1-year follow-up of 920 hip and knee arthroplasty patients after implementing fast-track

Good outcomes in a Norwegian university hospital

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Background — Fast-track has become a well-known concept resulting in improved patient satisfaction and postoperative results. Concerns have been raised about whether increased efficiency could compromise safety, and whether early hospital discharge might result in an increased number of complications. We present 1-year follow-up results after implementing fast-track in a Norwegian university hospital.

Methods — This was a register-based study of 1,069 consecutive fast-track hip and knee arthroplasty patients who were operated on between September 2010 and December 2012. Patients were followed up until 1 year after surgery.

Results — 987 primary and 82 revision hip or knee arthroplasty patients were included. 869 primary and 51 revision hip or knee patients attended 1-year follow-up. Mean patient satisfaction was 9.3 out of a maximum of 10. Mean length of stay was 3.1 days for primary patients. It was 4.2 days in the revision hip patients and 3.9 in the revision knee patients. Revision rates until 1-year follow-up were 2.9% and 3.3% for primary hip and knee patients, and 3.7% and 7.1% for revision hip and knee patients. Function scores and patient-reported outcome scores were improved in all groups.

Interpretation — We found reduced length of stay, a high level of patient satisfaction, and low revision rates, together with improved health-related quality of life and functionality, when we introduced fast-track into an orthopedic department in a Norwegian university hospital.

The health service in Norway has been reorganized in the last decade. The number of available beds and the length of stay (LOS) in somatic hospitals have been reduced. Patients are increasingly being treated as outpatients rather than being admitted to hospital (SSB 2011). Changes in treatment modalities have contributed to this reorganization. Within elective surgery, the "fast-track" principles are increasingly being adopted, although there is still potential for improvement regarding both treatment and clinical results (Rostlund and Kehlet 2007, Kehlet and Soballe 2010). Fast-track originated in Denmark-in gastrointestinal surgery-and has been further developed and documented in joint replacement surgery in hospitals in Denmark over the last decade (Rasmussen et al. 2001, Husted et al. 2010a,d, 2012, Leonhardt et al. 2010). The fast-track concept is an evidence-based multimodality treatment that reduces convalescence time and improves clinical results, including reduction in morbidity and mortality (Kehlet and Wilmore 2008, Schneider et al. 2009). The particularly important elements are: anesthesia, fluid therapy, pain therapy, and early postoperative mobilization (Husted and Holm 2006, Husted et al. 2010a, 2011a, 2012, Khan et al. 2014) as well as preoperative information and supervision (Kehlet 1997, Andersen et al. 2007, 2009, Holm et al. 2010).

It has been said that fast-track may result in increased complication rates and re-admissions (Mauerhan et al. 2003). However, several studies have found that reduced length of stay does not compromise patient safety (Pilot et al. 2006, Mahomed et al. 2008, Schneider et al. 2009) or increase complication rates compared to conventional treatment methods (Husted et al. 2010b). Also, it has been shown that fast-track surgery with early mobilization and short deep-vein thrombosis prophylaxis results in low rates of deep-vein thrombosis and pulmonary embolism (Husted et al. 2010c, Jorgensen et al. 2013).

A reorganization in the orthopedic department at Trondheim University Hospital in 2010 led to an increased number of

knee and hip arthroplasty patients, from 7 to 17 a week (Egeberg et al. 2010). Based on the successful implementation of fast-track in several hospitals in Denmark (Husted et al. 2008, Kehlet and Wilmore 2008), this model was adopted in our department. To be able to continually monitor treatment quality and process data, we established an internal quality register (Bjorgen et al. 2012). We now present the 1-year follow-up results after implementation of this fast-track procedure.

