Possible beneficial effect of bisphosphonates in osteonecrosis of the knee

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Abstract Osteonecrosis (ON) in the knee occurs as a localized inflammatory disease in relation to spontaneous or non-traumatic ON. Conservative treatment possibilities are limited, and prognosis appears to be poor; in most cases, ON results in knee arthroplasty. Bisphosphonates are suggested to prevent bone resorption and collapse of necrotic bone. In this observational, prospective study we investigated the effect of bisphosphonate treatment in patients with spontaneous or arthroscopy-induced ON of the knee. Twenty-eight patients with osteonecrotic lesions and bone marrow oedema in the knee were included. In 22 patients (80%), ON was identified after arthroscopic surgery of the knee; six patients were diagnosed with spontaneous ON. Patients were initially given pamidronate 120 mg i.v. divided in 3–4 perfusions over 2 weeks, followed by oral bisphosphonate treatment with alendronate 70 mg weekly for 4–6 months. Bisphosphonate treatment resulted in a rapid pain relief, VAS decreasing from 8.2 ± 1.2 at baseline to 5.02 ± 0.6 after 4–6 weeks ($p < 0.001$). After 6 months, the VAS decreased by 80% ($p < 0.001$). At the 6-month follow-up, symptoms had resolved completely in 15 patients out of 28; in 6 patients, minimal symptoms (VAS 1–2) remained. In two patients,

treatment effect was unsatisfactory, and surgical intervention was needed (arthroplasty). Bone marrow oedema on MRI resolved completely in 18 patients out of 28 with substantial reduction in the remaining. Furthermore, osteonecrotic area resolved completely or demarcation with sclerotic changes of the necrotic area could be observed. Bisphosphonate treatment in patients with osteonecrosis of the knee was associated with a rapid improvement in pain score and radiological consolidation of the area of osteonecrosis. Further randomized, controlled trials are warranted to confirm the potential beneficial role of bisphosphonates in the treatment of osteonecrosis of the knee. Level of evidence: observational study, level IV.

Keywords Osteonecrosis · Bisphosphonates · Arthroscopy · Knee

Introduction

Osteonecrosis (ON) is a localized inflammatory disease primarily affecting bone tissue. In its early stage, the overlying articular cartilage is intact; however, with progression of the disease, structural failure in subchondral bone and overlying cartilage tissue occurs [4, 7, 32, 40, 51]. Osteonecrosis of the knee is generally classified into two groups based on their underlying pathophysiological mechanisms [7, 15, 24, 40]. Spontaneous osteonecrosis of the knee (SONK) is characterized by a sudden onset of knee pain and affects predominately postmenopausal women [24, 40]. Although many causes have been proposed, the exact aetiology of ON of the knee is not entirely elucidated [17, 35, 39, 42, 46]. Primary or secondary vascular insufficiency leading to infarction of bone and minor, unrecognized traumatic incidents producing microfractures

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in the subchondral bone have been proposed [17, 35, 39, 42, 46]. Currently, the mechanical theory is favoured largely based on the finding of meniscal tears at the site of osteonecrotic condyle lesions. Histopathological studies support the theory of subchondral insufficiency fractures [23, 34, 38, 52]. The second type of osteonecrosis, secondary or atraumatic osteonecrosis, is in most cases related to medical conditions such as corticosteroid treatment, sickle cell anaemia, systemic lupus erythematosus, hyperbarism and alcohol abuse [24, 40]. The onset of pain is rather gradual, and the disease occurs usually in patients under 50 years of age, predominantly women. In most cases, other joints are also involved, bilateral affection is very common. Similar to avascular necrosis of the hip, the underlying cause seems to be systemic and mostly of vascular origin.

Osteonecrosis of the knee after arthroscopy is a relatively new diagnostic entity and has been described in several reports [6, 10, 12, 13, 20, 23, 26, 35, 39, 42, 46]. In 49 cases, in which ON has been reported after arthroscopy, the development of osteonecrosis was diagnosed secondary to treatment of meniscal and chondral lesions or reconstruction of an intraarticular ligament. In some, bone lesions were thought to be in direct relation with meniscal or chondral lesions [10, 13, 15, 20, 26, 35, 39, 42, 46]. The prognosis of ON after arthroscopic surgery remains unclear but appears to be poorer than in ordinary SONK resulting in knee arthroplasty in many cases. Furthermore, conservative treatment options are limited and mostly already exhausted.

