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Adverse events in total knee arthroplasty: Results of a physician independent survey in 260 patients

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Abstract

Purpose: Identification of all common and potentially avoidable adverse events is crucial to further improve the quality of medical care. The intention of the current study was to evaluate a standardized physician independent survey format on adverse events in total knee arthroplasty. The protocol for reporting adverse drug events following the International Conference of Harmonisation of technical requirements for registration of pharmaceuticals for human use (ICH) was adopted for adverse events occurring during surgical interventions.

Material and methods: Data of a prospective sequential cohort trial introducing a clinical pathway for total knee arthroplasty was analysed. Reporting of adverse events was done by a physician independent study nurse using the modified ICH-Good Clinical Practice (GCP) format (Structure and Content of Clinical study reports - E3) in 260 patients. The adverse events were graded to their severity and their potential relation to surgical treatment.

Results: A total of 55 patients (21%) suffered from an adverse event and 16 (6%) from a serious adverse event. In 38 patients' one adverse event occurred, 12 patients showed 2 adverse events and 5 patients suffered from a combination of an adverse and a serious adverse event. A serious adverse event alone occurred in 11 patients. The incidence of adverse events (Fisher $p = 0.448$) and serious adverse ($p = 0.126$) events showed no significant difference between the two cohorts. The most common adverse events were deep vein thrombosis (8% and 5%) followed by wound healing problems (1% and 0%) and haematoma (1% and 3%). A wide range of non surgical adverse events were recorded with low incidence levels.

Conclusion: The use of the modified ICH-GCP format supports standardization of adverse event reporting. Routine assessment of adverse events by a study nurse revealed higher incidence rates of adverse events in total knee arthroplasty. We recommend the implementation of trained paramedical staff for the documentation of adverse events in routine clinical care.

Introduction

Total knee replacement provides safe and effective treatment for patients suffering from end stage osteoarthritis [1]. Improvement in medical care offers total knee replacement to an increasing number of older patients even with severe comorbidities[2]. However, for the treatment of older and sicker patients an increased rate of medical complications has to be assumed,[3] but nevertheless the reported rate of adverse events is low, however, especially for the data reported to the national public

external quality assuring system in Germany[4]. The reporting of adverse events in former papers on total joint replacement mainly focuses on major surgical adverse events like wound infection, fracture or loosening of the implant[5-10]. Therefore medical adverse events are likely to be underreported in these investigations [11]. In an evidence report on total knee arthroplasty [12] a more complete list of adverse medical as well as surgical events is published. Alfonso et al. [13] reviewed the non surgical complications after total hip and total knee arthroplasty and also used a more detailed listing of adverse events.

Beside this tendency in the orthopaedic literature there is an ongoing debate about the reliability of

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measurement and monitoring of adverse events[14]. Identification of all common and potentially avoidable adverse events is crucial to further improve the quality of medical care. Routine reporting by physicians identified fewer events than medical record review[15]. Therefore a different workflow to monitor adverse events in surgical patients may be advisable. A possible entry into an increased accuracy in monitoring may be the introduction of a trained study nurse or medical documentation assistant into the clinical routine surveillance system to thoroughly monitor adverse events in surgical patients undergoing total knee replacement; the well-established report instruction (Structure and Content of clinical study reports - E3) of the Good Clinical Practice (GCP) Guideline from the International Conference on Harmonisation of technical requirements for registration of pharmaceuticals for human use (ICH GCP[16] and ICH E3) may provide an efficient and standardized rationale for this additional stage in quality assessment.