Patients and methods

Patients were recruited between September 1, 2010 and December 1, 2012. From September 2010, all elective patients (ASA classification I, II, and stable III) with a diagnosis of arthritis and scheduled for primary total hip replacement surgery (THA) or total knee replacement surgery (TKA) were enrolled in the fast-track system. From May 1, 2012, all elective primary patients (ASA I–IV) and all elective hip and knee revision patients were also enrolled. Acute events such as fractures or 2-stage revisions due to infection were excluded. At discharge, all fast-track patients were scheduled for 2 follow-up consultations with a physiotherapist (knee patients at 2 months and 1 year; hip patients at 3 months and 1 year). Data concerning LOS prior to the fast-track implementation were obtained from the hospital patient administration system.

The fast-track procedure

The fast-track procedure is based on principles previously described (Kehlet and Wilmore 2008, Handley 2009, Husted et al. 2010d, Kehlet and Soballe 2010, Husted 2012), focusing on standardization and evidence-based care in all parts of the treatment chain. Patients receive oral and written information about the whole procedure, from surgery until follow-up. Patients and relatives are also invited to a multidisciplinary education class arranged at the hospital shortly before admission. An orthopedic surgeon, an anesthetist, a nurse, and a physiotherapist present each part of the treatment chain from admission until discharge. Patients practice walking with crutches and are shown the fast-track unit in which they will be staying during hospitalization. All patients are admitted on the day their surgery is scheduled.

The patients receive optimized pain relief with spinal anesthesia, local infiltration analgesia, and systemic analgesics. The multimodal analgesic regimen is standardized as far as possible; pre-medication consists of paracetamol (1.5–2 g), dexamethasone (16–20 mg), and etoricoxib (90 mg). Benzodiazepines are not used. Operations are done under spinal anesthesia with 2.5–3.0 mL bupivacaine (0.5% plain), preferably at the L2/L3 or alternatively at the L3/L4 vertebral interspace. Propofol infusions are administered for sedation if needed. A standardized program for intraoperative fluid administration is followed consisting of 1–1.5 L Ringer's acetate, 15 mg/kg transexamic acid (max. 1.5 g), and 2 g cephalothin. Following surgery, patients are transferred to the recovery unit and later to a specialized hip and knee arthroplasty unit with a welldefined and experienced program for multimodal rehabilitation. Multimodal, orally administered opioid-sparing analgesia is given to all patients: etoricoxib (90 mg) and paracetamol (1–1.5 g). Oxycodone (5–10 mg) is given if needed (pain score on the numeric rating scale (NRS) > 4).

All patients are mobilized out of bed in the intensive care unit, as soon as the block from the spinal anesthesia disappears. They are encouraged to be as physically active as possible during the hospital stay, to wear their own clothes, and to have their meals in the dining room in the patient ward. Patients participate in group physiotherapy and are also instructed to individually perform specific joint and muscle exercises several times a day according to an exercise guide. Pain experienced at rest and mobilization is registered on an 11-point NRS with a maximum score of 10 where 0 is no pain and 10 is the worst pain imaginable. Before discharge from hospital, specific criteria must be fulfilled: when mobilized, the pain level must be 3 or less on the NRS, the wound must be dry, and the patient must be able to walk on stairs with crutches. Patients are mainly discharged to their homes and given advice about physiotherapy according to conventional rehabilitation procedures.

Registration

2–12 weeks before surgery, patients are screened by nurses and physiotherapists in the outpatient unit. During hospitalization, data are registered by nurses peroperatively and on a daily basis at the fast-track unit. At the follow-up consultations, the physiotherapist registers follow-up data, provides information about current expectations, and gives advice concerning further rehabilitation. If any adverse results are revealed during these consultations, the physiotherapist refers the patient to a surgeon for consultation.

