Osteoarthritis (OA) is characterised by progressive erosion of the articular cartilage as well as subchondral bone remodelling. It is now believed that subchondral bone modifications may be intimately linked to the actual progression of cartilage erosion in OA. The structural changes of this disease include the progressive erosion of the articular cartilage, the formation of osteophytes, and, at the clinical stage of the disease, a variable degree of synovial inflammation. A significant remodelling of the subchondral bone is believed to be predominantly an excessive bone resorption in the early stages of the disease, followed by excessive bone formation, leading to bone sclerosis and an increased thickening of the subchondral bone. This is believed, in turn, to explain the increase in subchondral bone density seen in patients with OA at the clinical stage of the disease. However, recent findings indicate that OA subchondral bone tissue comprises an excessive extracellular osteoid matrix that is undermineralised. Moreover, this matrix is composed of abnormal collagen type I heterotrimers with a predominance of collagen type 1 α1 chains over α2 chains.

The exact pathways that are involved in the pathophysiology of the disease remain largely unknown. In particular, whether subchondral bone changes in human OA precede or are simultaneous with cartilage lesions is still an unanswered question, although in animal models where OA develops spontaneously, bone changes precede cartilage lesions. The hypothesis of a link between subchondral bone changes and cartilage lesions also implies that local factors produced and/or modified by bone cells can alter articular chondrocyte function. The pathway(s) involved in such a cross communication between these two tissues also remain unresolved at present. Nevertheless, recent findings clearly indicate significant modifications of the metabolic activities of osteoblasts from both human primary OA cell cultures and in an experimental OA dog model. Several factors produced by osteoblasts might potentially affect cartilage chondrocytes. In particular, the local bone tissue production of insulin-like growth factor-I (IGF-I), urokinase plasminogen activator (uPA), and prostaglandin E2 (PGE2) may play a part in the attempt of the cartilage to promote the formation of a new extracellular cartilage matrix such as that seen in OA. Cytokines/growth factors/prostaglandins produced by subchondral bone cells may seep through the bone-cartilage interface to stimulate cartilage breakdown. Indeed, increasing evidence indicates the presence of channels and fissures between cartilage and bone that might provide a route for signals between these two tissues.

The experimental OA dog model is a reliable model that allows the study of early changes of the disease as well as several pathophysiological pathways. Reports show that in this model of OA the subchondral bone is the site of important morphological changes. As early as two to three months after surgery, the subchondral bone plate is osteopenic, and this phenomenon reversed with the appearance of bone sclerosis between 18 and 54 months after the anterior cruciate ligament (ACL). The development of osteoarthritis was also confirmed by histomorphometric studies in this model.}

**Abbreviations:** ACL, anterior cruciate ligament; COX, cyclo-oxygenase; DMEM, Dulbecco’s modified Eagle’s medium; ELISA, enzyme linked immunosorbent assay; FBS, fetal bovine serum; IGF-I, insulin-like growth factor-I; ITS, insulin-transferrin-selenium; 5-LO, 5-lipoxygenase; LTb4, leukotriene B4; NSAID, non-steroidal anti-inflammatory drug; OA, osteoarthritis; Ob, osteoblast cell cultures; 1,25(OH)2D3, 1,25-dihydroxyvitamin D; PGE2, prostaglandin E2; TGFβ, transforming growth factor β; uPA, urokinase plasminogen activator.
subchondral osteoblasts from an experimental OA dog model showed raised uPA and IGF-I levels, but did not present altered alkaline phosphatase activity or osteocalcin release.\textsuperscript{15}

Licofelone (previously named ML-3000), a new anti-inflammatory drug with combined 5-lipoxygenase (5-LO) and cyclo-oxygenase (COX)-I/COX-II inhibitory activity, has been previously shown to reduce the development of cartilage lesions in experimental dog OA.\textsuperscript{30} Here we evaluated the long term effects of treatments of experimental dog OA with licofelone on the synthetic abilities of isolated subchondral osteoblasts from these animals.

