The Journal of Maharashtra Orthopaedic Association June - 2006

Biopsy For Musculoskeletal Tumors -An Orthopaedic Surgeons Guide

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A biopsy is perhaps the most important step in the workup of bone tumors. A histological diagnosis establishes the identity of the neoplasm before any treatment is offered. In a way it is like the final conclusive identification of the "criminal" where clinical and imaging clues help us narrow down the possibilities. A badly performed biopsy is the most common iatrogenic complicating factor in this era of limb salvage surgery. Mankin¹ reported a 15.9% complication rate with 3% unnecessary amputations in his analysis of data of 597 patients from questionnaires sent to members of the musculoskeletal society. While the procedure by itself is not technically demanding, it has to be meticulously planned in concordance with the final procedure to be carried out. Again, in this modern era the histopathology is not an end in the diagnosis. Immunohistochemistry, microscopy and cytogenetics are additional tools in complicated cases. This chapter focuses on the practical aspects of the biopsy and attempts to answer common questions like how, when and which tumors to biopsy. It is important that a clinician knows how to obtain adequate material in the least traumatic way and without jeopardising the limb salvage procedure. Ill-advised and improperly performed biopsy frequently reduces the patient's chance of a limb saving surgery or even cure. Infact a poorly placed biopsy incision, a badly placed drain, or the complications of a biopsy are the commonest cause for an amputation instead of an otherwise possible limb salvage surgery.

When should the biopsy be done?

"Biopsy should be regarded as the final

diagnostic procedure, not as a shortcut to diagnosis."

This statement by Jaffe² sums up the answer to the question posed above. The biopsy should be done only when all imaging studies have been completed. The optimum integration of clinical and radiographic information prior to biopsy has important implications for the diagnosis of bone tumors, and is necessary for accurate pathologic interpretation.^{3,4}. A surgeon must resist the temptation of getting to an instantaneous diagnosis by a hasty biopsy. In such a situation the pathologist has the most difficult task of giving a diagnosis purely from the microscopic appearance. This can be dangerous especially for the bone and soft tissue tumors which are known to be extremely heterogenous. The correct approach is to narrow down the differential diagnosis. The histopathology merely confirms the strongly suspected diagnosis. This is where the mutidiciplinary cooperation between radiologist, clinician and the pathologist becomes vital. Prior imaging helps to narrow down the differential diagnosis. Lesions like lipomas, hemangiomas and aneurysmal bone cysts can be confidently identified by imaging, especially an MRI and may not require a formal biopsy prior to intervention. Imaging can also help identify the best areas to biopsy. The sclerotic, ossified, calcified or necrotic areas will not yield tissue for a diagnosis. In most instances the diagnosis is obvious on imaging and a biopsy merely confirms the diagnosis.

Do all radiological lesions need a biopsy ?

As a rule all suspected malignant tumors and aggressive benign tumors should be biopsied prior

to treatment. it is often difficult to be certain about the nature of some tumors. For example, some osteosarcomas resemble a classical giant cell tumor on xray. Conversely some giant cells appear extremely aggressive resembling malignant tumors on imaging. The surgeon as well as the pathologist must be familiar with radiology of bone tumors. Osteochondromas and enchondromas or fibrous cortical defects which are asymptomatic or latent do not need a biopsy. In certain conditions, the final treatment can be carried out at the same time as a biopsy; e.g., in a unicameral bone cysts (UBC) and histiocytosis, which are benign and have a characteristic appearance on xray, once the needle biopsy confirms the diagnosis (on frozen section), a steroid can be injected through the same needle. Osteoid osteomas can be treated by coring or surgery without a histological diagnosis. Some typical lesions like a GCT in the distal radius or a non-ossifying fibroma(NOF) in the distal femur can be treated by biopsy and surgery in the same sitting if the surgeon is experienced in tumors and the pathologist is certain about the diagnosis on frozen section. This may be the situation in just a handful of centres today and it may be safest to biopsy and get a firm diagnosis prior to final management. In cases of a local recurrence or metastatic disease where the primary diagnosis is known, one may safely proceed with management after a frozen section confirms the diagnosis. Even in situation of metastases, in particular solitary metastases in the proximal femur, a primary tumor like a chondrosarcoma must be ruled out by a frozen section biopsy prior to final management. A common error is to assume a diagnosis of chronic osteomyelitis and proceed with debridement without a biopsy only to find out later that it was a sarcoma.

Who should do the biopsy?

