# CLINICAL ARTICLE

# Evaluation of peri-articular aggressive benign bone tumours treated with cementation and a locked plate

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# **Abstract**

### **Background**

Treatment of aggressive benign bone lesions with curettage, burring, cementation and plate augmentation is a widely accepted treatment. We have used the above method utilising a locked plate, rather than conventional plating methods, facilitating immediate stability and early mobilisation. We hypothesised that this joint-preserving method would provide an economic alternative to megaprosthetic joint replacement without compromising good functional outcomes.

#### **Methods**

Patients with peri-articular aggressive benign bone lesions were treated with marginal excision, curettage, burring and cementation. This was augmented with a locked plate. Liquid nitrogen was used as an adjunct where feasible. Functional assessment and radiological recurrence was assessed at follow-up using the Musculoskeletal Tumour Society Score (MSTS) and routine X-rays.

#### Result

The average MSTS score was 88%. Average follow-up time was 21 months. To date, no revisions have been performed for recurrence or pain and no significant complications have occurred.

# Conclusions

Our early results in a small series make us optimistic that this may be an alternative to immediate megaprosthetic reconstruction; furthermore, this may provide a useful intermediate step in the treatment of aggressive benign peri-articular bone tumours with some degree of associated joint deformation without significantly compromising later prosthetic reconstruction.

Key words: aggressive benign tumours, bone cementation, locked plate

# Introduction

The treatment of aggressive benign peri-articular bone tumours is challenging. Close proximity to joint surfaces makes excision and reconstruction difficult and functional impairment likely. This, together with the fact that the majority of these tumours occur in younger patients who have a potentially long economically active life span, makes it imperative that optimal postoperative function is achieved.

Limb salvage options include:

- wide resection and endoprosthetic reconstruction of the joint
- · wide resection and bulk allograft reconstruction
- wide resection and excision arthroplasty or arthrodesis
- marginal excision, cementation +/- plate augmentation +/- chemical adjuncts (e.g. liquid nitrogen or phenol)
- marginal excision, bone graft +/- plate augmentation +/- chemical adjuncts (e.g. liquid nitrogen or phenol) Marginal excision and cementation is the treatment of choice for primary as well as recurrent peri-articular benign bone lesions.1 Furthermore, it is an acceptable surgical option in the treatment of aggressive benign bone lesions. 1,2,3 Gitelis et al.1 compared en-bloc resection of giant cell tumours and arthroplastic reconstruction, allograft reconstruction or arthrodesis of the joint to intralesional excision and cementation. He noted that all were excellent procedures but better functional results were achieved where the joint surface had been preserved. Although recurrence rates are somewhat higher with a limited resection,2 the better functional outcomes achieved with joint preservation surgery may be preferable.1,4 Indeed, consistently better functional outcomes have been noted in limb salvage procedures where the joint is preserved in comparison to those where endoprosthetic reconstruction has been performed. MSTS functional scores greater then 90% have consistently been achieved where the joint is preserved. 1,3,4 as opposed to MSTS scores of 70-80% with joint replacement .5,6

Wide resection margins and allograft reconstruction is associated with a number of problems. In addition to relatively compromised function, there is a reported higher incidence of complications.<sup>7</sup> These include a very high infection rate, neurovascular damage as well as non-union and pathological fracture.<sup>8,9</sup>

The safety of polymethylmethacrylate (PMMA) cement has been proven, although potential chondrolysis and cement fracture have resulted in some surgeons replacing PMMA cement at 2 years post-op or using bone graft or bone graft substitutes instead.<sup>2</sup> As well as its structural properties, PMMA cement has the advantage of causing tumour necrosis by means of the exothermic polymerisation process or the presence of toxic monomers.<sup>1</sup> It also facilitates easier detection of tumour recurrence on plain X-rays and MRI.<sup>10</sup>

Endoprosthetic reconstruction following tumour resection is associated with several disadvantages. Extensive soft tissue dissection may result in delayed mobilisation and even instability. Prolonged postoperative physiotherapy and hospitalisation are often required. Despite this, full range of motion may not be regained and subsequent functional impairment results. Difficulty in aligning megaprosthetic implants may result in early failure. Younger age group, accelerated implant wear and aseptic loosening may necessitate several revision surgeries. Arthroplasty is very expensive and the complication rate tends to increase with each revision operation, especially infection and wound complications. Post-surgical infection rates of up to 16%, following megaprosthetic reconstruction, have been reported.11 This is particularly evident when wide excision has been performed as the initial treatment.