**METHODS**

**Experimental groups**

Tibial plateaus obtained from dogs from different experimental groups included in our previous study were used.\textsuperscript{30} In short, 21 adult crossbred dogs 2–3 years old, weighing 20–25 kg each, were used in this study. Surgical sectioning of the ACL of the right knee was performed on 21 dogs through a stab wound, as previously described.\textsuperscript{29–32} Before surgery, the animals were anaesthetised intravenously with pentobarbital sodium (25 mg/kg) and intubated. After surgery, the dogs were kept in animal care facilities for one week then sent to a housing farm. Dogs were housed in a large pen and were exercised under supervision to ensure that they were bearing weight on the operated knee as approved by the Institutional Committee for Animal Protection of our institution.

These dogs were randomly separated into three groups of seven: group 1, dogs with OA that received placebo (encapsulated methylcellulose); group 2, dogs with OA given encapsulated licofelone (2.5 mg/kg daily by mouth) (Merckle GmbH, Ulm, Germany); group 3, dogs with OA given encapsulated licofelone (5.0 mg/kg daily by mouth). Treatments began the day after surgery. These doses were selected based on the doses of the drug given to patients for the treatment of symptomatic OA. Licofelone was given twice daily (8 am and 4 pm) with food at the total daily dose of 2.5 and 5.0 mg/kg. All dogs were killed eight weeks after surgery. Gross morphological changes in these animals, including the presence of osteophyte formation and cartilage lesions, were reported previously.\textsuperscript{30}

**Primary bone cell cultures**

The specimens obtained from the dogs with OA represented moderate OA according to macroscopic criteria. Sections from the weightbearing areas of the subchondral bone plate of tibial plateaus were processed as previously described.\textsuperscript{15} Briefly, specimens were obtained from plug explants of the tibial plateaus collected at an equivalent anatomical site. Both the overlying cartilage and the trabecular bone sections were removed under a magnifying microscope to ensure total removal of both tissues. The remaining subchondral bone plate was then cut into small pieces (2 mm\(^2\)) before sequential digestion using 1 mg/ml of collagenase type I (Sigma-Aldrich Canada, Oakville, ON, Canada) in Ham's F12/Dulbecco's modified Eagle's medium (DMEM; Sigma) without serum, at 37°C for two periods of 30 minutes and, lastly, for 240 minutes. This removed both adherent and remaining bone marrow cells from the subchondral bone pieces. These bone pieces were then washed extensively with Ham’s F12/DMEM media without serum containing 5% Pen-Strep before being cultured in Bigger, Gwatkin, Jackson (BGI; Sigma) medium containing 20% fetal bovine serum (FBS; Wisent, St-Bruno, Quebec, Canada). This medium was replaced every two days until cells appeared in Petri dishes. At this point, it was replaced by the same medium containing 10% FBS. At confluence (days 14–18), cells were passaged once at a ratio of 25 000 cells/cm\(^2\) and grown in 24 well plates (Falcon, Lincoln Park, NJ, USA) for five days before assay. Cells prepared under these conditions show an osteoblast-like phenotype.\textsuperscript{15, 33–34} Cells from dogs with OA treated or not with licofelone grew at similar rates (not shown), indicating that cell viability was not affected by prior surgery or by treatment. Conditioning was performed for the last two days of culture, and two sets of experimental conditions were performed. Half of the cells were incubated in the presence or absence of 50 nM 1,25-dihydroxyvitamin D (1,25(OH)\(_2\)D\(_3\); generous gift of Dr M Uskokovic, Roche, Nutley, NJ, USA) for maximal stimulation, in Ham’s F12/DMEM containing 2% charcoal-stripped FBS to determine alkaline phosphatase activity and osteocalcin release. The medium was collected at the end of the incubation and frozen before assay at -80°C. Cells were washed twice with phosphate buffered saline, pH 7.4, and solubilised in alkaline phosphatase buffer (100 mM glycine, 1 mM MgCl\(_2\), 1 mM ZnCl\(_2\), 1% Triton X-100; pH 10.5) for 60 minutes with agitation at 4°C. The second half of the cells were incubated for their last two days of culture with Ham’s F12/DMEM media without FBS containing 1% insulin-transferrin-selenium mix (ITS; Sigma) and were used for the determination of uPA activity, and the determination of IGF-I and PGE\(_2\) levels. Under these conditions cell viability is maintained, and previous studies have shown that osteoblasts can still respond to exogenous modulation.\textsuperscript{33–34}