Patients and Methods

The intention of the current study was to propose and illustrate a standardized documentation and reporting assessment tool for (serious) adverse event patterns in Orthopaedic Surgery; furthermore the putative effect of the implementation of a physician independent survey on adverse events in total knee arthroplasty had to be evaluated. For that purpose data of a prospective sequential cohort trial introducing a clinical pathway for total knee arthroplasty was re-analysed. Two sequential cohorts were recruited with the aim of simultaneously estimating the process costs from the health care providers' perspective and the patient-related benefit of TKA. A total of 260 patients with the clinical indication for TKA were enrolled consecutively. Data on the present state was collected with a process documentation of the clinical course within a first cohort of 132 patients. The recorded data was used to derive and then implement an interdisciplinary clinical pathway on TKA; after a three months period of confirming this pathway proposal within clinical routine environments, a second independent cohort of 128 patients undergoing TKA was recruited and documented at the same hospital. Further sociodemographic details of both study cohorts are given in table 1. Furthermore all study patients were invited for a three months recall. The underlying prospective cohort design was ratified by the local Independent Ethics Committee by June 25th 2005. All patients gave informed consent to take part in this trial.

Surgical procedure

The study population consisted of patients with primary or secondary knee osteoarthritis grade 3 and 4 according to Kellgren and Lawrence[17]. Surgery was performed in

nearly all patients under regional anaesthesia and a tourniquet was applied after admission of preoperative antibiotic prophylaxis (third generation cephalosporine). Through a straight skin incision and a medial parapatellar approach a cemented Natural Knee II (Fa. Zimmer, Germany) total knee arthroplasty without resurfacing of the patella was implanted. All patients undergoing total knee replacement were graded to be at high risk for deep venous thrombosis. All patients received Arixtra 1 × 1 s. c. for 6 weeks. Postoperatively all patients were allowed for immediate full weight bearing. After discharge from the hospital the patients were transferred to an inpatient rehabilitation unit for additional three weeks.

Monitoring of adverse events and data collection

All data collection preoperatively and at follow-up was done by a trained medical documentation assistant. Functional ability and health-related quality of life were assessed by means of appropriate instruments (WOMAC[18], EQ-5D[19] and Knee Society Knee Score)[20]. For monitoring of adverse and serious adverse events the definitions of the ICH GCP were adapted for surgical interventions and implemented as described below. The original wording is given in square brackets followed by the new text in italic letters.

Definition of adverse events (AE): any unexpected medical occurrence in a patient or clinical investigation subject [administered a pharmaceutical product] *undergoing surgical treatment*, which does not necessarily have a causal relationship with this treatment. An adverse event (AE) can therefore be any unfavourable and unintended sign (including an abnormal laboratory finding), symptom, or disease temporally associated with [the use of a medicinal (investigational) product] *surgical treatment*, whether or not related to [the medicinal (investigational) product] *surgical treatment*.

definition of serious adverse events (SAE): any unexpected medical occurrence [at any dose] *in the operative period*, which resulted in death, went life-threatening, required inpatient hospitalization or prolongation of existing hospitalization, or resulted in persistent or significant disability/incapacity.

Adverse events and serious adverse events were recorded at time of clinical appearance or at time of first available information to the Orthopaedic surgery department. In any case of a suspected deep vein thrombosis the patients were admitted to the university vascular centre and further sonographic diagnostics were carried out. There is no general screening for DVT in patients undergoing TKA. A medical documentation assistant interviewed all patients at follow up and actively screened any adverse event possible after knee arthroplasty. In order to grade the adverse events

Table 1 Distribution characteristics for sociodemographic cofactors assessed in 132 patients, who underwent total knee arthroplasty (TKA) before implementation of a critical pathway on TKA, versus 128 patients, who underwent TKA after path implementation

	Before pathway implementation (n = 132)	After pathway implementation (n = 128)
<i>Age: median and range [years]</i>	68 (43 - 88) years	70 (46 - 85) years
<i>Females</i>	64%	56%
<i>living alone</i>	32%	28%
<i>under employment</i>	8%	9%
<i>graduate</i>	20%	17%

according to their severity and their potential relation to surgical treatment all (S)AE reports by the medical documentation assistant were reviewed by the coordinating clinical investigator. Almost all patients suffering from an adverse event were consulted by a specialist. The (S) AE events were graded “surely, possibly, probably, non related”. Only in the case of a undoubtedly pre-existing disease like a Hyperhomocystinemia the (S)AE was graded “non related”. In all other cases a possible relation to surgical treatment was considered. At the time of the three months follow-up investigation the impact of the adverse event on mid-term outcome after total knee replacement was estimated. No structured interviews were done afterwards.