Bisphosphonates inhibit bone resorption and are widely used in metabolic bone disease characterized by increased osteoclast activity (i.e. osteoporosis, Paget's disease) [45]. In ON, structural failure is the result of resorption of necrotic bone during revascularisation, before new bone has formed. It can be hypothesized that if accelerated bone resorption could be reduced during the revascularization process until sufficient new bone has been formed, it would appear that structural failure with subsequent osteoarthritis could be avoided. In animal studies, it has been shown that bisphosphonate treatment prevents resorption of necrotic bone during revascularization and bone deformity [8, 9, 27, 30]. In humans, bisphosphonate treatment has been used successfully in bone marrow oedema and avascular necrosis of the femoral head [3, 28, 29, 44, 49]. In these reports, the use of bisphosphonates was followed by a rapid improvement of clinical symptoms and radiological findings. To date, there is no report on the use of bisphosphonate treatment in ON of the knee. This observational, prospective study summarizes the effect of bisphosphonate treatment on the clinical and radiological course in patients with spontaneous or postarthroscopic ON of the knee.

Materials and methods

Study population

Twenty-eight patients with the radiologic finding of osteonecrotic lesions and bone marrow oedema in the knee were recruited into this open-label, prospective observational study. In 22 patients (80%), ON was identified after arthroscopic surgery of the knee. Six patients were diagnosed with spontaneous osteonecrosis of the knee (SONK).

All patients included in the study had to be symptomatic and to meet the MRI diagnostic criteria. The criteria for diagnosis of ON on MRI included normal high-intensity signal of fat in the bone marrow replaced by a discrete low-intensity signal in the centre of the lesion and high-intensity signal area about the margin representing bone marrow oedema (Fig. 1). The size of the lesion was measured on T1-weighted images as the area of low signal intensity as described by Lotke et al. [32]. Informed consent to participate in the study was obtained from all patients, according to the rules of the local ethics committee.

Intervention

Patients were initially given pamidronate 120 mg i.v. divided in 3–4 perfusions over 2 weeks, followed by oral bisphosphonate treatment with alendronate 70 mg weekly for 4–6 months. This treatment regimen was chosen in analogy to experimental animal studies in which resorption of necrotic bone with alendronate treatment was prevented by doses 4–50 times higher than the dose used in patients with osteoporosis. It was felt that an IV loading with bisphosphonate would therefore be more effective [8].

All patients were given non-steroidal anti-inflammatory or analgesics as needed. Weight-bearing was not forbidden but was usually avoided by the patients because of pain.

Calcium (500–1,000 mg/day) and vitamin D (400–800 IU/day) supplements were provided throughout the whole treatment period.

Measurements

The initial evaluation included magnetic resonance imaging (MRI) in all patients and radiographs in a subset of patients, as well as a clinical and laboratory assessment.

Clinical assessments were made before treatment, 1 month later and thereafter at 6–12 weeks intervals. A further MRI scan was performed 3 months after initiating treatment.

Blood and urine samples were taken in the morning after an overnight fast. The baseline laboratory assessment

Fig. 1 X-ray (a) and magnetic resonance image (b) of a 62-year-old female patient from the series showing typical osteonecrotic lesion and oedema of the femoral condyle

included c-reactive protein, complete blood count, serum calcium, liver- and kidney function parameters, glucose, haemoglobin A1c and screening for coagulopathies.