Pain is registered preoperatively, on a daily basis during hospitalization, and at follow-up. An anonymous patient satisfaction form is filled out by the patient throughout the fasttrack course using an 11-point scale (where 0 is worst and 10 is best). LOS is reported as the number of nights hospitalized after the operation. Time to mobilization is defined as the number of hours from the end of surgery to mobilization. Patient-reported outcome scores (PROMS) are measured using (1) the self-administered generic health-related quality of life (HRQoL) questionnaire EQ-5D (Rabin and de Charro 2001), a standardized instrument for use as a measure of health outcome, with a score from 0.50 to 1.00, where 1.00 is the maximum score representing perfect health, for both hip and knee patients, (2) the self-administered disease-specific questionnaires Hip Disability and Osteoarthritis Outcome Score -Physical Function Short Forms (HOOS-PS) for hip patients (Davis et al. 2008), and (3) Knee Injury and Osteoarthritis Outcome Score - Physical Function Short Forms (KOOS-PS) for knee patients (Perruccio et al. 2008). The 2 latter forms have a score from 0 to 100 with 0 being the optimal score, representing no difficulty in performing specific tasks. These are designed for objective assessment of physical function.

All patients are examined by a physiotherapist and physical function is measured by the disease-specific Harris hip score (HHS) and the American Knee Society score (KSS). Maximum score for the HHS is 100 points, with a score of > 70 meaning poor, 70–79 fair, 80–89 good, and 90–100 excellent. The

maximum KSS values (knee score and functional score) are 100 points each, with a score of > 60 meaning poor, 60–69 fair, 70–79 good, and 80–100 excellent. All questionnaires are filled out preoperatively and twice postoperatively, knee patients after 8 weeks and hip patients aafter 12 weeks and after 1 year. 1 year after surgery, the patients are asked 2 specific questions: (1) "How does the leg that was operated on work today compared to before surgery?", and (2) "Based on your experience to date, would you go through the surgery again?".

Ethics

Before registration, all patients are informed about the registry and asked to sign consent forms allowing their data to be used for scientific purposes. According to our regional ethics committee, the present study was a quality assurance study that did not require any approval in order to be performed or published.

Statistics

The data distribution was evaluated by visual inspection of histograms. Normally distributed data are presented as mean with standard deviation (SD) or range. Since only 13 of the 920 patients (1.4%) had bilateral procedures (bilateral observations), the within-individual correlations were not taken into account in the statistical analyses. Data concerning complications are presented as relative risk with 95% confidence interval (CI) (Hagen 1998). The analyses were performed using SPSS version 21.0 and Microsoft Excel.

Results

Before the implementation of fast-track, 7 hip or knee arthroplasty patients were operated every week at Trondheim University Hospital and 8 were operated on in a satellite clinic. Mean LOS was 8.1 (5.3) days for primary hip patients and 8.1 (5.1) days for primary knee patients. After fast-track implementation, all hip and knee arthroplasty surgeries were performed at the university hospital, with a total number of 17 surgeries a week. 1,069 fast-track patients were included in

Table 1. Patient characteristics in the different groups (primary total hip arthroplasty and primary total knee arthroplasty; revision total hip arthroplasty and revision total knee arthroplasty)

| | Primary s | urgery | Revision surgery | | | |
|----------------------------------|------------|------------|------------------|------------|--|--|
| | THA | TKA | THA TKA | | | |
| Mean age (range), years | 65 (17–90) | 66 (36–89) | 68 (39–87) | 67 (46–82) | | |
| No. of men/women | 212/407 | 137/232 | 21/33 | 11/16 | | |
| Mean BMI (SD), kg/m ² | 27 (4) | 29 (5) | 27 (5) | 29 (5) | | |
| ASA I, % | 17 | 13 | 6 | 11 | | |
| ASA II, % | 68 | 67 | 69 | 59 | | |
| ASA III, % | 15 | 18 | 26 | 30 | | |

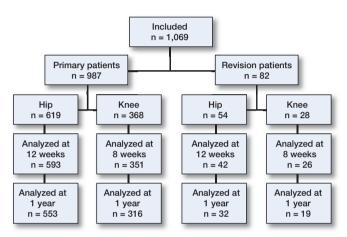


Figure 1. Participants from inclusion until 1-year follow-up.

