision-making regarding the potential impact of non-surgical management for these patients based on our previous study.

**Potential explanatory patient variables assessed for their relationship with outcome (independent variables)**

Patient characteristics recorded at the initial consultation with the service leader included routine demographic and clinical information from the patient interview, self-report questionnaires capturing condition specific, as well as general health and psychosocial information (that had been premailed and completed by patients prior to the initial session as is the standard procedure in these clinics), information from standardised physical tests and radiological imaging findings. In cases of a referral for bilateral KOA, participants were asked to nominate the most severe/problematic knee, which, for the purposes of this study, was recorded as the affected knee.

Variables included:

**Demographic and social measures:** This includes age (years), sex (% male), English first language (% yes), education level (% yes; school incomplete, completed secondary school, completed Technical and Further Education/trade/university), work status (% yes; employed (full or part-time), unemployed, retired), marital status (% yes; married/defacto, single), dependents (% yes).

**General and global health measures:** This includes body mass index (k/m²), disability benefits (% yes), smoking status (% yes), comorbidities (number/18 listed conditions), total body pain areas (number/18 segmented body chart), health-related quality of life (utility score/1) measured with the Assessment of Quality of Life Questionnaire (AQoL-6D).[^17]

**Psychological measures:** General psychological distress was evaluated with the Depression, Anxiety and Stress Scale (DASS-21) with each of the three dimensions of depression (score/42), anxiety (score/42) and stress (score/42) scored separately.[^18] Pain-related psychological measures included the Pain Self-Efficacy Questionnaire (score/60) and the Örebro Musculoskeletal Pain Screening Questionnaire (ÖMPSQ–short) (score/100).[^20] The patient’s expectation of benefit (score/10) from the non-surgical management approach was also recorded using a visual analogue scale anchored by the terms ‘No benefit’ and ‘Extreme benefit’.

**Condition-specific symptoms and signs:** The Knee Injury and Osteoarthritis Outcome Score (KOOS) including separate scores for symptoms and stiffness (score/100), pain (score/100), function and daily living (score/100) and quality of life (score/100).[^21][^22] Participant-nominated functional deficits were evaluated with the Patient Specific Functional Scale (score/10).[^23][^24] The potential presence of neuropathic pain was evaluated with the Leeds Assessment of Neuropathic Symptoms and Signs (S-LANSS) pain scale (% yes, score of ≥12 points).[^25] Other self-reported information regarding the condition included symptom duration (% yes; >12 months), traumatic onset (% yes), previous surgery (% yes) for the same condition and the presence of associated symptoms such as bilateral knee pain (% yes), instability/giving way (% yes), true locking (% yes), swelling (% yes) and joint noises (% yes).

Physical examination measures conducted by the advanced physiotherapist included knee flexion and extension (lack) range of motion (°) measured with a goniometer, observed lateral thrust of the affected knee during the examination of gait (% yes) and a reliable weight-bearing frontal knee alignment measure in standing using a goniometer. In this method, the goniometer axis is positioned over the centre of the patella with arms aligned to the anterior superior iliac spine and the centre of the ankle joint with the resultant knee angle recorded as either a valgus (−°), neutral (0°) or varus (+°) angle. Findings from four knee ligament stress tests (valgus, varus, anterior, posterior) were judged (normal or altered) and individually recorded (% altered). A measure of symptom modification during the initial examination was also evaluated. In this test, patients rated any perceived improvement (%) (taking all things into consideration such as pain, mobility, effort) in the replication of a self-nominated problematic activity (eg, squatting), comparing the overall experience before and then during a modification of the activity guided by the therapist (eg, corrected movement pattern instructions, manual assistance from therapist, use of strapping tape).

Up to three modification trials were permitted with improvement needing to be repeatable over at least two repetitions.

Radiological findings (X-ray, MRI, CT) of degenerative change in the medial, lateral and patellofemoral compartments of the knee were recorded based on either a specific radiological report or radiological findings specified in the medical notes by the medical consultant. Degenerative change was recorded as either absent, mild, moderate or severe. This approach was based on consensus by the service leaders that these ratings were consistently used in radiological reporting for imaging across all sites. Relevance of the imaging with regard to how recently it was taken was at the service leader’s discretion.