At several centres in the world, the needle biopsies, especially the ones requiring image guidance, are done by the interventional radiologists. At other places the pathologist does the biopsy to ensure adequacy of material. In both these situations tremendous cooperation and teamwork is necessary to ensure correct placement and technique. The biopsy is always done in consultation with the surgeon who will be deciding the final management. In most centres including ours, the biopsy is done by the surgical team. This is ideal as the site is chosen correctly and biopsy done in a manner not to compromise the final limb salvage procedure. A surgeon who has first seen the patient should resist the temptation of doing a biopsy himself unless he is familiar with the limb salvage procedures. As alluded to earlier, Mankin's article points out that even in advanced centres in the west, biopsy related complications have often been the cause of an unnecessary amputation¹. The rate of these complications is higher when the biopsy is done at centres not specialized in oncology¹.

What is the correct site ?

Regardless of the type of biopsy it's placement is critical. For appropriate placement of the biopsy, the surgeon needs to know the probable diagnosis and the extent of the tumor and should have established an operative plan prior to biopsy. He should not be concerned only with obtaining a tissue diagnosis but should also think about the definitive operative procedure.

Transverse incisions in the extremities are almost always contraindicated because the site of the incision cannot be excised en bloc with the longitudinally directed segments of bone or musculo aponeurotic compartments. Therefore, a longitudinal biopsy incision must always be used in the extremity. The major neurovascular structures should be avoided because if they are contaminated during the biopsy they may have to be sacrificed during the definitive procedure that follows. The biopsy tract also should not traverse a normal anatomical musculoskeletal compartment in order to reach a compartment that is involved by tumor, so that it will not be necessary to remove both compartments at the time of the definitive procedure. It is best to violate only one compartment. The joint should never be violated; attention to this is important especially around the knee. As a rule no biopsy should be done through the joint or arthroscopic for any aggressive tumor.

Standard operative approaches employed in orthopaedic procedures may prove inappropriate for a biopsy. As an example, biopsy of the humerus through the deltopectoral interval causes dissemination of tumor cells at a distance through normal neurovascular planes. It would be more appropriate to biopsy the tumor through the anterior deltoid (Fig. 1) so as to contain the hematoma and then to resect en bloc the biopsy-contaminated deltoid with the humerus during the definitive procedure. Similarly, an anterior midline approach is not the preferred approach for knee tumours. We prefer either a medial or lateral approach as we usually preserve the rectus femoris (Fig. 2). The recommended sites are given in the table 1 (Fig. 3, 4, 5). We wish to point out that these recommendations are based on the currently used limb salvage procedures by us. The final decision is of course that of the surgeon going to do the final procedure.

What part of the tumor should be biopsied?

Sarcomas grow centripetally and therefore have viable cells in the periphery. The centres of many rapidly growing tumors are necrotic. The best

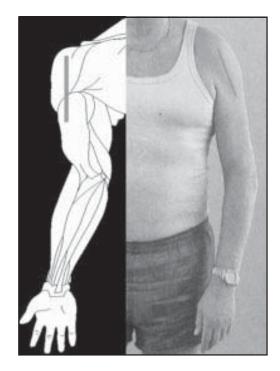


Fig. 1 : Correct site for biopsy from the proximal humerus. It should be through the anterior deltoid so that the hematoma can be contained by the muscle. The standard deltopectoral site would allow tracking of hematoma and tumor cells along loose areolar planes.

Clavicle	Through an incision that is parallel to the long axis of the clavicle
Scapula	An oblique incision which passes from superior lateral over the acromian to distal medial near the level of the inferior corner of the scapula. (Fig. 3)
Proximal Humerus Distal Humerus	Through anterior deltoid (Fig. 1) Posterior arm in midline
Pelvis (Fig. 4)	Along a line passing from the pubic tubercle to the anterior superior iliac spine over the iliac crest and to the posterior superior iliac spine depending upon the location of the lesion. Another incision line from ASIS to greater trochanter. Never through the buttock.
Hip & proximal femur	Through the lateral approach. Do not cross IM septum.
Distal femur (Fig. 2)	Anterior, medial or lateral and proximal to the suprapatellar pouch. Never through the rectus femoris Do not enter the joint
Proximal tibia (Fig. 5)	Anteromedial or anterolateral aspect. Do not enter the joint Never midline exposing through the patellar tendon.

Table 1 : Correct Biopsy Site.

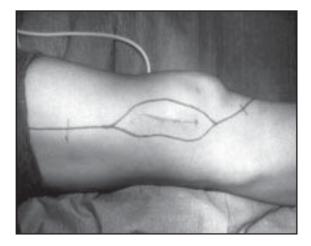


Fig. 2 : The correct site for biopsy of distal femur tumor is medial or lateral depending on where the tumor has maximum extent. The biopsy scar is excised enbloc with the tumor.