**Determination of alkaline phosphatase activity and osteocalcin release**

Cell lysates from cells treated or not with 1,25(OH)\(_2\)D\(_3\) were used for the determination of alkaline phosphatase activity as the release of p-nitrophenol hydrolysed from p-nitrophenyl phosphate (12.5 mmol/l) final concentration) at 37°C for 30 minutes. Protein determination was performed by the bicinechonic acid method on the same cell lysates.\textsuperscript{30} Osteocalcin release was measured in conditioned Ham’s F12/DMEM media (1:1) prepared for the last two days of culture of osteoblasts treated or not with 1,25(OH)\(_2\)D\(_3\). Nascent dog osteocalcin was determined by a specific radioimmunoassay (Biomedical Technologies, Stoughton, MA, USA). The detection limit of this assay is 0.5 ng/ml, and 2% charcoal-stripped FBS contains <0.1 ng/ml osteocalcin. Determinations were performed in triplicate for each assay.

**Determination of uPA activity, and of IGF-I, PGE\(_2\), and LT\(_B_4\) levels**

uPA activity was determined by the procedure of Leprince et al.\textsuperscript{35} This assay determines the activity of uPA by the hydrolysis of the specific substrate p-Val-Leu-Arg-p-nitroanilide (Sigma), which releases p-nitroaniline detectable at 405 nm. IGF-I levels were determined by a high sensitivity enzyme linked immunosorbent assay (ELISA; Diagnostic Systems Laboratories, Webster, TX, USA) that does not cross react with insulin. Samples were treated according to the method described by Mohan et al\textsuperscript{[40] to detect total IGF-1 levels. Internal control studies were performed with the media alone containing 1% ITS, and values were below the limit of detection for IGF-I. PGE\(_2\) and leukotriene B\(_4\) (LT\(_B_4\)) levels were determined by a highly specific ELISA from Cayman Chemicals (Ann Arbor, MI). The limit of quantification was 15 pg/ml and 5 pg/ml respectively for PGE\(_2\) and LT\(_B_4\). This ELISA is a very specific assay that does not cross react with related prostaglandins when tested at saturating concentrations. Determinations were performed in triplicate for each cell culture preparation.

**Statistical analysis**

Values are expressed as the mean (SEM). Statistical analysis was done with the Mann-Whitney U test. Values of p <0.05
were considered significant. Only significant values were included in the figures.

RESULTS
Macroscopic findings
Macroscopic findings for these animals have been reported previously. Only significant values were included in the figures.

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Macroscopic findings
Macroscopic findings for these animals have been reported previously. In brief, osteophytes were seen in 93% of the condyles of the OA placebo treated group, and their mean (SEM) width was 4.50 (0.66) mm. Osteophytes were still present in licofelone treated groups at 93% and 86% in the dogs treated with 2.5 mg/kg/day and 5.0 mg/kg/day, respectively and their widths were not statistically smaller in the licofelone treated groups (3.57 (0.56) and 3.86 (0.66) mm, respectively). Cartilage lesions of moderately severe grade were seen in both condyles and plateaus, with more severe lesions on the plateaus. Licofelone reduced the size of lesions significantly on the plateaus of both groups by 39% and 64% in the 2.5 and 5.0 mg/kg/day groups, respectively (p < 0.05 and p < 0.04, respectively compared with the OA placebo treated group).