**Procedure**

All potential explanatory patient characteristics were recorded during the initial consultation with the service leader following screening and consent procedures. Participants then underwent their normal care within the service at the respective sites. Each patient’s management (duration of management period, disciplines consulted) was pragmatically based on the initial examination findings of the service leader and the clinical discretion of the involved discipline-specific treatment providers. The study research officer contacted the participant at the 6-month follow-up period and mailed them the clinical outcome measure (GROC) independently of the service at all sites. Participants may or may not have been still
receiving management within the service at this time point, according to their individual treatment plan.

Sample size estimation
It was estimated that 224 cases would be required to identify explanatory variables with a strong relationship to a poor response based on an expected 50% non-responder rate, the inclusion of up to seven explanatory variables in a final multivariable (logistic family) model and a conservative number of non-responders (n=16) per variable based on prior simulations. To account for up to 15% dropout, it was estimated that a minimum number of 264 participants were required to be recruited for this study.

Statistical analysis
Analyses were conducted using Stata V.13 (StataCorp, 2013, Stata Statistical Software: Release 13, College Station, Texas, USA: StataCorp LP). Participant characteristics and outcomes were described using conventional descriptive statistics with number (%) for categorical data, mean (SD) for normally distributed data and median (IQR) for non-normally distributed data. Generalised linear models with binomial family and logit link with site as a random effect were used to examine the association between potential explanatory variables with patients being classified as either having a poor response (GROC score ≤1) or positive response (GROC score 2–7) to multidisciplinary non-surgical management in this sample. To select the most appropriate explanatory variables to carry forward to a multivariable model, univariate analyses were initially conducted, and an unadjusted OR (95% CI) was calculated for each variable’s relationship with a poor response. Due to the high number of potential explanatory variables and the study’s intention to identify a clinically relevant and preferably quantifiable parsimonious selection of explanatory variables associated with a poor response, only explanatory variables that displayed a univariate relationship of p≤0.1 with the reference score (GROC score ≤1) were considered for the multivariable analyses. Eligible variables were then further screened as to their relative level of potential clinical impact and priority to take to the final model. This was particularly the case if multiple variables from a similar domain (eg, psychological measures, physical measures) were eligible. Potential multicollinearity issues between eligible explanatory variables were then evaluated with a Spearman’s rho (rs) correlational coefficient as it is appropriate for both continuous and discrete ordinal variables. If variables were shown to have significant moderate (rs=0.4–0.6) or strong (rs=0.7–0.9) correlations, only one variable was selected (investigators choice based on clinical reasoning) to be carried forward for the final model. After accounting for potential non-independence associated with site by including site as a random effect, and the aforementioned variable selection process to avoid multicollinearity, we did not detect any breaches of model assumptions in the final model. The c-statistic was also calculated for the final model, and in the final model, ORs were expressed per 5° for range of motion variables, per 10 points on the KOOS and per 10% for symptom modification test. In addition, to examine whether findings were robust against this variable selection process which included investigator judgement, any correlated variables that were not selected were substituted into the multivariable analysis in place of the selected variable as sensitivity analyses to determine if findings were impacted by the choice of included variable.

RESULTS
The flow of participants through the study is depicted in figure 1. Of the 375 patients deemed eligible following screening and invited to participate, 286 participants completed written consent forms and baseline measures. At the 6-month follow-up period, GROC measures were received from 238 participants representing a 17% dropout rate; however, exploratory independent t-tests suggest that those who did/did not complete their outcomes were not different in their baseline level of knee disability (KOOS Function—Daily Living, p=0.43), suggesting the completers were a representative sample.