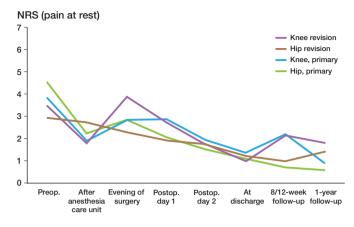
the present study: 619 primary THA patients, 368 primary TKA patients, 54 revision THA patients, and 28 revision TKA patients (Table 1).

For THA, the number of 12-week follow-ups was 593 (96%) for primary patients and 42 (78%) for revision patients. For TKA, the number of 8-week follow-ups was 351 (95%) for primary patients and 26 (93%) for revision patients. At the 1-year follow-up, 553 (89%) primary, and 32 (59%) revision hip patients were analyzed, and 316 (86%) primary and 19 (68%) revision knee patients were analyzed. A flow chart of the participants is given in Figure 1.

Mean patient satisfaction with the LOS for all patients was 8.9 (1.7), and with the fast-track course it was 9.3 (1.2). Pain scores at rest and during mobilization from before operation until the 1-year follow-up are presented in Figure 2. PROMS, physical function scores, and complications for primary and revision patients are given in Tables 2, 3, and 4, respectively.

Primary surgery

Of the THA patients who underwent primary surgery, 94% were mobilized in the recovery unit and the mean time from surgery to mobilization was 3.5 (1.6) h. Mean LOS was 3.1 (0.8) days. 82% of the patients were discharged directly to their homes. Total re-admission rate within one year was 5.7%



NRS (pain during mobilization)

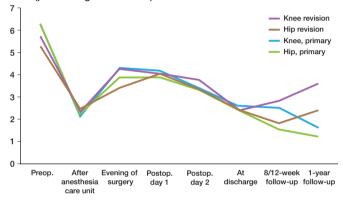


Figure 2. Pain at rest (upper panel) and mobilization (lower panel) at 8 time points, from preoperatively until 1 year postoperatively. Lines represent mean pain score for each patient group.

Table 2. Patient-reported outcome scores (PROMS) in the 4 different groups: EQ–5D, HOOS–PS, KOOS–PS, and the physical function scores HHS and KSS. For abbreviations, see text. All scores were obtained preoperatively, 8 and 12 weeks postoperatively, and 1 year postoperatively. Data are mean (SD)

| | Primary s | urgery | Revision surgery | | | |
|----------------------------|-------------|--------------------|------------------|--------------------|--|--|
| | THA | ТЌА | THA | TKA | | |
| Preoperatively | | | | | | |
| EQ-5D index score | 0.42 (0.30) | 0.49 (0.29) | 0.42 (0.32) | 0.38 (0.36) | | |
| HOOS-PS/KOOS-PS | 45 (16) | 47 (13) | 49 (18) | 49 (11) | | |
| HHS | 53 (14) | () | 52 (24) | | | |
| KSS knee score | | 41 (14) | - () | 49 (19) | | |
| KSS function score | | 56 (22) | | 43 (23) | | |
| 8/12 weeks postoperatively | | · · / | | . , | | |
| EQ-5D index score | 0.76 (0.21) | 0.68 (0.24) | 0.62 (0.32) | 0.53 (0.32) | | |
| HOOS-PS/KOOS-PS | 22 (14) | 37 (11) | 33 (16) | 41 (12) | | |
| HHS | 82 (13) | | 69 (20) | | | |
| KSS knee score | | 61 (19) | | 64 (15) | | |
| KSS function score | | 59 (25) | | 45 (28) | | |
| 1 year postoperatively | 0.70 (0.04) | 0.70 (0.000) | 0.07 (0.07) | | | |
| EQ-5D index score | 0.79 (0.24) | 0.76 (0.230) | 0.67 (0.27) | 0.62 (0.25) | | |
| HOOS-PS/KOOS-PS | 17 (15) | 30 (15) | 27 (15) | 34 (15) | | |
| HHS KSS knee score | 89 (14) | 75 (10) | 77 (20) | 60 (10) | | |
| KSS function score | | 75 (19) 78 (24) | | 62 (13) 65 (25) | | |
| NGG TUTICUOIT SCOLE | | 78 (24) | | 65 (25) | | |

(4.3-7.0). The revision rate was 2.9% (2.0-3.9), of which 1.6% (0.9-2.3) was caused by infections and 1.0% (0.4-1.5) by dislocations. At 1-year follow-up, 83% reported improved functionality in the operated limb and 85% reported that they would have been willing to have the surgery all over again.