Phenotypic markers of osteoblasts
Osteoblasts from all groups of dogs had normal microscopic appearance. The level of basal alkaline phosphatase activity in primary dog osteoblasts was not increased significantly in response to 1,25(OH)2D3 treatment (fig 1), a situation previously observed with dog osteoblasts. Licofelone failed to alter this activity under either basal or 1,25(OH)2D3 induction. In contrast, treatment with 1,25(OH)2D3 stimulated osteocalcin release from relatively low basal levels in all groups (fig 2); however, licofelone did not significantly alter osteocalcin release compared with placebo. There was a tendency for the 2.5 mg/kg/day group to show higher levels of 1,25(OH)2D3 induced osteocalcin release, but this failed to reach significance.

Figure 1 Evaluation of alkaline phosphatase activity of in vitro osteoblasts after in vivo treatment with placebo or with 2.5 mg/kg or 5.0 mg/kg licofelone daily for eight weeks of dogs with OA. Confluent primary osteoblasts were incubated for their last two days of culture in the presence of 50 nM 1,25(OH)2D3. Alkaline phosphatase activity was determined by the hydrolysis of p-nitrophenyl phosphate into p-nitrophenol. Values are the mean (SEM) of seven cell cultures for each experimental group. No statistical differences were seen between the groups.

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Figure 2 Evaluation of osteocalcin release of in vitro osteoblasts after in vivo treatment with placebo or with 2.5 mg/kg or 5.0 mg/kg licofelone daily for eight weeks of dogs with OA. Confluent primary osteoblasts were treated with the same experimental protocol as in fig 1. Osteocalcin release was determined on aliquots of conditioned media by radioimmunoassay. Values are the mean (SEM) of seven cell cultures for each group. No statistical differences were seen between the groups. OA compared with controls. In vivo licofelone treatment was able to reduce this in vitro activity in long term culture of OA Ob by about 50% and 60% for the 2.5 and 5.0 mg/kg/day groups, respectively (fig 3). Likewise, licofelone reduced total IG-F-1 synthesis in these cells by 30% and 50% for the 2.5 and 5.0 mg/kg/day groups, respectively (fig 4). PGE2 levels were still very high in the placebo group despite long term in vitro Ob, and licofelone reduced these levels significantly. Indeed, both doses of licofelone also had a profound effect on PGE2 production, especially in the 2.5 mg/kg/day group where it reduced PGE2 production by about 80% (fig 5). In contrast with PGE2, LTB4 levels were very low in the placebo group (28.6 (2.6) pg/mg protein) and remained unchanged in response to licofelone treatment (25.9 (10.2) and 39.4 (3.0) pg/mg protein for the 2.5 and 5.0 mg/kg/day groups, respectively). The reduction in uPA activity, IG-F-1, and PGE2 levels was paralleled by the reduction seen in the size of the lesions of the plateaus in these dogs (fig 6). Indeed, licofelone reduced the size of plateaus by about half in both groups; however, the difference between these two groups was not statistically significant.

Figure 3 Evaluation of urokinase plasminogen activator (uPA) activity in dog osteoblasts. Dogs were treated with placebo or with 2.5 mg/kg or 5.0 mg/kg licofelone for eight weeks before the preparation of primary osteoblast cell cultures. Confluent primary dog osteoblasts were incubated for their last two days of culture in the presence of Ham’s F12/DMEM without serum containing 1% ITS mix. uPA activity was determined on aliquots of conditioned media using a chromogenic substrate. Values are the mean (SEM) of seven cell cultures for each group.
DISCUSSION