For the 238 patients who completed the 6-month follow-up GROC measure, the median (IQR) GROC score was +2 (–2 to +4) with the range of scores from –7 to +7. Overall, 114 (47.9%) patients fitted the criteria as a poor responder. The characteristics of participants classified as poor responders or positive responders are described in table 1. Twenty potential explanatory variables demonstrated a univariate relationship (p≤0.1) with a poor response to

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Figure 1  Participant flow diagram for patients with knee osteoarthritis (KOA) and Global Rating of Change (GROC) completions.
Table 1  Means (±SD) (normally distributed continuous data), medians (IQR) (non-normally distributed continuous data) and row participant percentages (n) (categorical data) for the independent variables grouped for participants dichotomised as a poor response (GROC ≤+1) or a positive response (GROC ≥+2) to the non-surgical multidisciplinary management of KOA

| Variables                                      | Poor response (n=114) | Positive response (n=124) | Unadjusted OR (95% CI) | P value |
|------------------------------------------------|-----------------------|---------------------------|------------------------|---------|
| **Demographic and social measures**           |                       |                           |                        |         |
| Age (years)                                   | 63.45±9               | 62.27±12.17               | 1.01 (0.99 to 1.04)    | 0.31    |
| Sex: Male (% yes (n))                         | 54% (46)              | 46% (40)                  | 1.5 (0.87 to 2.6)      | 0.15    |
| English first language (% yes (n))            | 49% (110)             | 51% (116)                 | 2.03 (0.57 to 7.22)    | 0.28    |
| **Education level**                           |                       |                           |                        |         |
| School incomplete (% yes (n))                 | 52% (53)              | 48% (49)                  | Referent               |         |
| Completed secondary school (% yes (n))        | 40% (23)              | 60% (34)                  | 0.64 (0.33 to 1.24)    | 0.18    |
| Completed TAFE/Trade/ University (% yes (n))  | 48% (33)              | 52% (36)                  | 0.88 (0.47 to 1.64)    | 0.68    |
| **Work status**                               |                       |                           |                        |         |
| Employed (% yes (n))                          | 48% (54)              | 52% (59)                  | Referent               |         |
| Unemployed (% yes (n))                        | 50% (5)               | 50% (5)                   | 1.04 (0.28 to 3.85)    | 0.95    |
| Retired (% yes (n))                           | 45% (44)              | 55% (53)                  | 0.89 (0.51 to 1.56)    | 0.69    |
| Marital status: Married/de facto (% yes (n))  | 51% (64)              | 49% (62)                  | 1.34 (0.79 to 2.26)    | 0.28    |
| Dependents (% yes (n))                        | 38% (11)              | 62% (18)                  | 0.62 (0.28 to 1.38)    | 0.25    |
| **General and global health measures**        |                       |                           |                        |         |
| Body mass index (kg/m²)                       | 34.84±7.7             | 33.68±8.22                | 1.02 (0.98 to 1.05)    | 0.35    |
| Disability benefits (% yes (n))              | 46% (22)              | 54% (26)                  | 0.83 (0.43 to 1.61)    | 0.58    |
| Smoking status (% yes (n))                   | 57% (17)              | 43% (13)                  | 1.44 (0.65 to 3.17)    | 0.36    |
| Comorbidities (number)                        | 2 (1–3)               | 2 (1–3)                   | 1.03 (0.89 to 1.2)     | 0.65    |
| Total body pain areas (number/18 body regions)| 5 (3–8)               | 4 (2–6)                   | 1.1 (1.01 to 1.19)     | 0.02*   |
| Quality of Life (utility score/1)             | 0.63 (0.45–0.74)      | 0.69 (0.51–0.8)           | 0.2 (0.05 to 0.82)     | 0.03*   |
| **Psychological measures**                    |                       |                           |                        |         |
| DASS-21                                        |                       |                           |                        |         |
| Depression score (score/42)                   | 6 (2–16)              | 6 (2–14)                  | 1.02 (0.99 to 1.05)    | 0.19    |
| Anxiety score (score/42)                      | 6 (2–14)              | 4 (0–10)                  | 1.04 (1.01 to 1.08)    | 0.03*   |
| Stress score (score/42)                       | 10 (2–16)             | 8 (2–14)                  | 1.01 (0.98 to 1.04)    | 0.56    |
| **Condition specific symptoms and signs**     |                       |                           |                        |         |
| KOOS                                           |                       |                           |                        |         |
| Symptoms and stiffness (score/100)            | 39.52±17.24*          | 51.05±18.8                | 0.97 (0.95 to 0.98)    | <0.001* |
| Pain (score/100)                              | 38.02±16.02*          | 50.41±16.21               | 0.95 (0.94 to 0.97)    | <0.001* |
| Function—Daily Living (score/100)             | 45.72±16.13*          | 55.97±16.31               | 0.96 (0.94 to 0.98)    | <0.001* |
| Quality of life (score/100)                   | 22.8±17.14*           | 32.31±16                  | 0.97 (0.95 to 0.98)    | <0.001* |
| Patient Specific Functional Scale (score/10)  | 3.83±1.74             | 4.45±1.56                 | 0.79 (0.67 to 0.94)    | 0.01*   |
| S-LANSS pain score (% ≥12 points (n))         | 51% (30)              | 49% (29)                  | 1.24 (0.67 to 2.27)    | 0.49    |
| Symptom Duration (% >12 months (n))           | 53% (94)              | 47% (85)                  | 2.02 (1.08 to 3.78)    | 0.03*   |
| Traumatic onset (% yes (n))                   | 44% (40)              | 56% (51)                  | 0.76 (0.44 to 1.3)     | 0.32    |
| Previous surgery (% yes (n))                  | 51% (29)              | 49% (28)                  | 1.19 (0.65 to 2.19)    | 0.57    |
| Associated symptoms                           |                       |                           |                        |         |
| Bilateral knee pain (% yes (n))               | 51% (35)              | 49% (33)                  | 1.16 (0.63 to 2.14)    | 0.63    |
| Instability/giving way (% yes (n))            | 54% (56)              | 46% (47)                  | 1.54 (0.89 to 2.66)    | 0.13    |
| True locking (% yes (n))                      | 50% (10)              | 50% (10)                  | 1.05 (0.41 to 2.67)    | 0.92    |
| Swelling (% yes (n))                          | 51% (49)              | 49% (48)                  | 1.15 (0.68 to 1.97)    | 0.6     |