For TKA patients who underwent primary surgery, 96% were mobilized in the recovery unit and the mean time from surgery to mobilization was 3.2 (1.4) h. Mean LOS was 3.1 (0.8) days. 82% of the patients were discharged directly to their homes. Total re-admission rate within 1 year was 10.1% (8.4–11.8). The revision rate was 3.3% (2.3–4.3), of which 1.4% (0.7–2.0) was caused by infections and 1.4% (0.7–2.0) by mechanical causes. At 1-year follow-up, 72% reported improved functionality in the operated limb and 73% reported that they would have been willing to have the surgery all over again.

Revision surgery

For THA patients who underwent revision surgery, 91% were mobilized in the recovery unit and the mean time from surgery to mobilization was 4.6 (2.2) h. Mean LOS was 4.2 (1.6) days and 44% of the patients were discharged directly to their homes. Total re-admission rate within 1 year was 5.6% (4.3– 6.9), all of which was caused by infections. At 1-year followup, 39% reported improved functionality in the operated limb and 43% reported that they would have been willing to have the surgery all over again.

For TKA patients who underwent revision surgery, 89% were mobilized in the recovery unit and the mean time from surgery to mobilization was 4.9 (1.9) h. Mean LOS was 3.9

(2.2) days and 67% of the patients were discharged directly to their homes. Total re-admission rate within 1 year was 7.1% (5.7–8.6), of which 3.6% (2.5–4.6) was caused by infection and 3.6% (2.5–4.6) was due to mechanical causes. 1 year postoperatively, 48% reported improved functionality in the operated limb and 52% reported that they would have been willing to have the surgery all over again.

Discussion

After implementation of the fast-track course, the number of weekly hip and knee arthroplasty surgeries was increased from 7 to 17. Patient satisfaction was high in all parts of the treatment chain, with a mean score of 9.3 (1.2) out of a maximum of 10. 85% of hip patients and 73% of knee patients were satisfied with the results 1 year