Fig. 3 : The incision for scapulectomy extends from angle of acromion laterally to inferior pole of scapula medially. The site for biopsy should be along this line.

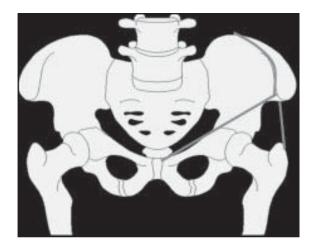


Fig. 4 : General incision for pelvic resections : line joining pubic symphysis to anterior superior iliac spine(ASIS) and then along iliac crest to posterior superior iliac spine. The second part of the incision begins just behind the ASIS and moves along the lateral thigh towards the greater trochanter and moves along the gluteal crease upto ischial tuberosity. The biopsy should be along these lines.

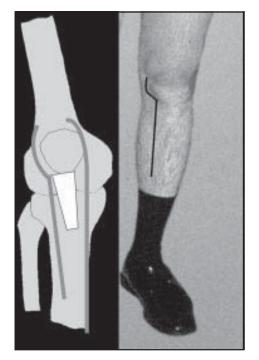


Fig. 5 : Incision for the proximal tibia is anteromedial or anterolateral, a safe distance from the margin of the patellar tendon. Utmost care should be taken to prevent contamination of the patellar tendon.

material for diagnosis is therefore from the periphery. Biopsy from the soft tissue component of a bone tumor is as representative as that from the bone and does not risk a fracture. Obviously ossified or calcified tissue should be avoided as these areas are paucicellular. Lytic areas provide the most representative tissue. Image guidance with a c-arm or CT scan is often useful when imaging points to the most suitable region to biopsy. This is most important for a needle biopsy where the maximum number of errors are due to wrong targeting.

Needle biopsy or Open biopsy ?

Traditionally an open biopsy has been used. The tissue is obtained by an operative procedure which involves an incision into the tumor. The material obtained is generally adequate in quantity and less challenging to the skills of the pathologist. The matrix, cells as well as the architecture of cells can be studied and if necessary special tests can be done. The errors in diagnosis are therefore less.

An open biopsy has to be scheduled as any other operative procedure. General anaesthesia is usually required. It is therefore a more expensive affair than a needle biopsy which is done as a percutaneous outpatient procedure. An open biopsy is also a more traumatic procedure. It involves greater tissue trauma, more blood loss and a higher risk of complications such as hematoma, infection and pathologic fracture. If a tourniquet is used there is always a fear that the oozing from tumour vessels after the tourniquet is released may contaminate large areas of the limb. All this makes an open biopsy a less forgiving procedure and a correct technique is of utmost importance if limb salvage is considered. The skin removed at final procedure is more and may compromise closure during salvage surgery.

In contradistinction to fine-needle aspiration biopsy, large gauge percutaneous cutting-needle or core-needle biopsy yields solid specimens that are amenable to histologic analysis. Multiple samples can be obtained from the same puncture site by slightly changing the angle of approach (Fig. 6). Core biopsy is especially helpful in difficult areas, such as the spine, pelvis and hips. Here it can be done image intensifier guided or CT guided.

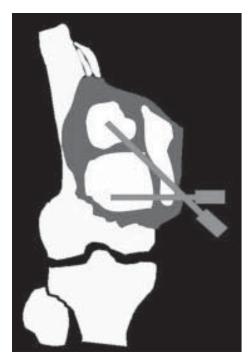


Fig. 6 : Through a single entry site, the biopsy needle should be directed in multiple directions inorder to sample different areas within the tumor. This reduces sampling error in heterogenous sarcomas.

Percutaneous biopsy of bone offers several advantages compared with open procedures. The methods are simple and economical; the biopsy can be done as an outpatient procedure, saving the time and extent of hospitalization. With the current impetus towards cost containment, core needle biopsy offers major savings compared with open biopsy. Healing of a wound is not endangered and thus, treatment with radiation and chemotherapy can be started appropriately. The opportunities of a limb salvage are improved, since there is less disruption of soft tissue and fascial planes. There is less risk of increased stress risers and pathological fractures. The rate of complications has been less than 1 percent in most series⁵⁻⁷. Needle biopsies can reach deep areas of the skeleton that are otherwise accessible only by open operation and multiple specimens can be obtained without increasing morbidity. When a needle biopsy is non-diagnostic, it can easily be repeated, or an open biopsy can be performed without major morbidity to the patient. It does however require a skilled and experienced pathologist to reach

an accurate diagnosis from the small sample of tissue. This may be possible only in specialized centres. The error rates are higher than with an open biopsy.