Continued
non-surgical management of KOA, as shown in table 1. Of these 20 variables, 9 were retained in the final multivariable model following exclusion of variables based on likely clinical utility/priority and potential multicollinearity (or both). The specific rationale for excluding the following variables was KOOS subscales (symptoms and stiffness; pain; quality of life) were moderate to strongly correlated with the KOOS Function—Daily Living subscale (rs=0.6–0.8, p<0.001) which was retained as it was considered most clinically informative based on its measure of pain-related function; PSFS was excluded as it was moderately correlated (r=0.4, p<0.001) with the KOOS Function—Daily Living subscale which already provided a measure of function; the QOL, PSEQ and ÖMPSQ measures were moderately correlated with the KOOS Function—Daily Living subscale (r=0.4–0.5, p<0.001); the DASS Anxiety subscale was excluded as findings for the poor responder group (median score 6/42 points) was below the lowest end of the mild anxiety category (8–9 points);18; flexion knee range was excluded on the basis of the very small difference between poor/positive responders and negligible OR (95% CI) (OR 0.97–1); the quantifiable frontal knee alignment measure was maintained instead of the clinician-rated measures of gait lateral thrust and valgus stress test. Additionally, findings of the valgus stress test were considered to reflect pseudolaxity associated with medial compartment degenerative change as supported by their weak but significant correlation (r=0.17, p<0.02).
In addition, when the variables that were not selected (eg, other subscales of KOOS, ÖMPSQ, PSEQ) were substituted into the multivariable model as sensitivity analyses, findings were consistent regardless of which of the variables were used (not presented).