| | Primary THA | | Primary TKA | | | |
|--------------------------------|-------------|-----------------|-------------|-----|-----------------|---------|
| | n | RR ^a | 95% CI | n | RR ^a | 95% CI |
| Total | 619 | | | 368 | | |
| Re-admissions | 35 | 5.7 | 4.3-7.0 | 37 | 10.0 | 8.4–12 |
| Revisions | 18 | 2.9 | 2.0-3.9 | 12 | 3.3 | 2.3-4.3 |
| Infection | 10 | 1.6 | 0.9–2.3 | 5 | 1.4 | 0.7-2.0 |
| Dislocation | 6 | 1.0 | 0.4–1.5 | - | _ | _ |
| Impaired mobility | _ | _ | - | 1 | 0.3 | 0.0-0.6 |
| Fracture | 1 | 0.2 | -0.1-0.4 | _ | _ | _ |
| Mechanical causes | 1 | 0.2 | -0.1-0.4 | 5 | 1.4 | 0.7–2.0 |
| Reoperations | 7 | 1.1 | 0.5-1.7 | 12 | 3.3 | 2.3-4.3 |
| Dislocation | 5 | 0.8 | 0.3–1.3 | 3 | 0.8 | 0.3–1.3 |
| Gluteal insufficiency | 1 | 0.2 | -0.1-0.4 | _ | _ | _ |
| Quadriceps rupture | _ | _ | _ | 1 | 0.3 | 0.0-0.6 |
| Impaired mobility | _ | _ | _ | 6 | 1.6 | 0.9-2.3 |
| Fracture | 1 | 0.2 | -0.1-0.4 | 1 | 0.3 | 0.0-0.6 |
| Other causes | _ | _ | _ | 1 | 0.3 | 0.0-0.6 |
| Other causes | 10 | 1.6 | 0.9-2.6 | 13 | 3.5 | 2.5-4.6 |
| DVT | 1 | 0.2 | -0.1-0.4 | _ | _ | _ |
| Hematoma | 1 | 0.2 | -0.1-0.4 | _ | _ | _ |
| Stroke | 1 | 0.2 | -0.1-0.4 | _ | _ | _ |
| lleus | 1 | 0.2 | -0.1-0.4 | _ | _ | _ |
| Superficial infection | _ | _ | _ | 1 | 0.3 | 0.0-0.6 |
| Peroneus palsy | _ | _ | _ | 1 | 0.3 | 0.0-0.6 |
| Quadriceps rupture | _ | _ | _ | 1 | 0.3 | 0.0-0.6 |
| Urinary infection | _ | _ | _ | 1 | 0.3 | 0.0-0.6 |
| Gluteal insufficiency | 2 | 0.3 | 0.0-0.6 | _ | _ | _ |
| Other causes | 4 | 0.6 | 0.2-1.1 | 9 | 2.4 | 1.6-3.3 |
| Leg length difference (> 2 cm) | 3 | 0.5 | 0.1-0.9 | 1 | 0.3 | 0.0-0.6 |
| Superficial infection | 13 | 2.1 | 1.3–2.9 | 15 | 4.1 | 3.0-5.2 |
| Urinary infection | 27 | 4.4 | 3.2-5.5 | 22 | 6.0 | 4.6-7.3 |
| Pneumonia | 2 | 0.3 | 0.0-0.6 | | _ | _ |
| Impaired mobility | _ | - | _ | 12 | 3.3 | 2.3-4.3 |
| Peroneus palsy | _ | _ | _ | 1 | 0.3 | 0.0-0.6 |
| Gluteal insufficiency | 2 | 0.3 | 0.0-0.6 | _ | - | _ |
| Other causes | 15 | 2.4 | 1.6-3.3 | 11 | 3.0 | 2.0-4.0 |
| | | | | | 0.0 | |

Table 3. Complications within 1 year of primary total hip arthroplasty or primary total knee arthroplasty

^a Relative risk, %.

Table 4. Complications within 1 year of revision surgery for hip and knee patients

| | Revision THA | | n THA | Revision TKA | | |
|----------------------------|--------------|------|---------|---------------------|------|---------|
| | n | RR ª | 95% CI | n | RR ª | 95% CI |
| Total | 54 | | | 28 | | |
| Re-admissions Revisions | 3 | 5.6 | 4.3–6.9 | 2 | 7.1 | 5.7–8.6 |
| Infection | 2 | 3.7 | 2.6-4.8 | 1 | 3.6 | 2.5-4.6 |
| Mechanical causes | - | - | - | 1 | 3.6 | 2.5-4.6 |
| Other causes | - | - | - | - | _ | - |
| Sub dislocation | 1 | 1.9 | 1.1–2.6 | - | - | - |
| Superficial infection | 1 | 1.9 | 1.1–2.6 | - | - | - |
| Urinary infection | 2 | 3.7 | 2.6-4.8 | 1 | 3.6 | 2.5-4.6 |
| Pneumonia | - | - | - | 1 | 3.6 | 2.5–4.6 |

after the operation. In addition, LOS was reduced by approximately 5 days for both hip patients and knee patients. The satisfaction with the LOS had a mean score of 8.9 (1.7) out of a maximum of 10. These findings are in line with other

publications concerning fast-track arthroplasty (Husted et al. 2008, 2010a). Hospital logistics and clinical features are crucial for the LOS. and reduced LOS reduces costs without compromising treatment quality (Husted et al. 2008, 2010d, 2012). The numbers of re-admissions and revisions in the present study were lower than those previously reported (Husted et al. 2008, 2010b). These results demonstrate that even though the treatment is more effective, as indicated by the increased number of patients operated annually and with a reduced LOS, it does not compromise patient satisfaction or treatment quality. Fast-track also provided good results in non-septic revision surgery, as has been reported previously by others (Schneider et al. 2009, Husted et al. 2011b, Jorgensen et al. 2013).