As we previously showed that licofelone reduced the progression of experimental OA in our dog model,\(^{30}\) we wanted to determine whether this new combined inhibitor of 5-LO and COX might influence osteoblasts from the subchondral bone plate and grown in in vitro culture. As this technical approach requires long term culture of these cells and we could expect a washout of the drug, it is noteworthy that licofelone could still exert an effect in vitro under these conditions. This indicates that either the drug itself has a long half life in vivo as well as in vitro, or that the effects it initiated in vivo in the bone tissue were carried over into in vitro experiments. This is similar to the observation we previously made in this same experimental OA dog model with another non-steroidal anti-inflammatory drug (NSAID), carprofen.\(^{15}\) Both studies suggest that osteoblasts are intimately involved in the development/progression of OA, and that any intervention that modifies abnormal bone cell metabolism under these conditions could contribute to preventing the onset or reducing the progression of OA. In this animal model of OA, morphological subchondral bone changes were observed consistent with altered bone remodelling. Indeed, the bone surface and trabecular width were reduced compared with normal dog bone tissues, whereas the number of osteoclasts per mm\(^2\) was raised in OA bone tissue compared with normal. Licofelone treatment of dogs with OA significantly improved all these measures (manuscript in preparation).

Because licofelone was very effective and is a combined inhibitor of 5-LO and COX, this raises the possibility that both pathways may be involved in the onset/progression of OA. This last hypothesis is reinforced by our recent observation that both PGE\(_2\) and LTB\(_4\) levels are raised in human OA subchondral osteoblasts, and that licofelone reduced the synthesis of both eicosanoids in in vitro experiments.\(^{23}\) Moreover, the observation that chronic reduction of PGE\(_2\) synthesis by COX-2 inhibition in human OA subchondral osteoblasts leads to increased LTB\(_4\) synthesis argues in favour of both pathways being involved in this process.\(^{23}\)

Our study indicated that licofelone had no adverse effects on cell proliferation or differentiation of osteoblasts as it did not significantly modify either alkaline phosphatase activity or osteocalcin release. The lack of effect of 1,25(OH)\(_2\)D\(_3\) on alkaline phosphatase activity is specific for dog osteoblasts as this activity is usually stimulated by 1,25(OH)\(_2\)D\(_3\) treatments in other osteoblast-like cells.\(^{4,5,13,18}\) We previously observed similar behaviour of dog osteoblasts in vitro.\(^{15}\) However, this situation is not related to a lack of response to 1,25(OH)\(_2\)D\(_3\) because osteocalcin release, which is strictly dependent on 1,25(OH)\(_2\)D\(_3\) induction in osteoblasts,\(^{18}\) was stimulated by vitamin D in this study.

The fact that licofelone did not modify any of these activities indicates that this drug does not modify the osteoblast phenotype, a situation also observed with another NSAID, carprofen.\(^{15}\) Indeed, neither alkaline phosphatase activity nor osteocalcin release from OA dog osteoblasts was modified compared with normal dog osteoblasts or after carprofen treatment in this last study. This observation with the in vivo dog model of OA is different, however,
from the situation with human OA subchondral osteoblasts, which showed increased alkaline phosphatase activity and osteocalcin release compared with normal.46 This may be related to the naturally occurring disease state in humans as opposed to its induction in the experimental OA dog model.

In contrast, uPA activity and IGF-I levels were raised in vitro osteoblasts isolated from this experimental OA dog model. We previously showed similar data for OA dog osteoblasts compared with normal dog osteoblasts,44 a situation also similar to that observed in human OA subchondral osteoblasts, indicating that these metabolic modifications are important pathways in disease progression. This is further indicated by the fact that licofelone caused a decrease in both uPA activity and IGF-I levels, reducing them to levels similar to those seen in normal dog osteoblasts.45 The effect of licofelone on uPA activity and IGF-I levels may be related to its reduction of PGE2 levels. Indeed, we and others previously showed that PGE2 can regulate IGF-I synthesis in osteoblasts, whereas its effect on uPA activity was never assessed. The observation that the modifications of both uPA and IGF-I follow the inhibition of PGE2 synthesis observed with licofelone suggests a possible link between uPA and PGE2. However, this will have to be tested directly. Abnormal bone remodeling sites.21 As licofelone is a specific inhibitor for 5-LO and COX-I/II, it may not be able to modulate in vivo TGF

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