Of the nine variables retained in the final model, only four were observed to be independently associated with a poor response to the non-surgical management of KOA, as shown in table 2. Patient characteristics associated with a lower odds of a poor response included higher patient expectations of benefit with the OR 0.74 (0.63 to 0.87) per 1/10 point score increase (p<0.001), indicating the odds of a poor response reduces with higher patient expectations; higher self-reported knee function (KOOS Function—Daily Living) with the OR 0.67 (0.51 to 0.89) per 10/100 point score increase (p<0.01), indicating the odds of a poor response reduced with higher self-reported knee function. Patient characteristics associated with greater odds of a poor response included greater varus frontal knee alignment in standing with the OR 1.35 (1.03 to 1.78) per 5° increase in varus angle (p=0.03), suggesting increased odds of a poor response with greater recording of knee varus angle; and reported severe medial compartment degenerative change on imaging with the OR 3.11 (1.04–9.3) (p=0.04), suggesting a severe rating on imaging is associated with greater odds of a poor response than a reported mild degenerative rating. Service site was not a significant factor in the model (p=0.64).

**DISCUSSION**

Study findings support our hypothesis that a mix of patient characteristics including cognitive (expected benefit of management, perceived daily function), physical (frontal knee alignment) and radiological (degenerative severity) characteristics may be associated with a poor response to non-surgical multidisciplinary management of KOA within the tertiary care service under study. Collectively, these findings describe poor responders within a biopsychosocial context as generally being more severely disabled, matched by more severe knee pathology, and more negative outlook regarding their likely capacity to benefit from non-surgical management. These findings provide clinicians within these tertiary settings some scope to identify patients at the initial consultation who may be more unlikely to benefit from this line of care. This knowledge provides opportunity for early management planning that may include more finely tailored non-surgical interventions to specifically address identified risk factors, or earlier referral for specialist medical consultation if not benefiting.

We evaluated the patient’s perceived expected benefit from the recommended multidisciplinary management using a simple visual analogue scale. With higher patient
expectation of benefit, the odds of patients reporting a poor response to management is reduced (OR 0.74 (0.63 to 0.87) per 1 point increase). These findings are consistent with previous studies in KOA and other conditions showing negative/positive patient expectations about treatment benefits to influence outcomes. An individual’s cognitive actions in response to recommended treatments are thought to reflect his or her beliefs about the potential benefits of recommended management approaches, potentially impacting his or her motivation to engage in interventions. Older adults with KOA have been reported to have low expectations of benefit from non-surgical interventions such as exercise. Patient expectancy may be shaped by multiple factors including psychological variables such as fear avoidance, disease severity and X-ray findings, confidence in exercise capacity, and pain and disability level. Lower reported knee function at initial consultation was associated with a poorer response in both this current study and our previous study in KOA. This current study suggests that the odds of a poor response reduced (OR 0.67 (0.51 to 0.89) per 10 point increase) with higher Daily Knee Function (KOOS) score signifying a better level of knee function. While lower perceived knee function may simply reflect a more severe disorder, cognitively it may also reduce the patient’s expectations that an activity-based management approach (such as that provided by the multidisciplinary service under study) will be beneficial. While factors influencing patient expectations vary between individuals, findings support recommendations for determining patient expectations in the initial examination as it may impact patient–therapist communication. Skilled clinicians may communicate management options with patients engaging them on a personal level, collaboratively problem solving personal perceived barriers to recovery and providing reassurance, and ultimately improving the patients’ motivation to participate in recommended management approaches.

Patients with a severe radiological rating of medial knee compartment degeneration had, on average, greater than three times the odds (OR 3.11 (1.04 to 9.3)) of reporting a poor response to management compared with those with a mild rating (table 2). Although the wide CIs suggest this relationship may be variable between patients. Traditionally, radiological degenerative severity has not been shown to be strongly associated with knee symptom level and functional impairment. However, more recent studies, including one incorporating five different racial/ethnic populations, have shown radiographic knee osteoarthritis to be strongly associated with the presence and severity of knee pain. This current study lends support to this more recent literature with a more severe level of medial knee degeneration increasing the odds of a poorer response. However, this was only for the medial compartment, as the severity of degenerative change was not related to response for the lateral or patellofemoral knee compartments. Furthermore, consistent with the wide 95% OR CIs for this measure (1.04–9.3), table 1 shows that 24% of participants who recorded a positive response to management also had a severe rating of medial degenerative change. This is important in the interpretation of findings as a severe rating of medial knee degeneration should not exclude a patient from a trial of non-surgical multidisciplinary management.