Measurement of pain with an NRS is a practical method to use, since it is easy to understand and the patient does not need clear vision or paper and pen to describe the degree of pain-in contrast to the visual analog scale (VAS). An NRS of 3 or less corresponds to "mild pain" on the VAS, whereas 4-6 corresponds to "moderate pain" and 7-10 to "severe pain" (Breivik et al. 2008). Pain, dizziness, and general weakness are main causes of prolonged postoperative hospitalization. Pain is also a limiting factor for early postoperative mobilization and physical activity (Husted et al. 2011a). In general, the patients in our study had only mild pain directly after or during the evening following surgery. We found that the maximum pain experienced was during rest in the evening after surgery, both for primary hip and knee patients and for revision hip and knee patients (2.9 (2.0), 2.9 (2.0), 2.3 (1.8), and 3.9 (2.6), respectively). From day 2 postoperatively to patient discharge, the pain both at rest and during mobilization was reduced and sustained at the "mild pain" level. These results suggest that pain was no limitation regarding early discharge.

For optimal rehabilitation and assured function, patients need continuous analgesic treatment after discharge (Andersen et al. 2009, Husted et al. 2011a). Holm et al. (2010) found that pain has a limited influence on functional recovery after the first postoperative day after

TKA, thereby allowing early physiotherapy. They found that 90% of the patients were mobilized after the first postoperative day, which agrees with our findings. We found that over 90% of the primary patients and about 90% of the revision patients were mobilized at the recovery unit, and the mean time from surgery until mobilization was about 3.5 h and 4.5 h, respectively. Patients who were not mobilized were either immobilized due to perioperative complications, illness, or inability to cooperate.

Early mobilization is important to reduce the risk of thrombosis and to initiate rapid recovery (Husted et al. 2008, 2010b,c). The numbers of deep-vein thrombosis and pulmonary embolism could be underreported in our study. These diagnoses had to be verified by ultrasonography or CT before being reported to the registry.

All the patients had to fulfill the fast-track discharge criteria before leaving hospital, either to their homes or to the rehabilitation institution. Before the fast-track implementation, there were no standardized discharge criteria, and the patients usually went to a rehabilitation institution. Approximately 80% of the primary patients in our study were discharged directly to their homes, which is a lower percentage than can be found in publications from Denmark (Husted et al. 2008). In Norway, it has been a tradition to discharge patients to rehabilitation institutions depending, for example, on the patient's housing conditions, care facilities, comorbidities, and distance from hospital. Now that fast-track has been implemented, this trend is about to shift. Only 44% of the revision patients were discharged to their homes due to their reduced functionality postoperatively-the result of greater surgical trauma, as full synovectomi and intramedular reaming for femoral and tibial components are used in the revision surgeries. At postoperative follow-ups for all groups at week 8 or 12, and at 1 year, the mean pain score was relatively low, as expected, with values of less than 3 on the NRS. These findings are in line with the results of other studies (Rostlund and Kehlet 2007, Holm et al. 2010), demonstrating low levels of postoperative pain after fast-track joint replacement surgery.