We hypothesised that these disadvantages may be minimised by rather utilising a marginal resection of the tumour, high-speed burring of the tumour bed and PMMA cementation augmented with a locked plate. This would confer several advantages:

- Early mobilisation and rapid return of function could be expected due to reduced soft tissue dissection and immediate postoperative stability.
- The locked-plate PMMA cement construct is relatively cheap when compared to joint replacement.
- · The incidence of infection is lower.
- · Conversion to a later endoprosthesis could still be possible.
- The fixation method would also suffice for the treatment of concurrent pathological fractures.

This study was devised in order to assess the functional outcome of patients suffering from aggressive benign peri-articular tumours treated in this manner.

### Materials and methods

A retrospective review of 15 patients, who presented to the Tumour and Sepsis Unit at the Charlotte Maxeke Johannesburg Academic Hospital between March 2005 and March 2009 suffering from aggressive benign periarticular bone tumours, was performed. Only 14 patients were available for follow-up, as one patient failed to attend any follow-up clinics. There were seven male and seven female patients. The average age of the patients was 28 years, with ages ranging from 14 to 63 years. All 14 patients presented with a single clinically benign bone tumour; 13 of the 14 tumours were graded as Enneking grade 3 and further classified as Campanacci grade 3 (in the case of the giant cells tumours). One patient was staged as Enneking grade 2. There were seven giant cell tumours (Figures 1 and 2), three chondroblastomas, one aneurysmal bone cyst, one chondromyxoid fibroma, one enchondroma and one fibrous dysplasia. Nine of the tumours were located around the knee, with five occurring in the proximal tibia and four in the distal femur.



Figure 1. Pre-operative AP X-ray demonstrating a distal femur giant cell tumour



Figure 2. MRI scan of giant cell tumour

Two were found in the proximal humerus, one in the distal radius, one in the proximal femur and one in the distal tibia. Eight of the lesions were associated with pathological fractures and six of the lesions revealed eroded articular cartilage and were, therefore, intra-articular. All of the tumours were larger than 10 cm in diameter, and occupied more than 50% of the width of the subchondral bone. Average follow-up time was 21 months (range: 3 months-54 months) (Table I).

All patients were treated with a marginal excision of the tumour followed by burring, curettage and PMMA cementation. This was augmented with a locked plate (Figures 3, 4 and 5). A variety of plates was used from different suppliers; however, all were similar in that they provided a rigid single construct by means of the capability of locking the screws within the plate. Liquid nitrogen was used as a further adjunct whenever feasible. In six of the 14 patients the tumour had eroded through the articular cartilage into the joint preventing the use of liquid nitrogen. All patients were operated on by the same surgeon. Physiotherapy was commenced on the second postoperative day. Patients who had tumours of the lower limb were encouraged to weight bear as soon as possible. Functional assessment was performed at follow-up using the Musculoskeletal Tumour Society Score (MSTS Score)11,12 at 6 weeks, 3 months, 6 months, 12 months and 6-monthly thereafter, and possible recurrence was assessed clinically and radiologically (Table II). The MSTS scoring system assesses the functional outcome of patients after tumour treatment. It consists of six categories. Both the upper and lower limb scores have categories of pain, emotional acceptance and function. The upper limb score has additional categories of hand-positioning ability, manual dexterity and lifting ability. The lower limb score has additional categories of walking ability, gait and use of supports. Points are given for abilities in each category with the total out of 30. This score is then converted to a percentage, with the higher the percentage the better the functional outcome (Figures 6 and 7). Radiological recurrence would be evident as an increasing radiolucent line at the bone-cement interface.

## Results

On functional evaluation at last follow-up the average MSTS score was 88% (range: 63%-100%). Nine of the 14 patients had a MSTS score of greater than 90%. All patients who had tumours of the lower limb were partially weight bearing at 2 weeks postoperatively and where these occurred around the knee, flexion of at least 90° was achieved by the time of discharge and an average MSTS score of 89%. The average MSTS score among the upper limb group was 84% with all patients able to return to their former occupations.

Radiographically there has been no evidence of recurrence in any of the patients. Twelve of the patients exhibit no radiolucent line at the bone-cement interface.

