



Detection of Bone Metastasis On Whole Body Bone Scan Among Sudanese Patients with Proven Cancer in Tumours Therapy and Cancer Research Center –Shendi –Sudan

Dr. Motwakil Imam Awad Elkareim Imam^{1*}, Taha Ali Mohammed Taha², Abdalla Atta Abdalla Abukleawa² and Mahmoud Babekir Hassan Mohammed²

^{1*}Associate Professor of Internal Medicine, Faculty of Medicine, Shendi University, Consultant physician, Elmek Nimer University Hospital

²Medical students, Faculty of Medicine, Shendi University, Sudan

Abstract: Bony metastasis is one of the most common causes of death in patients with proven cancer, most of the patients present by pathological fracture, and carry poor prognosis. The bone scan is the main investigation for diagnosis. Our objective is to determine the most common cancer that causes metastasis, the commonest site involved in the skeleton, and the metastatic pattern for each cancer. We studied 150 cases collected in tumor therapy and cancer research centers, those who underwent bone scans using a gamma camera. The most age of our patients ranged between (40 and 60) years old. The predominance of females was noticed (68.7%). 90% of the patients were married. The study showed that 35.3% with left breast cancer, 20% with right breast cancer, and 22% with prostate cancer. The study showed that 40% of the participants had bone metastasis, 41.7% with prostate cancer, 21.7% with left breast cancer, and 16.7% with right breast cancer. The study showed that the lumbar vertebra involved in 51.7% of bone metastasis, thoracic vertebra involved in (50%) with bone metastasis, the anterior ribs involved in 28.3% with bone metastasis, while posterior ribs in 25%, the right femur involved in 28.3 while the left femur engaged in 25% of bone metastasis and 25% metastasis to the skull. About 40% of the patients had bony metastasis. Breast and prostate are the most common cancers that cause bony metastasis. The spine is the most common site for metastasis. Prostate cancer metastasized mainly to the spine followed by the femur, ribs, skull, and shoulders

Keywords: bony metastasis, bone scan, cancer patient, Shendi –Sudan

*Corresponding Author

Dr. Motwakil Imam Awad Elkareim Imam , Associate Professor of Internal Medicine, Faculty of Medicine, Shendi University, Consultant physician, Elmek Nimer University Hospital

Received On 4 November 2024

Revised On 28 November 2024

Accepted On 12 December 2024

Published On 6 January 2025

Funding This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Citation Dr. Motwakil Imam Awad Elkareim Imam, Taha Ali Mohammed Taha, Abdalla Atta Abdalla Abukleawa , Mahmoud Babekir Hassan Mohammed , Detection of bone metastasis on whole body bone scan among Sudanese patients with proven cancer in Tumours Therapy and Cancer Research Center –Shendi -Sudan.(2025).Int. J. Trends in OncoSci.3(1), 21-39

This article is under the CC BY- NC-ND Licence (<https://creativecommons.org/licenses/by-nc-nd/4.0>)

Copyright @ International Journal of trends in OncoScience, available at www.ijtos.com

Int. J. Trends in OncoSci., Volume3., No 1 (January) 2025, pp 21-39



1. INTRODUCTION

Bone metastases are a frequent complication of cancer, occurring in up to 70% of patients with advanced breast or prostate cancer and approximately 15% to 30% of patients with carcinoma of the lung, colon, stomach, bladder, uterus, rectum, thyroid, or kidney. The exact incidence of bone metastasis is unknown, but it is estimated that 350,000 people die from bone metastases annually in the United States.¹⁻⁵ Once tumors metastasize to bone, they are usually incurable; only 20% of patients with breast cancer are still alive five years after the discovery of bone metastasis. The consequences of bone metastasis are often devastating. Osteolytic metastases can cause severe pain, pathologic fractures, life-threatening hypercalcemia, spinal cord compression, and other nerve-compression syndromes. Patients with osteoblastic metastases have bone pain and pathologic fractures because of the poor quality of bone produced by the osteoblasts. For all these reasons, bone metastasis is a serious and costly complication of cancer.⁶⁻¹⁵ Most patients with cancer do not die from primary cancer, but from secondary metastatic disease. The bone is the most common site of secondary metastases, secondary to the lung and liver. Bone metastasis is a malignant tumor of the extraosseous organ or tissue. These metastatic cells are transferred by the lymphatic blood system to the bone, which then continues to grow and form a tumor. According to the American Cancer Society, about 400,000 new cases of malignant bone metastasis are diagnosed in the United States each year. The incidence of advanced malignant tumors with bone metastasis is 30–75%, prevalent in patients with advanced prostate cancer and breast cancer. Bone metastases often cause limb dysfunction, pathological fractures, spinal cord compression, and severe pain, seriously affecting the quality of life of patients with advanced cancer and poor prognoses.¹⁶⁻²⁶ Despite the influence of bone metastases on cancer patients, large-scale research studies examining the incidence, prevalence, and outcomes of patients with bone metastases remain lacking.^{1, 20- 28} The skeleton of the human being is a unique structure that has adapted to the needs of bipedal locomotion and upright posture. The structural peculiarities of the human skeleton give human beings their characteristic appearance and facial geometry. The bony skeleton provides the shape and framework on which the human body is designed and functions. It houses and protects vital organs, contains bone marrow, which is the functional unit of the hematopoietic system; and it provides attachments and anchorage to muscles, ligaments, and joint capsules. Bones often act as levers, which, in conjunction with muscular contraction, initiate and sustain movement.^{2,29-35} When solid tumors metastasize to the skeleton, they cause a variety of alterations in bone cell function that may be represented as discrete osteolysis, diffuse osteopenia, osteoblastic lesions, or a combination of all of the above. All these effects are caused by the effects of tumor products on the normal bone remodeling sequence. The most common of these lesions is the destructive or osteolytic lesion. Osteolysis is characterized by a marked increase in osteoclast formation and osteoclast activity caused by tumor products. There may be a subsequent osteoblastic response, but this is often blunted and sometimes absent. Less commonly, solid tumors cause an increase in osteoblast activity. This may occur without obvious previous desorption; although it may also be associated with prior desorption at the same site, and the formation phase may be relatively exaggerated.^{7-9,36-40} Bone imaging continues to be the second-greatest-volume nuclear imaging procedure, offering the advantages of total body examination, low cost, and high