Calcium, creatinine and total alkaline phosphatase (ALP) were analysed by standard method on an autoanalyser (Hitachi System 704 analyser; Roche Diagnostics, Rotkreuz, Switzerland). Serum bone alkaline phosphatase (BAP) concentration was determined by ELISA (ALK-PHASE-B ELISA, Quidel) [22]. The parameters N-MID-Osteocalcin (OC) and intact parathyroid hormone (PTH) were measured in serum with electrochemiluminescence immunoassays (ECLIA) on the automated analyser Elecsys 2010 (Roche Diagnostics, Rotkreuz/Switzerland) [21, 47]. The intra- and interassay variation was 1.1–5.9% for OC, 2.6–6 and 1.7–5.5% for intact PTH. Urinary C-terminal teleopeptides of type I collagen was determined by ELISA (Nordic Bioscience Diagnostics A/S, Denmark) and corrected for creatinine [18]. The intra- and interassay CV's were 3.5 and 8.6%, respectively. Pyridinium collagen cross-links (pyridinoline [PYD] and deoxypyridinoline [DPD]) were measured by high performance liquid chromatography (HPLC) using a commercially available kit from Chromsystems Instruments and Chemicals, Munich (Germany), and the results were expressed in relation to the urinary creatinine. The intra- and interassay were 8.3 and 11.5%, respectively.

Pain was measured at baseline and during follow-up using a standardized visual analogue scale (VAS) [25].

Statistical analysis

All data are expressed as means $\pm$ standard errors of the mean (SEM). To compare study variables during study period, one-way ANOVA for repeated measures with post hoc comparisons corrected for multiple testing by least squares difference (LSD) was used. Significance was defined as $p < 0.05$. Data were analysed using Statistica for Windows (version 6.0; StatSoft, Tulsa, OK, USA).

Results

Baseline characteristics

The baseline characteristics are summarized in Table 1. Twenty-two patients had an arthroscopy for meniscal lesions or osteoarthritis, and in six patients, ON developed spontaneously. The median time interval between arthroscopy or occurrence of symptoms (in patients with SONK) and inclusion into the study was 4.0 months (range, 3 weeks to 24 months).

Laboratory investigations

Baseline laboratory investigations did not reveal any specific risk factor or disease associated with ON (Table 1). Bisphosphonate treatment resulted in an inhibition of bone

Table 1 Baseline characteristics

	Normal range	Results
Age (years)		57.7 ± 2.7
Height (cm)		164.6 ± 2.1
Weight (kg)		75.7 ± 3.3
Serum		
Calcium (mmol/l)	2.15–2.62	2.43 ± 0.02
Creatinine (μmol/l)	<97	73.4 ± 3.2
Total ALP (U/L)	35–104	79.3 ± 5.2
Bone ALP (μg/l)	7.4–15.7	13.8 ± 2.1
Osteocalcin (ng/ml)	7.7–32	21.5 ± 1.9
Intact PTH (pg/ml)	10–60	36.2 ± 5.8
25-OH Vitamin D (ng/ml)	>25	18.7 ± 3.6
Glucose (fasting) (mmol/l)	<6.1	5.3 ± 0.6
HbA1c (%)	<5.7	5.1 ± 0.7
Urine		
Calcium/creatinine (mmol/mmol)	0.1–0.32	0.25 ± 0.04
PYD/creatinine (mmol/mmol)	60–100	97.3 ± 12.7*
DPD/creatinine (nmol/mmol)	9–20	19.1 ± 2.4*
CTx/creatinine (μg/mmol)	80–320	281.6 ± 38*

Mean ± SEM/* $p < 0.05$ compared to age-matched controls

turnover, as evidenced by a significant decrease in urinary levels of CTx, PYD and DPD (already after 4 weeks) and a decrease in serum concentrations of OC, TALP and BAP (after 3 months; Table 2).

Pain and disability

The mean VAS at baseline was 8.2 ± 1.2 . As expected with such high VAS pain scores, quality of life and walking ability was severely reduced. Mobility was also affected, and patients could not walk a longer distance without walking aid.