Statistical analysis

The descriptive analysis of this embedded quality investigation was based on two-sided Fisher tests at a local 5% significance level to compare the cumulative incidences of adverse events and serious adverse events between the two sequential cohorts. The observed (serious) adverse events were listed according to the ICH report form.

To further investigate possible factors contributing to the observed (serious) adverse event patterns a severity score on the individual adverse event patterns was constituted: The appearance of two adverse events or a combination of an adverse event together with a serious adverse event was hypothesized to be a more severe impairment on individual health than a single adverse events or no adverse event at all. The severity score counted one point for every adverse event and two points for serious adverse events. Description of this score was based on relative frequencies. Furthermore a multiple logistic regression model was fitted for this score by means of a forward selection model and Likelihood Ratio tests at a local 5% significance level: A total score of 2 points or more was considered “severe” and applied as a surrogate endpoint of the overall (S)AE patterns; age at time of surgery, the patients’ individual comorbidity and complexity level (pccl, derived from the DRG calculation), gender, educational level, WOMAC index before surgery

and cut suture time were introduced into the model as possible explaining factors, the patients’ respective cohort - before versus after pathway implementation - was furthermore included.

All numerical evaluations were based on the software SPSS® (release 17.0 for Windows®).

Results

A total of 55 patients (21%) suffered from an adverse event and 16 (6%) from a serious adverse event. The improvement in terms of patient related outcome measures for WOMAC, EQ-5 D and Knee Society Knee and Function Score is given in table 2. In 38 patients’ one adverse event occurred, 12 patients showed 2 adverse events and 5 patients suffered from a combination of an adverse and a serious adverse event. A serious adverse event alone occurred in 11 patients. In the entire cohort at total of 68 adverse events and 16 serious adverse events were recorded, 30 + 5 of which reported from the first cohort (23% + 4%) and 37+11 from the second cohort (29% + 11%) after clinical pathway implementation. The incidence of adverse events (Fisher p = 0.448) and serious adverse (p = 0.126) events showed no significant difference between the two cohorts. The most common adverse events were deep vein thrombosis with an occurrence rate of 8% and 5% for the lower leg in the respective cohorts. Further surgical adverse events like wound healing problems (1% and 0%) and haematoma (1% and 3%) were observed less frequently. Falls (2%) were only reported for the second cohort. Further adverse events were reported at a lower level of incidence (Table 3).

In cohort 1 (before pathway implementation), a total of 10 patients (8%) showed a severity adverse event score of 2 or more, in cohort 2 (after pathway implementation) a total of 17 patients (13%). The logistic regression model could only identify the individual pccl (LR p < 0.001) and the a patient’s cohort designation (LR p = 0.036) as significantly associated with this endpoint; neither age at time of surgery (LR p = 0.062) nor gender (LR p = 0.106) nor educational level (LR p = 0.096) nor cut/suture time (LR p = 0.366) nor

Table 2 Medians and quartiles for the total WOMAC osteoarthritis index, EQ-5 D [%, 100% = optimum rating] and Knee Society Knee and Function Score before and three months after total knee arthroplasty (TKA)

	Before pathway implementation (n = 132)	After pathway implementation (n = 128)
EQ-5D		
<i>preoperative</i> [%]	45% (35 - 50%)	50% (31 - 50%)
<i>postoperative</i> [%]	75% (60 - 85%)	75% (60 - 80%)
WOMAC		
<i>preoperative</i> [%]	41% (32 - 48%)	44% (36 - 52%)
<i>postoperative</i> [%]	83% (68 - 91%)	82% (70 - 91%)
Knee Society		
Knee Score		
<i>preoperative</i> [pt]	43 (29-54)	46 (37-56)
<i>postoperative</i> [pt]	87 (69-94)	85 (65-92)
Knee Society Function Score		
<i>preoperative</i> [pt]	50 (40-60)	55 (45-60)
<i>postoperative</i> [pt]	60 (50-75)	65 (55-76)

WOMAC (LR $p = 0.344$) before surgery were found significantly associated with the occurrence of a severity score of 2 points and more.