sensitivity. Its strength rests in the physiological uptake and path physiologic behavior of 99m technetium (99m-Tc) diphosphonates. The diagnostic utility, sensitivity, specificity, and predictive value of 99m-Tc bone imaging for benign conditions and tumors were established when only planar imaging was available. Currently, nearly all bone scans are performed as a planar study (whole-body, 3-phase, or regional), with the radiologist often adding single-photon emission computed tomography (SPECT) imaging. Here we review many current indications for planar bone imaging, highlighting indications in which the planar data are often diagnostically sufficient, although diagnosis may be enhanced by SPECT. ¹⁸F sodium fluoride positron emission tomography (PET) is also re-emerging as a bone agent and has been considered alternate with 99m-Tc diphosphonates in the past. In addition to SPECT, new imaging modalities, including ¹⁸F fluorodeoxyglucose, PET/CT, CT, magnetic resonance, and SPECT/CT, have been developed and can aid in evaluating benign and malignant bone disease. Because ¹⁸F fluorodeoxyglucose is taken up by tumor cells and Tc diphosphonates are taken up in osteoblastic activity or osteoblastic healing reaction, both modalities are complementary.⁴¹⁻⁴⁵ CT and magnetic resonance may supplement, but do not replace, bone imaging, which often detects pathology before anatomic changes are appreciated. We also stressed the importance of dose reduction by reducing the dose of 99m-Tc diphosphonates and avoiding unnecessary CT acquisitions. Moreover, we propose an approach to image interpretation that emphasizes communication with referring colleagues and correlation with appropriate history to significantly improve our impact on patient care.^{8, 46-51}

2. METHODOLOGY

The present study was a prospective, cross-sectional, hospital-based study conducted in tumor therapy and cancer research center - Shendi University Sudan. We used the total coverage technique as the sampling method to collect data from the total number of included participants. Thus, the total number of included participants was 150 patients. The study was conducted between (2010- 2017). The center was established in 2010 to provide chemotherapy, nuclear medicine imaging, endoscopy services, laboratory services, radio-iodine therapy, early detection services, and Radiation therapy for cervical cancer (brachytherapy). The Teletherapy department is under establishment. This is the only center providing these services for the population in the River Nile state of Sudan. The center is located in Shendi town, it is about 150 km northern to Khartoum capital of the Sudan, and about 45 km Southern to the ancient city of Merwe. The center is located about 4km from the center of Shendi town, at the cross of highway road from Khartoum to Atbara across the main road inter center of Shendi town. The inclusion criteria of the patients were: being 18 years of age and older, having proven cancer, and a whole body bone scan was done to them for detecting bone metastasis.

2.1. Data collection method & tool

Data were collected using a structured questionnaire. The questionnaire was filled directly from the medical files of the patient's records. For all the patients involved in the study whole body bone scan was done . The questionnaire included items to measure sociodemographic characteristics (gender, age, residence, and marital status .

2.2. Statistical Data Analysis

Data was reviewed, ordered, and coded, and then Statistical Package for Social Sciences (SPSS) version 20 was used for data analysis. Descriptive statistics were used to analyze the participants' data... The data is presented in the form of figures and tables .

2.3. Ethical consideration

Ethical approval for this study was obtained from the scientific research commitment of Shendi University and the Ministries of Health in North Sudan (SMSB-E.C.66.2021), in

inconsistency with Helsinki's declaration of the international conference on harmonization, regulations, and laws of Sudan.

3. RESULTS

A total of 150 patients with different types of cancer were included in the study to bone scan was done by using a gamma camera in the tumor therapy and cancer research center at Shendi University Sudan. More than half of the participants were males, more than two-thirds of them were in the age group of 40 to 60 years and most of the females were housewife workers. Nearly two-thirds of them were from Shendi town and the majority of them (90%) were married . Table I

Table I: Distribution of Participants by Gender and Residency		
gender	N	%
male	47	31.3
female	103	68.7
Resident		
Shendi	78	52.0
Al matama	11	7.3
Al mesaiktab	6	4.0
Alshagaloa	6	4.0
Al trajma	3	2.0
Kabosheah	5	3.3
Algeliia	4	2.7
Dem algrai	4	2.7
Alseal	4	2.7
Alkimair	8	5.3
Alsloab	2	1.3
Ttaibhalkhoad	3	2.0
al mahmeia	2	1.3
Al damar	6	4.0
Atbara	3	2.0
abuhamaed	1	.7
Algraef	2	1.3
Banaga	1	.7
Barbar	1	.7
Occupation		
Housewife	97	64.7
Free business	44	29.3
Employee	9	6.0
Educational level		
Not educate	75	50.0
Primary	40	26.7
Secondary	28	18.7
University	7	4.7
Marital status		
Married	135	90.0
Single	15	10.0

Table 2: type of tumor among participants		
Type of tumor	Frequency	Percent%
breast cancer	53	35.3
Ca prostate	33	22.0
Ca ovary	5	3.3
Endometrial carcinoma	3	2.0
Thyroid cancer	3	2.0
Chronic lymphocytic leukemia	1	.7
Burkett's lymphoma	1	.7
Nasopharyngeal carcinoma	1	.7
Colorectal carcinoma	3	2.0

Esophageal carcinoma	2	1.3
Rhabdomyosarcoma	1	.7
Renal cell carcinoma	3	2.0
Synovial sarcoma	1	.7
Cervical carcinoma	2	1.3
Carcinoma of the head of pancreas	1	.7
Right breast cancer	30	20.0
Bilateral breast cancer	4	2.7
Uterine leiomyosarcoma	1	.7
Myxoid chondrosarcoma	1	.7
Ca bladder	1	.7
Total	150	100.0