Bisphosphonate treatment resulted in a rapid and substantial pain relief (Fig. 2). After 6 months of bisphosphonate therapy, the VAS decreased by 80% in average, and

Fig. 2 Change in visual analogue scale pain scores (VAS) in patients with osteonecrosis of the knee treated with bisphosphonates

all patients achieved a reduction of at least 50–60%. At the 6-months follow-up in 15 patients out of 27 symptoms had resolved completely and in 6 patients complained of minimal residual pain (VAS 1–2). At 1 year follow-up, clinical data on 18 patients were available, and VAS was still reduced significantly (2.2 ± 0.8 , $p < 0.001$), 14 patients reported to be free of symptoms. In 1 patient, there was evidence of moderate impression of the articular surface of the femoral condyle, however, as the patient was asymptomatic in his daily activities and in exercising (jogging) conservative treatment was continued. In two patients, treatment effect was unsatisfactory and surgical intervention (alloarthroplasty) was necessary.

Although no quantitative assessment of functional disability was performed, improvement in quality of life was apparent from the resolution of pain and considerable improvement in mobility was evident. All but two patients showed improvement in their walking ability without the needed of a walking aid.

Table 2 Biochemical parameters during follow-up

	Baseline	1 Month	3 Months	6 Months
Calcium (mmol/l)	2.44 ± 0.02	2.39 ± 0.02	2.35 ± 0.04	2.46 ± 0.03
Creatinine (μmol/l)	73.4 ± 3.3	75.2 ± 3.4	78.1 ± 5.7	81.9 ± 6.3
ALP (U/L)	79.3 ± 5.3	78.1 ± 7.2	80.8 ± 11.0	64.8 ± 4.5*
OC (ng/ml)	21.5 ± 2.0	18.9 ± 2.3	12.5 ± 1.7*	14.5 ± 2.3*
PYD (nM/mM creatinine)	97.3 ± 12.7	59.4 ± 13.0*	44.9 ± 4.2*	54.4 ± 4.4*
DPD (nM/mM creatinine)	19.0 ± 2.4	10.6 ± 2.4*	8.2 ± 0.7*	9.7 ± 0.9*
CTx (μg/mM creatinine)	231.6 ± 38.5	40.4 ± 14.0*	52.3 ± 14.7*	75.3 ± 15.0*

ALP total alkaline phosphatase; OC osteocalcin; PYD pyridinoline; DPD deoxypyridinoline; CTx c-terminal telopeptide

* $p < 0.05$ compared to baseline

Radiological findings

In all patients, the initial MRI scan showed typical appearance for ON with subchondral area of low signal and signal alteration indicative of perifocal bone oedema (low signal intensity on T1-weighted images corresponding to areas of high signal intensity on fat-suppressed T2-weighted images). At the follow-up MRI after 2–3 months of treatment bone marrow oedema resolved completely in 18 patients (out of 27), in the remaining substantial reduction in bone oedema could be observed. The osteonecrotic area resolved completely in five patients; in the others, there was demarcation with sclerotic changes of the necrotic area. No further collapse of articular surface occurred during the whole observation period.

Tolerability and side effects

In general, bisphosphonate treatment was well tolerated. Four patients experienced an acute phase reaction after IV pamidronate with arthralgias and myalgias. The symptoms, however, were moderate and the treatment course could be completed in all patients. Oral treatment with alendronate was tolerated without side effects in all patients.

Discussion

The most important finding of the present study was the rapid and sustained improvement in clinical symptoms and disability, as well as resolution of the bone marrow oedema indicating a possible beneficial effect of bisphosphonates in patients with osteonecrosis. Specifically, pain reduction (assessed by the VAS) of 30% after 4–6 weeks and of 80% after 6 months during bisphosphonate therapy could be observed. Complete or partial resolution of bone marrow oedema was documented in all cases. The size of the osteonecrotic area did not change significantly during the short observation period. However, in early stages of ON, complete resolution and in more advanced stages, demarcation of the osteonecrotic area was observed.

The development of osteonecrosis following arthroscopic knee-surgery is a rare event. In a series of publications, a total of 49 cases have been reported [6, 10, 12, 13, 20, 23, 26, 35, 39, 42, 46]. In 21 patients, in which treatment outcome has been reported surgical reintervention (9 alloarthroplasties, 1 tibial osteotomy, 1 core decompression) was needed in eleven patients. Only in five cases a favourable outcome on conservative treatment (mechanical unloading, anti-inflammatory drugs) could be observed. From these observations one might conclude that favourable treatment response on non-invasive treatment is limited.