Table I: Demographic data of enrolled patients								
Patient	Sex	Age	Diagnosis	Grade of Tumour	Site	Date of Surgery	Follow Up (Months)	MSTS Score
TS	F	33	Giant cell tumour	Campanacci III	Distal femur	03/03/2005	54	100%
MN	М	19	Giant cell tumour	Campanacci III	Proximal tibia	July 2005	44	97%
TM	М	23	Giant cell tumour	Campanacci III	Proximal tibia	24/2/2006	42	70%
GN	F	18	Chondroblastoma	Enneking III	Distal femur	13/7/2009	24	100%
AS	М	27	Fibrous dysplasia	Enneking III	Proximal femur	18/6/2007	24	70%
MS	F	63	Giant cell tumour	Campanacci III	Distal radius	26/11/2007	20	63%
CN	F	14	Chondroblastoma	Enneking III	Proximal tibia	11/12/2007	18	100%
GM	М	18	Giant cell tumour	Campanacci III	Distal tibia	21/4/2008	15	100%
TM	М	19	Chondroblastoma	Enneking III	Proximal tibia	12/5/2008	15	97%
MT	F	19	Giant cell tumour	Campanacci III	Distal femur	5/5/2008	13	77%
NA	F		Aneurysmal bone cyst	Enneking III	Proximal humerus	Feb 2009	6	90%
MM	F	52	Enchondroma	Enneking III	Proximal humerus	March 2009	5	100%
PN	F	37	Chondromyxoid fibroma	Enneking II	Proximal tibia	March 2009	5	97%
SM	М	33	Giant cell tumour	Campanacci III	Distal femur	15/5/2009	3	70%

The remaining two patients display a radiolucent line of 1 mm, which was evident on the immediate postoperative X-rays and remained constant in size. In fact, two of the patients displayed hypertrophy of subchondral bone.

To date there have been no significant complications. All patients have returned to their former occupations. One patient, however, was re-admitted for a week for intensive physiotherapy. Another patient, whose follow-up time was 4.5 years, is a potential candidate for revision surgery using a megaprosthetic knee replacement due to radiologically advanced osteoarthritic changes in his knee. He feels his functioning and pain levels are within acceptable limits and is refusing further surgery at present.

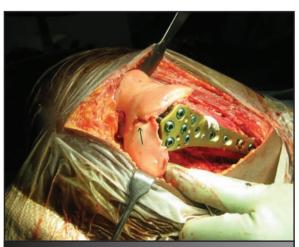


Figure 3. Intra-operative resection of giant cell tumour. Arrow indicates cartilage fracture and intra-articular extension.

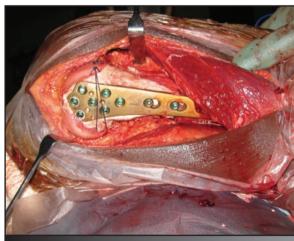


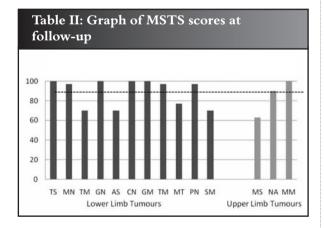
Figure 4. Final result of tumour resection, with cementation and locked plate augmentation

# Discussion

It is well documented that it is desirable to maintain the joint surface in patients with peri-articular bone lesions when performing limb salvage surgery.<sup>2-4</sup> This preservation of the joint is associated with better functional outcomes as well as avoiding the complications associated with megaprosthetic replacement. This was confirmed in our study with the average MSTS score of 88% as opposed to the studies of Hayashi and Per-Ulf Tunn,<sup>4-6</sup> where average MSTS scores of 88% and 74% respectively were documented. The average MSTS score achieved in our study is lower than that achieved by Gitelis *et al*.



Figure 5. Postoperative AP X-ray of patients in previous figures



The use of a locking plate confers immediate stability by providing a single rigid construct together with the adherent cement

However, he divided his patients into two groups. The first group consisted of patients with giant cell tumours staged as Campanacci grade 2 and 3, with the second group consisting only of tumours graded as Campanacci 2. The first group was treated with wide resection and joint reconstruction whereas the second group was treated with a joint-preserving surgery. All of our patients, except for one, were graded as Campanacci 3 or Enneking 3, with eight of the lesions associated with pathological fractures, and six of the lesions revealed eroded articular cartilage and were, therefore, intra-articular. The patient's tumour, graded as an Enneking grade 2, extended across more than 50% of the subchondral surface and was therefore included in the aggressive benign group of peri-articular tumours.