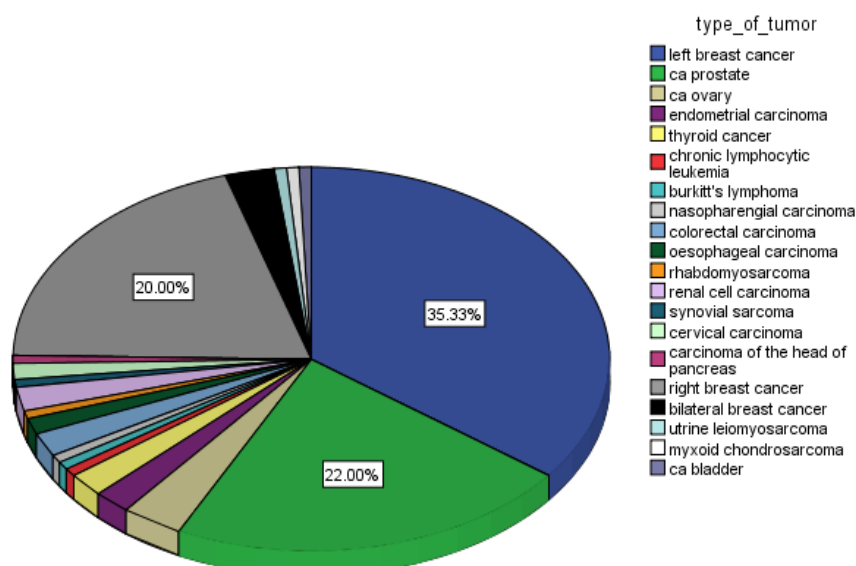


Fig 1: Distribution of study population according to their type of tumor

Table 3: Type of therapy among participants		
	Frequency	Percent%
Chemotherapy		
Yes	107	71.3
No	43	28.7
Radiotherapy		
Yes	52	34.7
No	98	65.3
Surgery		
Yes	106	70.7
No	44	29.3
Hormonal therapy		
Yes	24	16.0
No	126	84.0

Table 4: Frequency of bone metastasis among participants and duration of the disease		
Bone metastasis	Frequency	Percent%
Yes	60	40.0
No	90	60.0

Table 5: Frequency of the site of bone metastasis		
Duration from diagnosis to metastasis	Frequency	Percent%
no metastasis	90	60
one year	36	24
2 years	17	11.4
3 years	4	2.6
more than 3 years	3	2
Total	150	100.0

Table 6: Frequency and Percentage Distribution of Affected Sites

Site	Count	percent %
Skull	15	25.0%
Anterior ribs	17	28.3%
Posterior ribs	15	25.0%
Right shoulder	12	20.0%
Left shoulder	13	21.7%
Right clavicle	0	0.0%
Left clavicle	2	3.3%
Right humerus	6	10.0%
Left humerus	9	15.0%
Right scapula	1	1.7%
Cervical vertebra	10	16.7%
Left scapula	1	1.7%
Thoracic vertebra	30	50.0%
Lumber vertebra	31	51.7%
Sacral vertebra	6	10.0%
Coccyx	5	8.3%
Right sternoclavicular joint	2	3.3%
Left sternoclavicular joint	1	1.7%
Sternum	8	13.3%
Right pelvis	10	16.7%
Left pelvis	11	18.3%
Right sacroiliac	1	1.7%
Left sacroiliac	3	5.0%
Right hip joint	0	0.0%
Left hip joint	3	5.0%
Right acetabulum	0	0.0%
Left acetabulum	2	3.3%
Right femur	17	28.3%
Left femur	15	25.0%
Right knee	6	10.0%
Left knee	7	11.7%
Right tibia	4	6.7%
Left tibia	9	15.0%
Right tarsus	1	1.7%
Left tarsus	1	1.7%
Right toes	1	1.7%
Left toes	0	0.0%

Table 7: Frequency of bone metastasis according to type of cancer

Type of tumor	Frequency	Percent %
Left breast cancer	13	21.7%
Prostate cancer	25	41.7%
Ovarian cancer	0	0.0%
Endometrial carcinoma	0	0.0%
Thyroid cancer	0	0.0%
Chronic lymphocytic leukemia	0	0.0%
Burkett's lymphoma	0	0.0%
Nasopharyngeal carcinoma	1	1.7%
Colorectal carcinoma	0	0.0%
Esophageal carcinoma	1	1.7%
Rhabdomyosarcoma	1	1.7%
Renal cell carcinoma	2	3.3%
Synovial sarcoma	0	0.0%
Cervical carcinoma	1	1.7%
Carcinoma of the head of pancreas	0	0.0%
Right breast cancer	10	16.7%
Bilateral breast cancer	3	5.0%
Uterine leiomyosarcoma	1	1.7%

Myxoid chondrosarcoma	I	1.7%
Urinary bladder cancer	I	1.7%

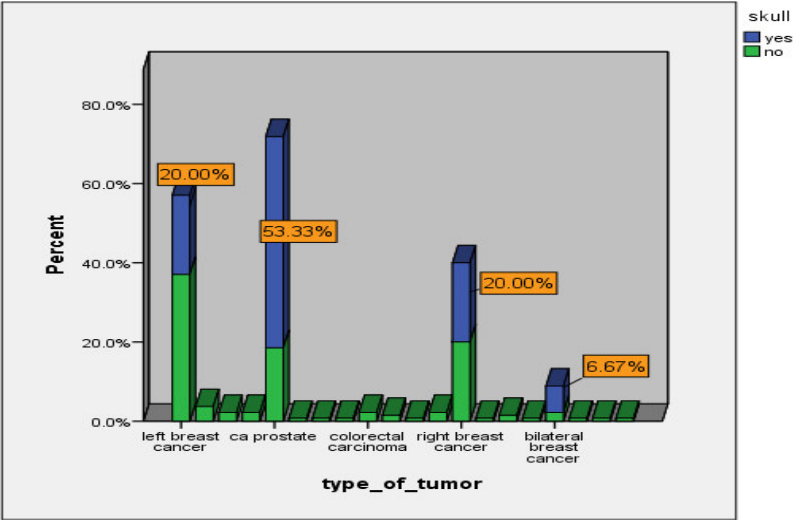


Fig 2: The skull is involved in (25%) of all bone metastases (53.33%) from prostate cancer, (20%) from right breast cancer, (20%) from left breast cancer, and (6.67%) from bilateral breast cancer

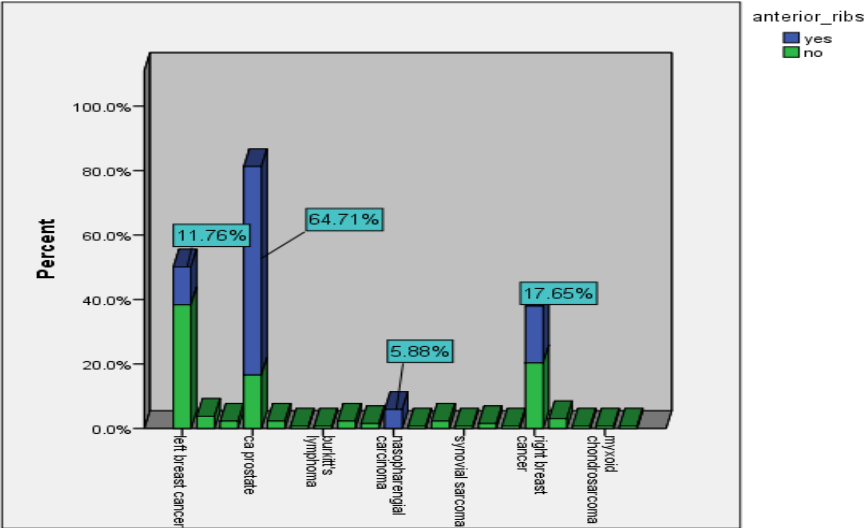


Fig 3: The anterior ribs are involved in (28.3%) of all bone metastasis (64.71%) from prostate cancer, (17.65%) from right breast cancer, (11.76%) from left breast cancer, and (5.88%) from nasopharyngeal carcinoma