Prognosis appears to be unfavourable in postarthroscopic ON, with progression to irreversible stages in most cases [39]. Reasons for the poor outcome could be delayed diagnosis and treatment because of existing degenerative changes before the arthroscopic intervention. Treatment options of ON depend in general on the size of the lesion, smaller lesion being managed by mechanical unloading, insole with lateral wedge and use of non-steroidal anti-inflammatory drugs [39]. Larger lesions (>50% of the femoral condyle or >5 cm²) in general require osteotomy or arthroplasty [31]. In early stages, treatment with core decompression of the bone has been successfully used [11, 19, 33]. Some authors report medical therapy with Iloprost, a prostacyclin analogue, which has been used in patients with bone marrow oedema and osteonecrosis, and in children with aseptic osteonecrosis of the proximal femur [5, 16, 41]. In these patients, treatment with Iloprost resulted in resolution of pain and rapid regression of the bone marrow oedema and subchondral lesions on MRI. In regard of the unfavourable prognosis and the limited non-surgical treatment possibility, other treatment possibilities are warranted in avoiding the need for arthroplasty.

In experimental animal studies, it has been shown that alendronate and more recently zoledronate treatment prevents resorption of necrotic bone during revascularization [8, 9]. There are several reports of successful bisphosphonate treatment of the transient osteoporosis of the hip or osteonecrosis of the femoral head [1, 2, 14, 29, 36, 37, 43, 50]. In an uncontrolled clinical study, 16 patients with avascular necrosis of the hip were treated with alendronate and an improvement in pain, disability and function compared to the expected natural history of the disease was found [3]. In a recently published observation from the same authors of 395 hips with avascular necrosis which were treated with alendronate for 3 years with a mean follow-up of 4 years improvement in the clinical function, a reduction in the rate of collapse and a decrease in the requirement for total hip replacement, compared with the findings of other studies in which no treatment was given [2]. In one of the few randomized controlled trial with alendronate treatment in 40 patients with atraumatic osteonecrosis of the femoral head with an involvement of at least 30% of the head, only one patient in the alendronate group needed a total hip arthroplasty whereas 16 subjects in the control group underwent total hip arthroplasty over an observation period of 28 months [29]. Thus, Alendronate treatment was effective in preventing early collapse of the femoral head in these patients. In another study using intravenous ibandronate in the treatment of transient osteoporosis of the hip, it has been shown that bone marrow oedema resolved completely in the MRI, and a significant pain relief could be observed [44].

In accordance with the findings in patients with avascular necrosis of the hip, bisphosphonates appear to be also beneficial in patients with osteonecrosis of the knee. The optimal dosage and duration of the bisphosphonate treatment in ON of the knee is not known. The dosage regimen used in the present study was chosen based on animal data which have shown that the prevention of resorption of necrotic bone during revascularisation is dose-dependent [8, 48]. Nevertheless, this study would support the use of bisphosphonates for the treatment of osteonecrosis and bone marrow oedema of the knee.

There are limitations in this study. First, the number of patients included in the present study is rather small, nevertheless significant reduction in pain score could be detected early after initiation of bisphosphonate therapy. Secondly, the lack of a control group does not allow comparing definitively beneficial effects of bisphosphonates against the natural course of disease. And thirdly, the follow-up period was only 6–12 months, which may have been too short to demonstrate treatment failure.

Conclusion

In conclusion, bisphosphonate treatment of osteonecrosis of the knee was associated with a rapid improvement of the clinical symptoms and disability. Also complete or partial resolution of bone marrow oedema and demarcation or resolution of the osteonecrotic area in MRI could be observed. A preventive effect against progressive destruction of the architecture of the knee following spontaneous and postarthroscopic osteonecrosis can be assumed. Further randomized, controlled trials are needed to confirm a beneficial effect of bisphosphonate therapy in osteonecrosis of the knee.

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