The use of a locking plate confers immediate stability by providing a single rigid construct together with the adherent cement. This facilitated early rehabilitation which we feel is associated with a better functional outcome. All of our patients with tumours affecting the lower limb were partially weight bearing at 2 weeks postoperatively. At last follow-up, only one patient utilised a ranger brace when walking more than 500 m; the rest mobilised completely independently. In addition, the apparent hypertrophy of the subchondral bone, clearly demonstrated in two of our patients, may further corroborate the functioning of this single rigid construct. This rigid construct enables appropriate loading of the joint and avoids shear forces, which may lead to premature failure. However, further investigation and follow-up is necessary to confirm this. This group of patients presented with tumours that had resulted in pathological fractures or extended intra-articularly or occupied more than 50% of the subchondral bone. They were therefore classified as aggressive in nature and benefited from the use of a locked plate construct. Smaller, less aggressive tumours or those not peri-articular will not necessarily benefit from a locked plate construct.

The patient series of De Camargo *et al.*<sup>2</sup> reported a 12% incidence of osteoarthritis at long-term follow-up of 24 years. This was evident within patients graded as Campanacci 3 tumour (n=25), who were treated with curettage, cauterisation and cementation. In our study one of our patients (1/14=7%) demonstrated evidence of osteoarthritis at 3.5 years post-op, notably within the lateral compartment of his knee, which had been destroyed by the tumour. De Camargo *et al.* also reported a 6% infection rate. Two other studies, one reporting the incidence of infection in allograft, and one reporting that occurring in megaprosthetic reconstruction, confirmed infection rates of greater than 10%. None of our patients developed any infection.

Four of the patients in our series had MSTS scores of 70% or less. Two of these patients presented with an intraarticular fracture, with one of them developing frank osteoarthritis, within the follow-up time period.

1. Pain score		
Description	Data	Numeric value
No pain	No medication	5
Intermediate		4
Modest / non disabling	Non-narcotic analgesia	3
Intermediate		2
Moderate / intermittently disabling	Intermittent narcotics	1
Severe / continuously disabling	Continuous narcotics	0
2. Function		
Description	Data	Numeric value
No restriction	No disability	5
Intermediate		4
Recreational restriction	Minor disability	3
Intermediate		2
Partial occupational disability	Major disability	1
Total occupational disability	Complete disability	0
3. Emotional acceptance	complete sections,	•
	P. I.	p
Description	Data	Numeric value
Enthused	Would recommend to others	5
Intermediate		4
Satisfied	Would do again	3
Intermediate		2
Accepts	Would repeat reluctantly	1
Dislikes	Would not repeat	0
4. Walking ability		
Description	Data	Numeric value
Unlimited	Same as preoperative	5
Intermediate		4
Limited	Significantly less	3
ntermediate		2
Inside only	Cannot walk outside	1
Not independently	Walk with assistance / wheelchair bound	0
5. Gait	Walk With addictation / Whodishair bound	•
Description	Data	Numeric value
Normal	No alteration	5
Intermediate		4
Minor cosmetic	Cosmetic alteration only	3
Intermediate		2
Major cosmetic	Minor functional deficit	1
Major handicap	Major functional deficit	0
6. Supports		
	Dete	Numerie value
Description	Data	Numeric value
		5
None	No support	
None Intermediate	Occasional use of brace	4
None ntermediate Brace	Occasional use of brace  Mostly brace	4 3
None Intermediate Brace Intermediate	Occasional use of brace  Mostly brace  Occasional cane / crutch	4 3 2
Description  None Intermediate Brace Intermediate 1 cane / crutch	Occasional use of brace  Mostly brace Occasional cane / crutch Mostly cane / crutch	4 3 2 1
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Intermediate		2
Partial occupational disability	Major disability	1
Total occupational disability	Complete disability	0
3. Emotional acceptance		
Description	Data	Numeric value
Enthused	Would recommend to others	5
Intermediate		4
Satisfied	Would do again	3
Intermediate		2
Accepts	Would repeat reluctantly	1
Dislikes	Would not repeat	0
4. Hand positioning		
Description	Data	Numeric value
Unlimited	180° Elevation	5
Intermediate		4
Not above shoulder / no pro- or supination	90° Elevation	3
Intermediate		2
Not above waist	30° Elevation	1
	0 Elevation	0
None	U Elevation	U
	o cievation	U
5. Manual dexterity		
5. Manual dexterity Description	Data	Numeric value
5. Manual dexterity  Description  No limitations		Numeric value 5
5. Manual dexterity  Description  No limitations  Intermediate	Data  Normal dexterity and sensation	Numeric value 5 4
Description No limitations Intermediate Loss of fine movements	Data	Numeric value  5 4 3
Description No limitations Intermediate Loss of fine movements Intermediate	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)	Numeric value  5 4 3 2
None  5. Manual dexterity  Description  No limitations Intermediate Loss of fine movements Intermediate Can't pinch	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation	Numeric value  5 4 3 2
Description No limitations Intermediate Loss of fine movements Intermediate Can't pinch Can't grasp	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)	Numeric value  5 4 3 2
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Description No limitations Intermediate Loss of fine movements Intermediate Can't pinch Can't grasp Can't grasp Can't grasp	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation	Numeric value  5 4 3 2
Description No limitations Intermediate Loss of fine movements Intermediate Can't pinch Can't grasp Can't grasp Can't grasp Can't grasp Can't grasp Can't grasp	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation  Anaesthetic hand	Numeric value  5 4 3 2 1
Description No limitations Intermediate Loss of fine movements Intermediate Can't pinch Can't grasp 6. Lifting ability Description Normal	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation  Anaesthetic hand  Data	Numeric value  5 4 3 2 1 0 Numeric value
Description No limitations Intermediate Loss of fine movements Intermediate Can't pinch Can't grasp Can't grasp Can't grasp Can't grasp Country Countr	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation  Anaesthetic hand  Data  Matches normal	Numeric value  5 4 3 2 1 0  Numeric value
Description No limitations Intermediate Loss of fine movements Intermediate Can't pinch Can't grasp 6. Lifting ability Description Normal Intermediate Limited	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation  Anaesthetic hand  Data  Matches normal  Mild limitation	Numeric value  5 4 3 2 1 0  Numeric value  5 4 4 4 4 4 4 4 4 4 6 6 7 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8
Description  No limitations Intermediate Loss of fine movements Intermediate Can't pinch	Data  Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation  Anaesthetic hand  Data  Matches normal  Mild limitation  Minor load	Numeric value  5 4 3 2 1 0  Numeric value  5 4 3 3
Description No limitations Intermediate Loss of fine movements Intermediate Can't pinch Can't grasp 6. Lifting ability Description Normal Intermediate Limited Intermediate	Data Normal dexterity and sensation  Can't button etc. or mild loss of sensation (Specify)  Major loss of sensation Anaesthetic hand  Data  Matches normal Mild limitation Minor load Gravity only	Numeric value  5 4 3 2 1 0  Numeric value  5 4 3 2 1 0  1 0  1 0  1 0  1 0  1 0  1 0 0  1 0