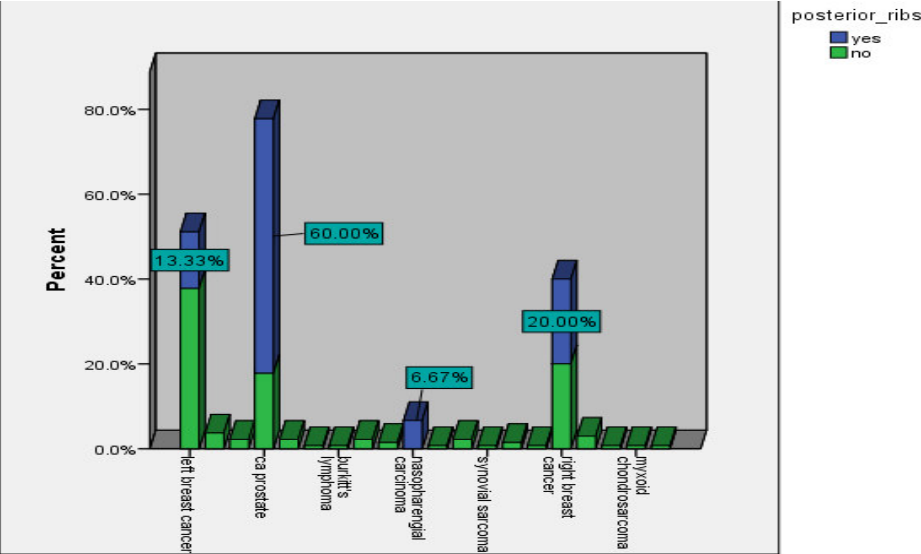


Fig 4: The posterior ribs were involved in (25%) of all bone metastasis (60%) from prostate cancer, (20%) from right breast cancer, (13.33%) from left breast cancer, and (6.67%) from nasopharyngeal carcinoma

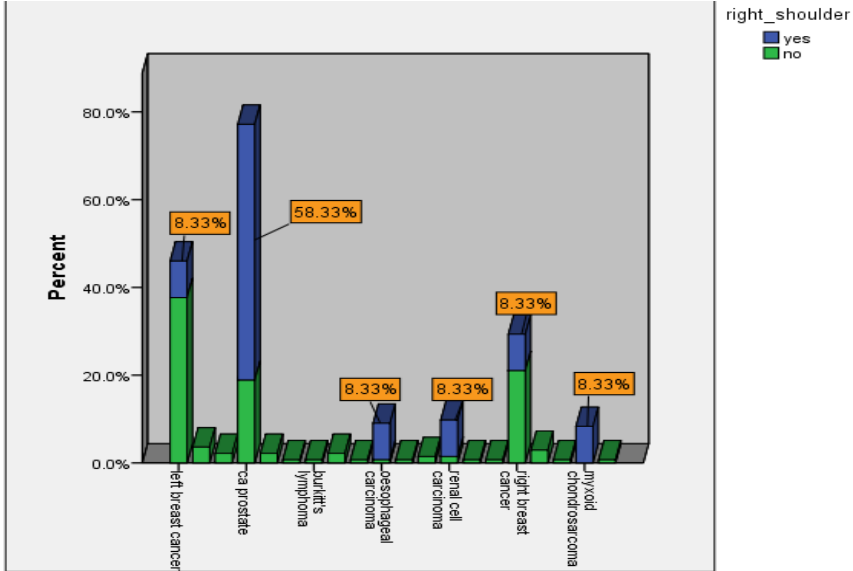


Fig 5: The right shoulder is involved in 20% of all bone metastasis (58.33%) from prostate cancer, 8.33%) from left breast cancer, 8.33%) from right breast cancer, 8.33%) from esophageal carcinoma, 8.33%) from renal cell carcinoma, and 8.33%) from myxoid chondrosarcoma

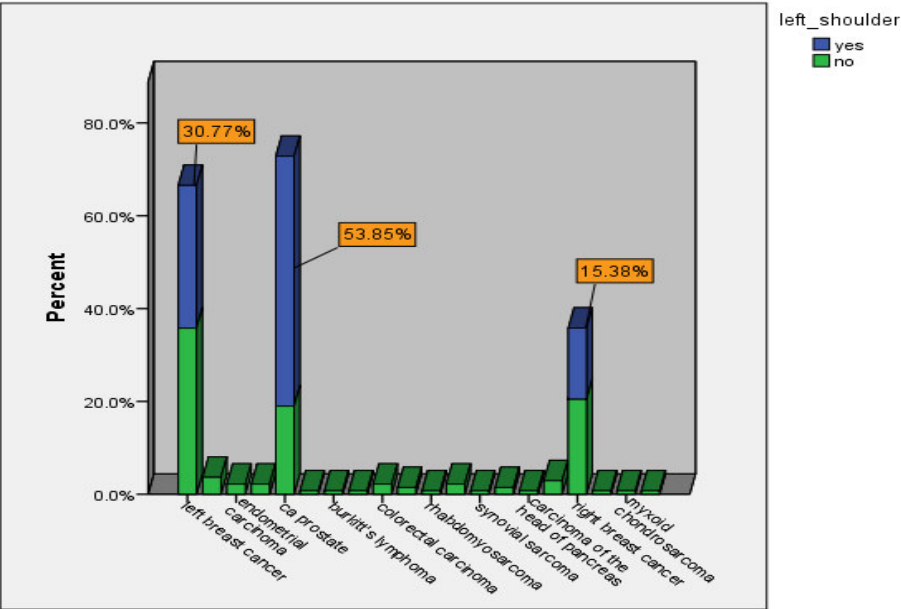


Fig 6: The left shoulder is involved in 21.7% of all bone metastasis (53.85%) from prostate cancer, 30.77%) from left breast cancer, and 15.38%) from right breast cancer