The radiological findings also complement the observed relationship between greater varus knee angle and poorer outcome. Degenerative changes and varus malalignment are thought to affect knee biomechanics, load distribution and quadriceps muscle strength, although the relationship between these variables and their impact on knee function is not entirely clear. We included the frontal knee alignment measure because of literature suggesting local knee mechanics may impact the therapeutic effect of interventions such as exercise. Our findings support this proposition. For every 5° increase in varus knee angle measured at the initial examination, the odds of later reporting a poor response increased notably (OR 1.35 (1.03 to 1.78)) (table 2). This rudimentary clinical measure of frontal plane knee alignment (recorded using a goniometer in standing) has been previously shown to be significantly related to radiological measures of alignment (r=0.67, p<0.0001). Although this clinical measure of frontal knee alignment is obviously influenced by other factors than just radiological knee alignment, it provides an accessible method of evaluating a potentially valuable aspect of frontal knee mechanics associated with a patient’s response to management, which does not rely on costly and probably clinically inaccessible motion analysis or radiological measures.

Limitations
This study has limitations. The 17% dropout rate from baseline may have affected results, although baseline levels of knee disability were similar between completers/non-completers. Findings are also specific to patients within the service studied, patient characteristics recorded and outcome used (ie, GROC). Patients referred to this tertiary care service may represent those at the more severe end of the clinical spectrum and findings may be different in patients managed in other settings (eg, primary care). The large number of service leaders collecting measures across sites potentially increased the likelihood of measurement error, but we attempted to minimise this by providing a training manual and training session before the commencement of the study. Radiological findings were limited to clinical descriptors (absent, mild, moderate, severe, degenerative change) to ensure measurement consistency across imaging types and service sites. Other radiological variables (eg, meniscus tear) requiring imaging techniques other than X-ray were not included as they were inconsistently available (ie, X-ray findings noted in 83% of participants, compared with MRI (21%) and CT (2%)). Other patient characteristics (eg, medication use) may have been relevant but were beyond the scope of this study to record in enough detail to not be potentially misleading.
CONCLUSIONS
This study identified four patient characteristics assessed at the initial consultation associated with a poor response to non-surgical multidisciplinary management of KOA in a tertiary care service. While low patient expectations, low self-reported knee function, a greater knee varus angle and severe radiological knee degeneration do not nullify a trial of non-surgical multidisciplinary management, their presence may signal to clinicians the need to further individualise care and judiciously monitor outcomes to avoid unproductive use of health resources and burden to the patient.

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Acknowledgements The investigators would like to thank the clinical and administrative staff of the Orthopaedic Physiotherapy Screening Clinic and Multidisciplinary Services, as well as the Physiotherapy Departments at the participating hospital sites; Royal Brisbane and Women’s Hospital, Ipswich Hospital, Mackay Hospital, Mater Hospital, The Prince Charles Hospital, Toowoomba Hospital, Townsville Hospital and Redcliffe Hospital.

Contributors SOL, MR, PSK, TC, MCot, AK, BV and SM contributed to the study design and preparation, data collection planning and process, data analysis and interpretation. PW, PSK, BE, IM, WOS, BP, AW, AR, HO, EJ, KG, AH, DL, LS, MCoC and DW contributed to the planning and undertaking of data collection. All authors contributed to and approved the final manuscript preparation.

Funding This study was funded by a Royal Brisbane and Women’s Hospital Research Project Grant.

Competing interests None declared.

Patient consent for publication Not required.

Ethics approval This project received multisite ethical approval from the Royal Brisbane and Women’s Hospital Medical Research Ethics Committee (HREC/13/QRWB/80). Subsequent site specific approval was then gained from each participating hospital site.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement The dataset from this study is not publicly available due to the data having been collated from multiple hospital health services each with individual data custodians that require further approval for access.

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