Discussion

Improvement in medical care extends the life expectancy of our patients and secondary increases the need for total joint replacement[21]. For surgical procedures with a high volume the introduction of clinical pathways was promoted[22]. The quality of medical care can be improved through the use of clinical pathways embedded within a quality management system by continuously measuring and evaluating process and outcome indicators[23]. Especially for elder patients this systematic and structured improvement process could be beneficial and illustrates again the relevance of the PDCA-cycle of Walter Deming. Therefore a vital part of continuous quality assurance is the monitoring of adverse events and to evaluate whether these were avoidable[24]. University hospitals are more likely to treat patients at higher age and with more complex comorbidities. Therefore a higher rate for adverse events has to be assumed in these institutions.

The rates for adverse events (21%) and serious adverse events (6%) in our patients appear relatively high when compared to the official data of the national external quality assurance system in Germany[4]. In this nation wide database for example the average incidence of general medical complications was reported to be 2% with a reference limit of 5%. Data published by Parvizi et al. [3] with 21% minor complications and 5% major complication, however, match better to our observations. Similar

results were found by Frosch et al.[25]. Alfonso [13] provided an extensive review of the non orthopaedic literature and confirmed a broad range of non surgical adverse events in patients undergoing total joint replacement. The importance to use accepted definitions was highlighted by Bruce et al.[14]. Especially in the context of suspected wound infection he found only little evidence for systemic measurement and monitoring of surgical wound. The strict definition for persisting wound drainage by Parvizi [3] as any secretion from the wound 48 hours post surgery is easy to use. In case of postoperative haematoma or suspected wound infection active treatment with revision of the wound is recommended. Timing of the intervention is important for further prognosis [26]. If in doubt a revision should be undertaken. The clear and comprehensible definition makes it easier for the patient to understand the situation and to agree to further surgery. We carried out 4 revisions for suspected early infection and one for haematoma.

Falls of patients occurring after surgical procedures performed under regional anaesthesia are reported in the literature[27]. We observed these only in the second cohort of our patients. During pathway implementation no changes concerning the use of regional anaesthesia were made. In contrast the patients of the second cohort were offered to attend further information about the disease, the operation and the pain management with regional anaesthesia thereafter. The increased rate of documented falls may be attributable to closer monitoring of our patients. Higher age as well as multiple comorbidities are a documented risk factors for the occurrence of adverse events[28,29]. We were able to

Table 3 Adverse event and serious adverse event counts, percentages given in brackets, for 132 patients, who underwent total knee arthroplasty (TKA) before implementation of a critical pathway on TKA, versus 128 patients, who underwent TKA after path implementation