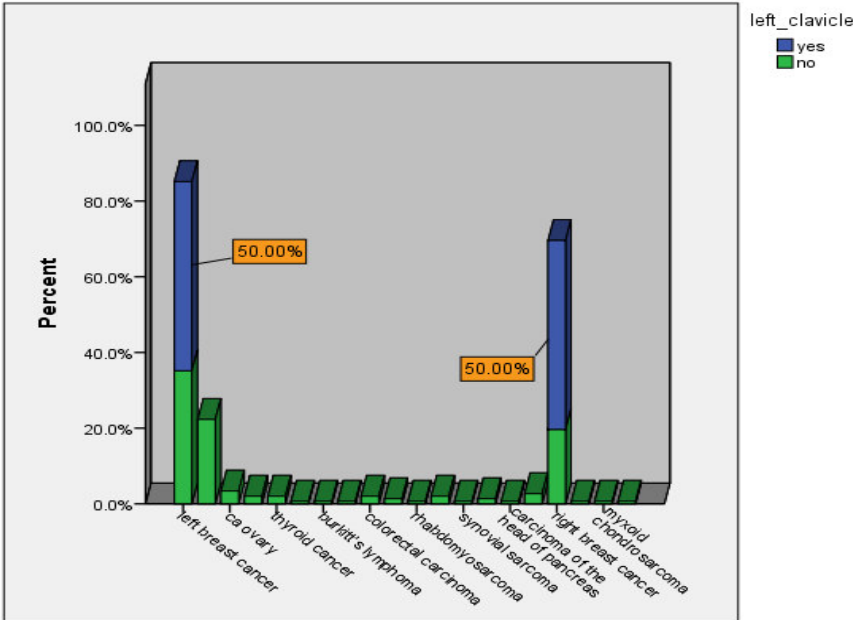


Fig 7: The left clavicle is involved in 21.7% of all bone metastases (50%) from left breast cancer and right breast cancer

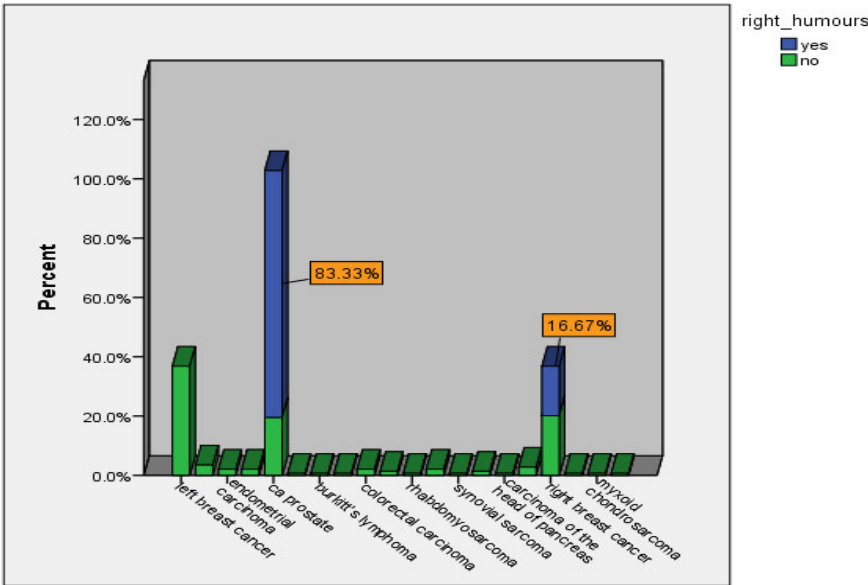
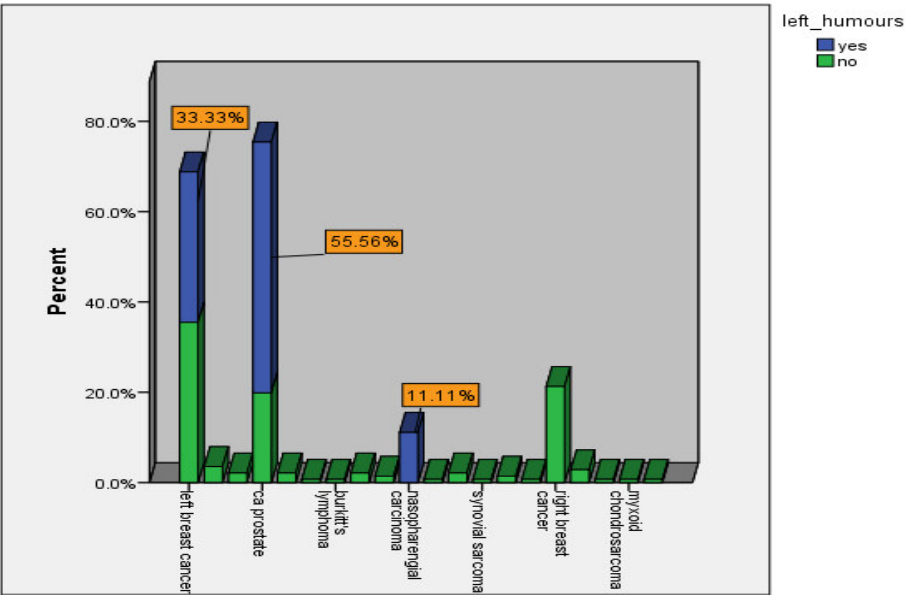


Fig 8: The left humerus was involved in 55.56% of all bone metastasis (33.33%) from left breast cancer, and 11.11% came from nasopharyngeal carcinoma