It is likely that those patients with intra-articular extension will develop some form of osteoarthritis and potentially require conversion to a megaprosthetic replacement in the future. The third patient with a score of less the 70% was the oldest patient in the series, who presented with a giant cell tumour of the distal radius. Fine motor co-ordination and dexterity of the hand are difficult to restore

In our opinion this form of treatment is best suited to lower extremity tumours, where the main function to be restored is weight-bearing and locomotion.

This treatment plan is not intended to replace megaprosthetic replacement of the joint. On the contrary, in several of our patients we anticipate the necessity for this at some stage. This method may assist in postponing arthroplasty and thereby gaining valuable years of bone and joint-preserving function. This is of extreme importance to young patients in whom the longevity of a megaprosthetic joint replacement may be suboptimal.

The use of a limited surgical dissection and smaller metal implant may be associated with lower rates of post-operative infection, as has been demonstrated in our study. In a country such as South Africa, where there is a high HIV prevalence, infection may be extremely difficult to treat and any method which could diminish infection rates is desirable in limb salvage surgery. There are also economic considerations, particularly in a developing world setting, with our construct being a far more affordable alternative.

Some of the drawbacks of our study are its small sample size, consisting of only 15 patients, as well as our relatively short follow-up of 21 months. In addition, although this was a retrospective analysis, no aggressive giant cell tumours have ever been treated in our unit with initial – or subsequent — megaprosthetic replacement, with this being our preferred method of treatment.

# Conclusion

Notwithstanding our small study and limited follow-up time, the results we have obtained to date appear to indicate that the use of a marginal excision, combined with a rigid locked-plate cement augment, may be effective in treating young patients suffering from peri-articular and even intra-articular aggressive benign bone tumours.

The content of the article is the sole work of the authors. No benefits of any form have been received from a commercial party related directly or indirectly to the subject of this article. Informed consent was obtained from all participants in this study.

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