Even though the main purpose of THA and TKA surgery is pain relief, regaining HRQoL and functionality is considered to be the ultimate goal after joint replacement (Woolf and Akesson 2001). Thus, to evaluate the treatment outcome, it is important to measure outcomes from the patient's perspective. In a fast-track treatment set-up without any formal intensive rehabilitation after discharge, it was found that THA and TKA patients at 1-year follow-up had regained health similar to that of an age- and sex-matched population in Denmark when measured by HRQoL (EQ-5D) (Larsen et al. 2010, 2012). We found similar improvements (using EQ-5D scores) one year after primary THA and TKA to those reported by Larsen et al. However, the patients in our study did not reach the same HRQoL levels 1 year postoperatively, which may have been due to the different demographics of the patient cohorts. Patients included in the hip study (Larsen et al. 2010) had a diagnosis of unilateral primary arthritis, and in the knee study (Larsen et al. 2012) patients with unicompartmental knee arthroplasty were included. In the present study, patients with bilateral hip arthroplasty were included and patients with unicompartmental knee arthroplasty were excluded. Thus, our patients may have been affected by the disease to a greater extent than the patients in the Larsen study, therefore resulting in both higher preoperative and 1-year HRQoL scores. EQ–5D results from the Swedish Arthroplasty Register showed that THA patients had mean scores of 0.42 preoperatively and 0.78 after 1 year (Rolfson et al. 2011), which is similar to what we found. A cohort study from Karolinska University Hospital in Sweden (Jansson and Granath 2011) also found similar EQ–5D results in THA patients pre- and 1 year postoperatively (0.49–0.80, and TKA patients pre- and 1 year postoperatively (0.51–0.73).

The THA patients in the study by Larsen et al. (2010) did not regain the same level of health as the age- and sexmatched normal population, which was based on the diseasespecific HHS outcome one year postoperatively (88 vs. 94). However, the results were similar to our findings (HHS = 89) 1 year postoperatively. Any HHS score within these values is close to "excellent" (i.e. close to > 90) on the HHS grading scale, which indicates that even though THA patients have good functionality at 1 year postoperatively in disease-specific terms, they do not reach the level of the normal population. In another study (Medalla et al. 2009), the postoperative KSS outcomes, knee score (81) and physical function score (71), were similar to the postoperative scores that we found (75 and 78, respectively). However, patients in the Medalla study gave their postoperative scores 2 years after surgery, as compared to 1 year in the present study. Also, the patients in the Medalla study generally had a higher knee score and a lower physical function score than in our study. This might be explained by the different postoperative testing times, since knee function is directly related to the surgery and physical function is related to the physical fitness of the patient, which is not directly influenced by the surgery. Scores between 70 and 80 correspond to a "good" knee and functionality score on the KSS grading scale. The knee and physical function scores in the present study were both > 75, which is close to "excellent" (> 80), indicating good functionality one year postoperatively.

In a study from Toronto, Canada (Davis et al. 2009), based on 201 THA and 248 TKA patients, the improvements in HOOS–PS (55%) and KOOS–PS (33%) from preoperatively to 6 months after surgery were similar to what we found. This similarity was apparent in HOOS–PS from before surgery to 3 and 12 months after surgery (51% and 62%, respectively), and in KOOS–PS from before surgery to 2 and 12 months after surgery (21% and 35%), where the patients in both studies reported less difficulty in performing daily activities postoperatively. In light of our findings and those previously reported by others, fast-track is a treatment that gives good postoperative results, both in generic and disease-specific terms.

The overall loss of participants at follow-up in our study 1 year postoperatively was 14%, which is less than that reported by others (26%) (Larsen et al. 2012). A higher proportion of revision patients than primary patients were lost to follow-up

at 1 year. Local hospitals often refer patients scheduled for revision surgery to the university hospital. The follow-ups are often done at the local hospital, which could explain the higher loss of patients observed in this group. Patients eligible for follow-up had different reasons for not turning up. In some cases the secretary had not scheduled an appointment, some patients had forgotten the appointment, some patients did not want the consultation, and patients with severe complications at the first follow-up were excluded from the second followup. However, all patients with complications resulting in revision were followed up, as they were re-admitted to the same university hospital.

In summary, we found improved efficiency, high patient satisfaction, and low revision rates together with improved health-related quality of life and functionality with the fasttrack course implemented in a Norwegian university hospital, indicating that fast-track is a favorable and feasible method of treatment for both primary and revision THA and TKA patients.

All the authors contributed to the literature search, study design, data collection, data analysis, and interpretation. All the authors contributed substantially to drafting of the article and they also revised it for important intellectual content.

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