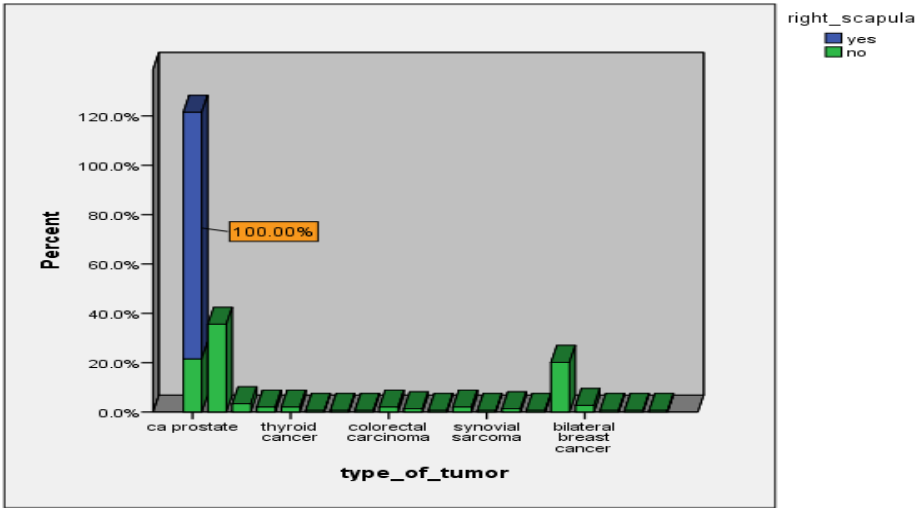


Fig 9: The right humerus is involved in (10%) of all bone metastasis (83.33%) from prostate cancer and (16.67%) from right breast cancer

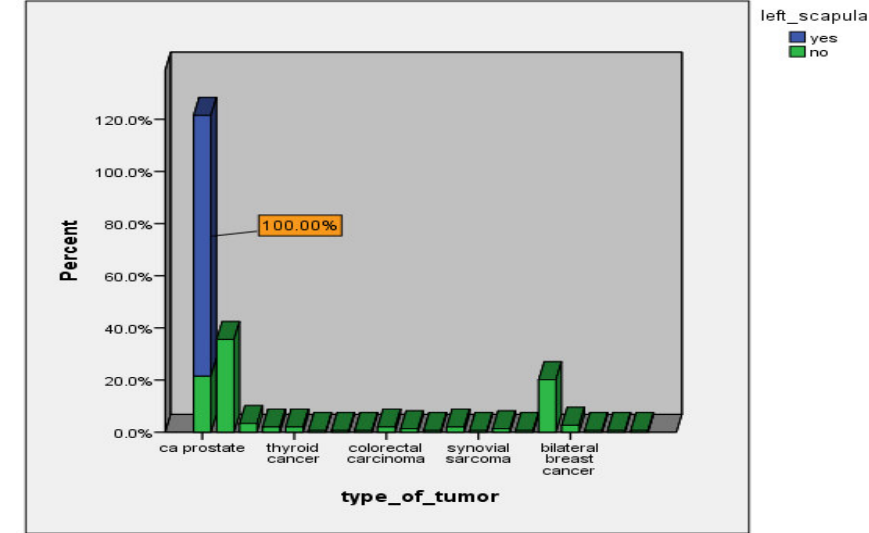


Fig 10: The right scapula is involved in (1.7%) of all bone metastasis (100%) from prostate cancer

Fig 11: The right scapula is involved in (1.7%) of all bone metastasis (100%) from prostate cancer

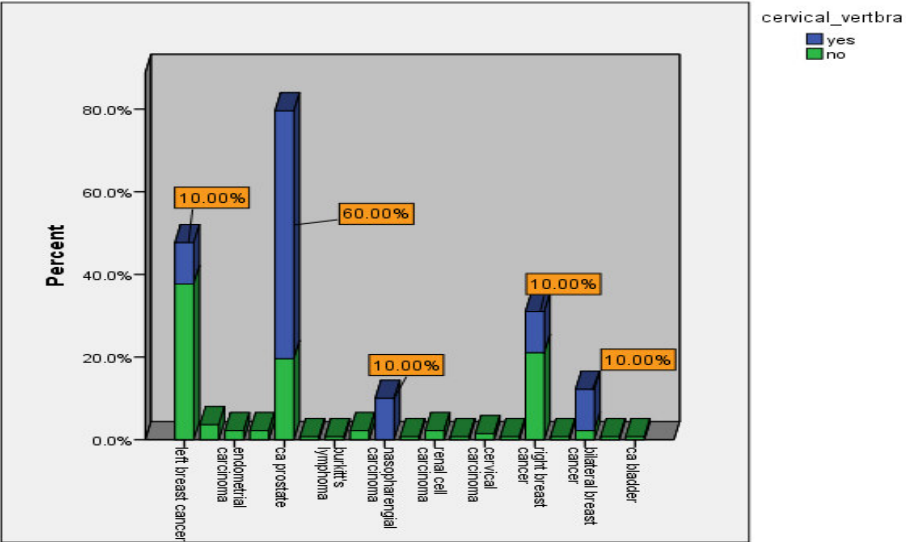


Fig 12: The cervical vertebra is involved in 16.7% of all bone metastasis (60%) of patients from prostate cancer, (10%) from left breast cancer, (10%) from right breast cancer, (10%) from bilateral breast cancer, and (10%) from nasopharyngeal carcinoma

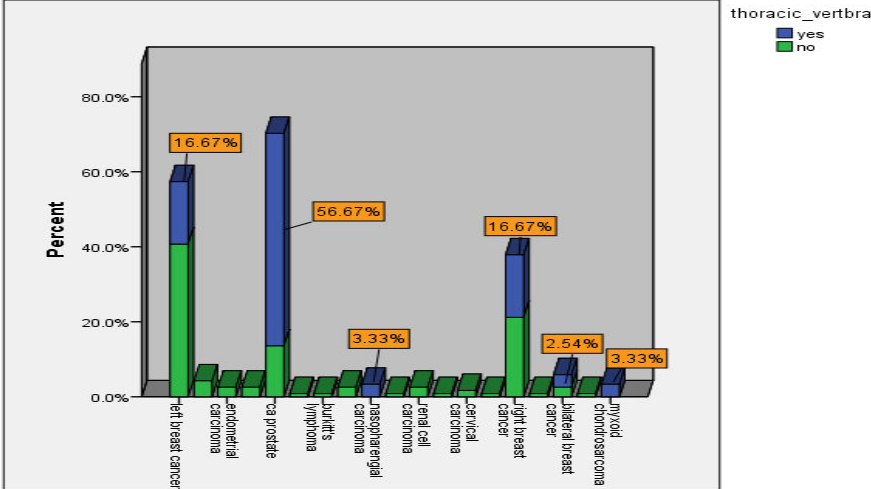


Fig 13: The thoracic vertebra is involved in (50%) of all bone metastasis (56.76%) from prostate cancer, (16.67%) from left breast cancer, (16.67%) from right breast cancer, (3.33%) nasopharyngeal carcinoma, (3.33%) from myxoid chondrosarcoma, and (2.54%) from bilateral breast cancer