What is the accuracy of a needle biopsy?

The diagnostic accuracy for trucut biopsy for soft tissue tumours approaches 96 percent.^{6,7} The diagnostic accuracy reported in literature for closed bone biopsy of bone tumors is about 80 percent.² In our experience⁸, histopathological diagnosis was obtained in 89% patients. The specimen was nonrepresentative in 11% patients. 96.9% patients who had a confirmed final diagnosis had an accurate diagnosis from the J needle biopsy.

What needle to use ?

To biopsy a soft mass we use the tru-cut needle-biopsy system consisting of a cannulated needle with an inner trocar that contains a specimen notch. We use a spring loaded system to reduce crushing artifacts and pain. The trucut needle mounts onto a spring loaded gun.(Fig. 7) Alternatively, disposable single use spring loaded guns for core biopsy are commercially available. (Fig. 8) For obtaining cores from bone we use the Jamshidi needle which consists of an outer cannula and an inner trocar. (Fig. 9) A stylet to remove the cores from the cannula completes the set. The needle is sharp enough to penetrate metaphyseal bone. For the spine a coaxial bone needle like the cook's needle is used. Generally 2-3 cores are adequate.



Fig. 7 : The biopsy gun used by us at TMH. The trigger(t) releases the spring(s) which drives the trucut needle(N) into the tumor to yield a core.



Fig. 8 : Commercially available disposable spring-loaded disposable trucut biopsy needles.

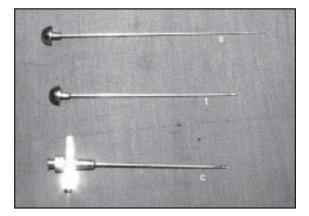


Fig. 9 : The Jamshedi Needle for bone biopsies. c : Cannula; t : Trocar; s : Stilllete.

Technique for needle biopsy

A trucut biopsy is done after infiltrating skin with local anaesthetic. Too much infiltration can increase the area of contamination. The needle is advanced till it enters the tumor. The trigger is released and the needle withdrawn to obtain the core.

For bone the Jamshidi needle is advanced through the stab incision until the trocar touches the bone. With rotatory motion the outer cortex is pierced and the trocar is withdrawn. The cannula is further introduced into the bone and rotated to core out the tissue. The cannula is withdrawn and the core removed from the cannula using a stylet. The trocar is replaced in the cannula and the procedure repeated from the same puncture site by slightly changing the angle of approach. The needle should penetrate different areas of the tumor from the same point of entry by changing the angle of approach. Sarcomas are often heterogenous; some areas appear low grade whereas some may be very high grade. Often necrotic areas are present and there may be areas of dedifferentiation. Sampling multiple areas can thus reduce the errors. After adequate cores are obtained (we usually take three), the wound is closed using a single skin suture. In general, the procedure is well tolerated with local anaesthesia. In our experience, in no patient did we fail to complete the procedure for any reason.

Principles of open biopsy technique

During an incisional biopsy attention to technical details is important for high specimen quality and reduced tumor spread. It is emphasized that the biopsy is a very important step in the evaluation of musculoskeletal lesions and that it requires careful planning and execution as in any surgery. Biopsy is not to be regarded as a simple minor procedure. Close attention to asepsis, skin preparation, hemostasis, wound closure is necessary to minimize complications.

1. Place the skin incision in such a manner so as not to compromise a subsequent definitive surgical procedure. (Avoid transverse incisions!) The incision should be in line with the planned final surgical incision and should be as small as compatible with the obtaining of an adequate tissue specimen.

2. No flaps should be raised. One should cut directly into the tumor. This reduces contamination and minimizes tissue loss at final surgery.

3. The periphery of any malignant tumor is its most viable, representative and diagnostic portion, whereas the central portion is often necrotic. The area of the Codman triangle should be avoided because of the risk that the reactive bone will be interpreted as an osteosarcoma.2 It is not necessary to biopsy the bone containing a malignant bone tumor unless there is no soft tissue extension. Violating the cortex of a bone that contains malignant tumour may lead to pathological fracture. If the bone must be opened a small circular hole should be made with

a trephine, so that only minimum stress-risers are created.23

4. Meticulous hemostasis is necessary so that substantial post-operative hematoma is prevented. 2,4,8 If a hole has been created in the bone, it should be plugged with Gelfoam or methylmethaacrylate to prevent bleeding into the soft tissues. 2,8

5. Be certain that an adequate amount of representative tissue is obtained and that the pathologist prepares the slides in a manner that will allow a definitive diagnosis.