adverse event	Before pathway implementation	After pathway implementation	Total
	n = 132	n = 128	n = 260
Generalised seizure due to local anaesthetic intoxication	0 (0)	1 (1)	1 (0)
DVT (lower leg)	10 (8)	6 (5)	16 (6)
DVT (knee)	4 (3)	2 (2)	6 (2)
DVT (thigh)	0 (0)	1 (1)	1 (0)
Haematoma	1 (1)	3 (2)	4 (2)
Wound healing problems	1 (1)	0 (0)	1 (0)
Stiff knee	0 (0)	1 (1)	1 (0)
Gastrointestinal bleeding	0 (0)	3 (2)	3 (1)
Dekubital ulcer	1 (1)	2 (2)	3 (1)
Postoperative cognitive disorder	1 (1)	0 (0)	1 (0)
Speech disturbance and Hyperkinesia in Parkinson disease	2 (2)	1 (1)	3 (1)
Postoperative hypertonia	1 (1)	3 (2)	4 (2)
Cardiac blocks	2 (2)	2 (2)	4 (2)
Diarrhoea	1 (1)	1 (1)	2 (1)
Obstipation	0 (0)	1 (1)	1 (0)
Erysipel	1 (1)	0 (0)	1 (0)
Exanthema	1 (1)	3 (2)	4 (2)
Otolaryngology consultation*	2 (2)	0 (0)	2 (1)
Dentist consultation*	1 (1)	0 (0)	1 (0)
Urinary tract infection	0 (0)	2 (2)	2 (1)
Urologic consultation*	1 (1)	2 (2)	3 (1)
Fall	0 (0)	2 (2)	2 (1)
Posterior collum lesion (myelon)*	0 (0)	1 (1)	1 (0)
Hyperhomocystinemia*	0 (0)	1 (1)	1 (0)
Total AE	30 (23)	38 (29)	68 (26)
serious adverse event			
Death during study period, unrelated	1 (1)	1 (1)	2 (1)
Resuscitation (cardiological cause)	1 (1)	0 (0)	1 (0)
Pulmonary embolism	1 (1)	0 (0)	1 (0)
Revision for infection	1 (1)	3 (2)	4 (2)
Revision for haematoma	1 (1)	0 (0)	1 (0)
Closed mobilisation	0 (0)	3 (2)	3 (1)
Opticusneuropathia	0 (0)	1 (1)	1 (0)
Myocardial infarction	0 (0)	1 (1)	1 (0)
Central vein thrombosis eye	0 (0)	1 (1)	1 (0)
Subcutaneous infection belly (related to DVT prophylaxis)	0 (0)	1 (1)	1 (0)
Total SAE	5 (4)	11 (9)	16 (6)

* Adverse events unrelated to surgical treatment

confirm the impact of comorbidities expressed in elevated pcc level in our patients.

Treatment of patients suffering from severe Parkinson disease with brain stimulators is a focus of the neurological and neurosurgical department at our university

[30]. In the postoperative period these patients are closely monitored by these specialists. The reported adverse events in these patients may be an effect of this intensive care. None of these patients showed worsened mental state at the time of discharge.

Among the reported serious adverse results the optic neuropathia as well as the thrombosis of the central eye vein is unrelated to treatment to our understanding. Major surgical adverse events like revisions for haematoma or suspected infection are similar in both cohorts. Treatment for decreased motion after total knee arthroplasty was more aggressive in the second cohort as part of the clinical pathway[31]. In consequence we observed three closed manipulations only in the second cohort. There was an increase in serious adverse events for cohort II, but the difference in incidence rates were not found significant at all.

There are some limitations of the current study. We reported the incidence of adverse events for a clinical trial by means of a rather small series of 260 patients only. Larger numbers of patients could probably identify more influencing factors like age for the occurrence of adverse events and identify additional (S)AEs with smaller incidences than being observable by means of the recent sample size. As the investigation was performed at a tertiary referral centre our cases might represent a selected cohort with a higher percentage of elderly, more extended use of anticoagulative drugs and higher prevalence of comorbidities. Generalization of the above quantitative results is therefore limited; however, bearing the primary intention of proposing and illustrating a standardized reporting system, the pilot investigation ended with a positive conclusion in confirming the feasibility of the proposal demonstrated here.

In summary, further improvement in the quality of medical care depends on full recognition of adverse events. Routine assessment of adverse events by a trained medical documentation assistant or study nurse could lead to a more complete documentation and therefore to a higher rate of reported adverse and serious adverse events. The independency of study nurse assessment from physicians and the nursing team seem to be useful to achieve a more constructive discussion culture as highlighted by de Vries[32]. We therefore recommend the implementation of trained paramedical staff for the documentation of adverse events in routine clinical care.

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Authors' contributions

SK Study design, Obtaining of funding, statistical analysis and interpretation of the data, drafting and revision of the article.
JL Contribution to the study design, discussion of the results, technical and logistic support.
KPG administrative, technical, and logistic support.
MEG Support in study design with special reference to quality management.
FK Study design, Obtaining of funding; statistical expertise; analysis and interpretation of the data; administrative, technical and logistic support.
All authors read and approved the final manuscript.

Competing interests

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