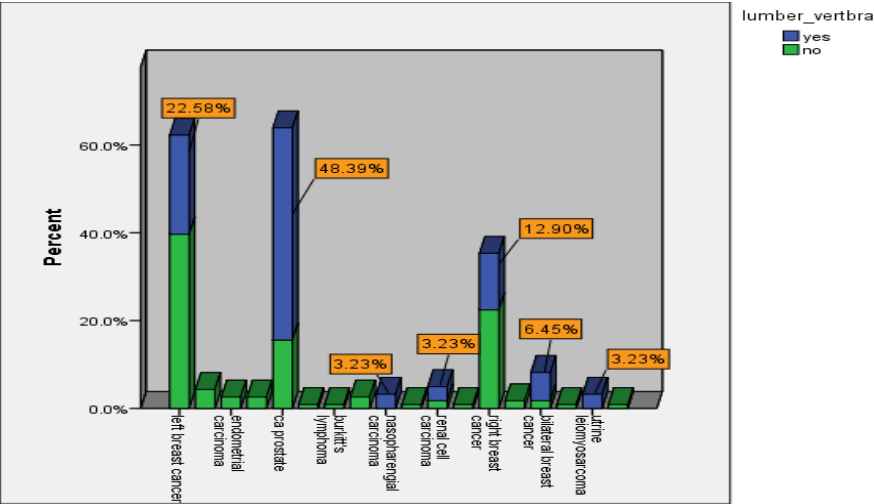


Fig 14: The lumbar vertebra is involved in 51.7% of all bone metastases (48.39%) from prostate cancer, (22.58%) from left breast cancer, 12.90% from right breast cancer, 6.45% from bilateral breast cancer, 3.23% from renal cell carcinoma, 3.23% from nasopharyngeal carcinoma, and 3.23% from uterine leiomyosarcoma .

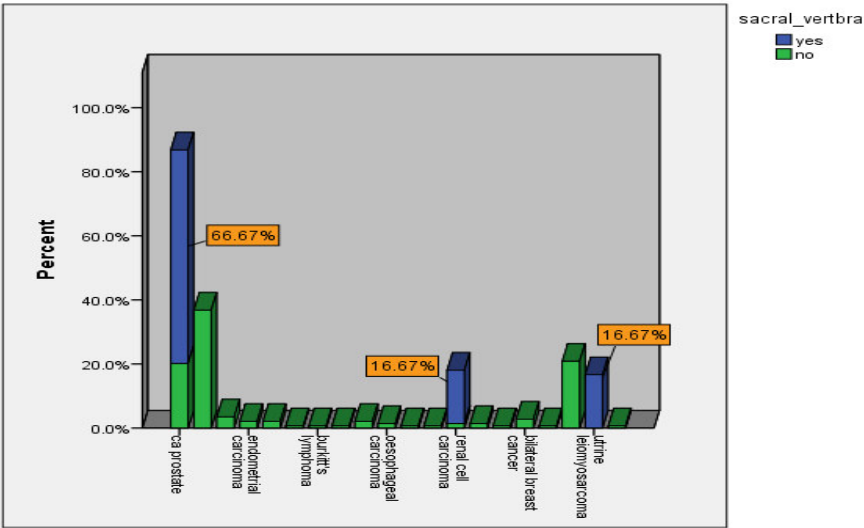


Fig15: The sacral vertebra is involved in 10% of all bone metastases (66.67%) from prostate cancer, 16.67% from renal cell carcinoma, and .16.67% from uterine leiomyosarcoma

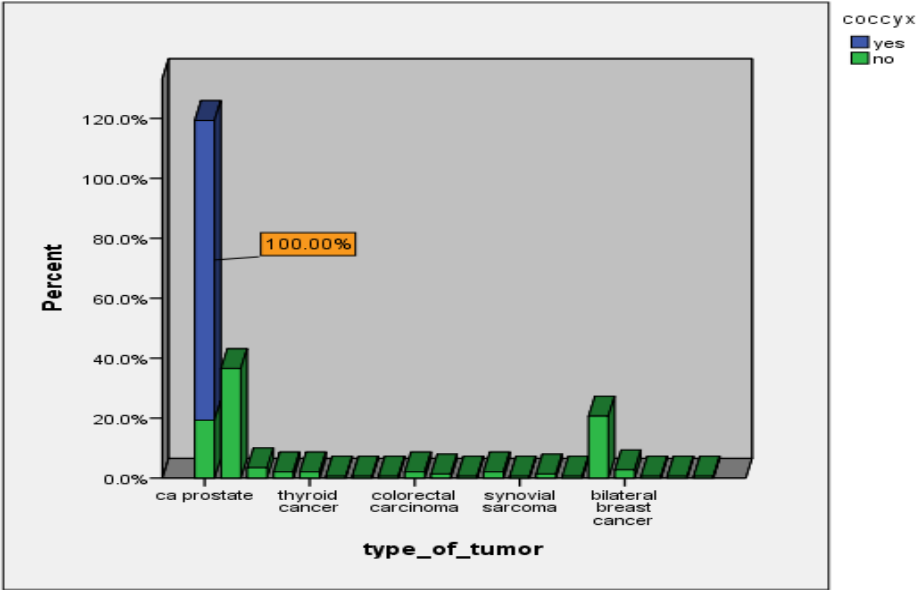


Fig 16: The coccyx is involved in 8.3% of all bone metastasis (100%) from prostate cancer .

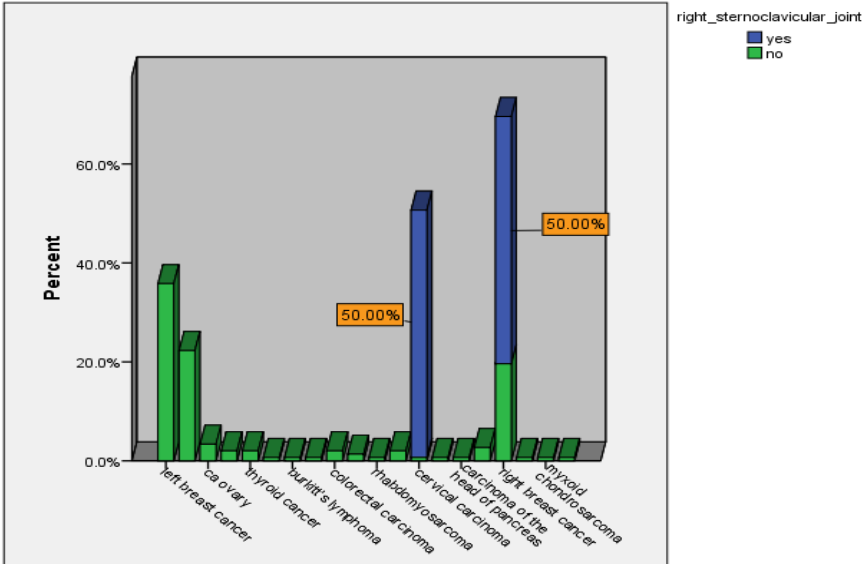


Fig 17: The right sternoclavicular joint is involved in (3.3%) of all bone metastases (50%) from cervical carcinoma and (50%) from right breast cancer