6. Biopsy site must be closed carefully to prevent necrosis.

7. Suction drains should not be used if malignant disease is likely, as the drainage tube tract can be a site for tumor spread and will have to be excised en bloc with the biopsy site. If a drain must be used, the tract should be adjacent to and in line with the biopsy incision. Drain exiting away from incision means more contaminated skin and more excision at time of resection. (Fig. 10).



Fig. 10 : example of a badly place drain site (white arrow). It should have been ideally situated at wound ends marked a and b.

The Pathology report ... read between the lines

1. Ensure that the material was adequate and representative.

2. No diagnosis does not always mean a failed biopsy. In lesions like UBC it can rule out other differential diagnoses.

3. In a needle biopsy the diagnosis is not often "confident" or complete. If the diagnosis is consistent with the clinico-radiologic picture then treatment can be started immediately. *e.g.*, a diagnosis of a sarcoma without clear osteoid is consistent with a diagnosis of an osteosarcoma in a 15 year old child with a classical xray picture.

4. If the pathologist cannot make a diagnosis because of unfamiliarity with bone and soft-tissue tumors, urge him/her to seek consultation promptly.

5. If the diagnosis does not match the clinicoradiologic picture then the pathologist should be asked to review the diagnosis. If necessary one can repeat the biopsy or do an open biopsy.

6. Aadditional tests such as immunohisto chemistry, electron microscopy and molecular cytogenetics can help diagnosis in some cases. For example these tests can help differentiate between lymphoma and Ewings sarcoma(both are small round cell tumors)

Some more rules....

1. If the orthopaedist or the institution is not equipped to perform accurate diagnostic studies or definitive surgery and adjunctive treatment, the patient should be referred to a treating center prior to performance of the biopsy.

2. All the material collected at biopsy should be processed at one place. *It is a bad idea to divide the material and send it to different pathology laboratories.* Due to the heterogenous nature of many sarcomas this can lead to different reports and more confusion. For any discrepancy, it is better to seek a second opinion on the same material by sending slides and blocks.

3. The pathologist should be provided with all the clinical and imaging information. Without this background the diagnosis can be seriously wrong.

4. If the needle biopsy yields only fluid or blood an FNAC can be done to establish the diagnosis.

5. Beware of heterogenous or dedifferentiated tumors. A needle biopsy may sample only one area

and give an incorrect diagnosis. The dedifferentiated component may not be sampled or only a low grade area may be sampled from a tumor which is otherwise high grade.

Summary

The biopsy should be done only after appropriate and adequate imaging. It should be done at the centre by the team which will perform the final procedure. A Needle biopsy is safer but requires a skilled pathologist for interpretation. A clinicoradiologic correlation is necessary for final pathologic diagnosis and calls for co-ordination between the surgeon, radiologist and the pathologist. Additional diagnostic tools like immunohistochemistry and cytogenetics can aid differential diagnosis. The surgical principles and technique should be meticulously adhered to failing which an unnecessary amputation may result.

Before doing a biopsy think. After reading a biopsy report, think again.

References

1. Mankin, Henry J., Mankin, Carole J., Simon, Michael A.: The Hazards of the Biopsy, Revisited. J Bone Joint Surg [Am] 1996; 78-A; 656-63

2. Jaffe HL: Introduction: Problems of Classification and Diagnosis. In Jaffe HL (ed). Tumors and Tumourous Conditions of the Bones and Joints. Philadelphia, Lea and Febiger 9-17, 1958.

3. Enneking WF: The issue of the biopsy [editorial]. J Bone Joint Surg 64-A: 1119-1120 1982.

4. Simon MA: Current concepts review. Biopsy of musculoskeletal tumors. J Bone Joint Surg 64-A: 1253-1257,1982

5. Barth RJ Jr., Merino MJ, Solomon D, Yang JC and Baker AR: A prospective study of the value of core needle biopsy and fine needle aspiration in the diagnosis of soft tissue masses. Surgery 112: 536-543, 1992

6. Ball ABS, Fisher C, Pittam M, Watkins RM and Westburg G: Diagnosis of soft tissue tumours by Tru-cut biopsy. British J Surg 77: 756-758, 1990

7. Kissin MW, Fisher C, Carter RL, Horton LWL and Westburg G: Value of Tru-Cut Biopsy in the diagnosis of soft tissue tumours. British J Surg 73: 742-744, 1986

8. Pramesh CS, Deshpande MS, Pardiwala DN, Agarwal MG, Puri A.: Core needle biopsy for bone tumours. Eur J Surg Oncol. 2001 Nov;27(7):668-71

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