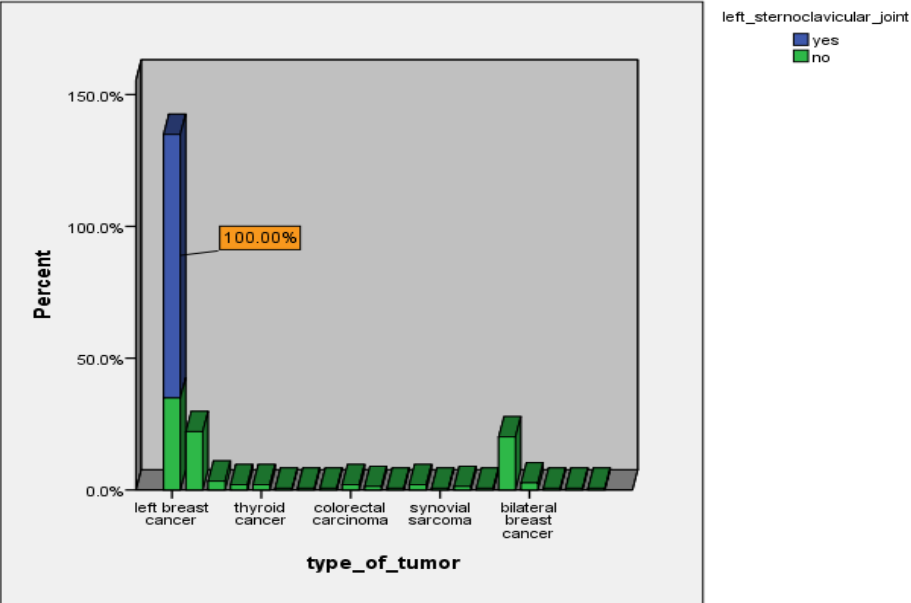


Fig 18: The left sternoclavicular joint is involved in (1.7%) of all bone metastases (100%) from left breast cancer

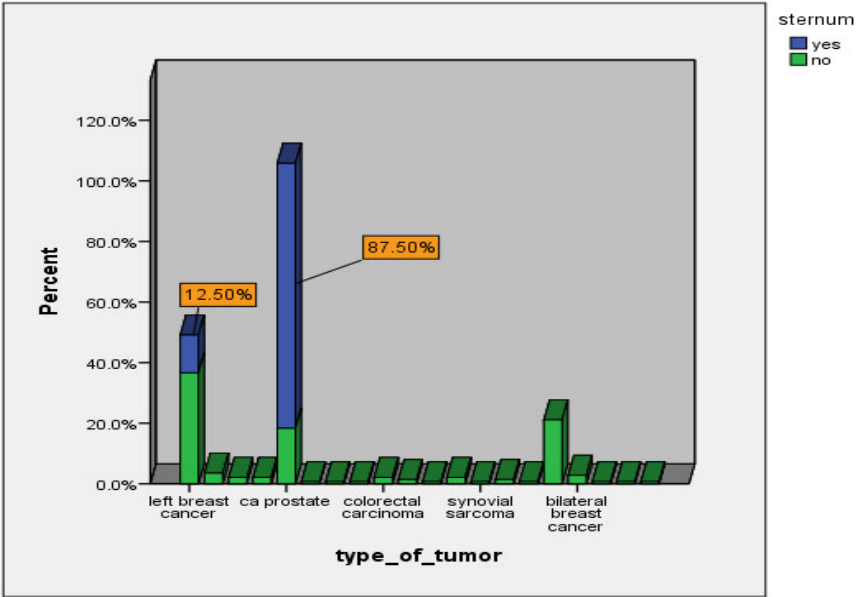


Fig 19: The sternum is involved in 13.3% of all bone metastases (87.5%) from prostate cancer and 12.50% from left breast cancer

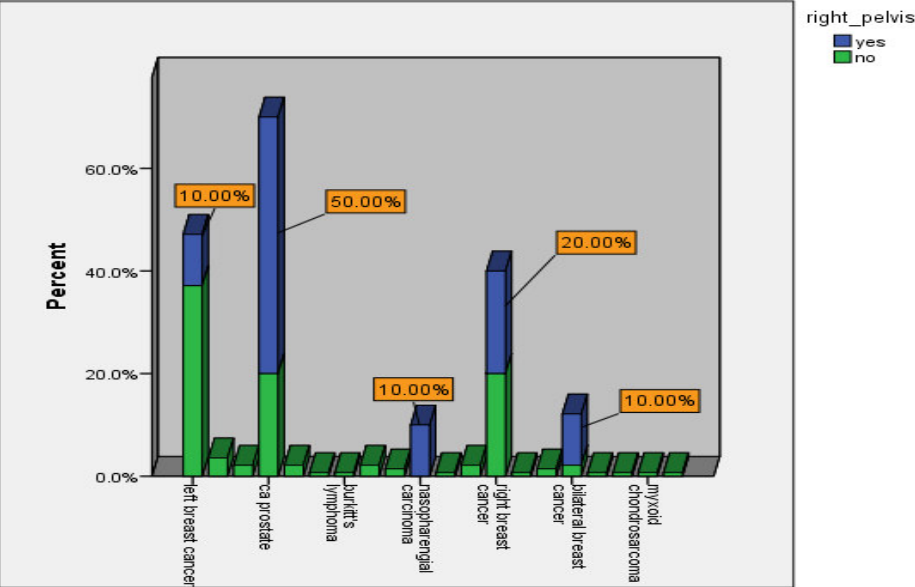


Fig 20: The right pelvis was involved in 16.7% of all bone metastasis (50%) from prostate cancer, (20%) came from right breast cancer, (10%) came from left breast cancer, (10%) from bilateral breast cancer, and (10%) from nasopharyngeal carcinoma.

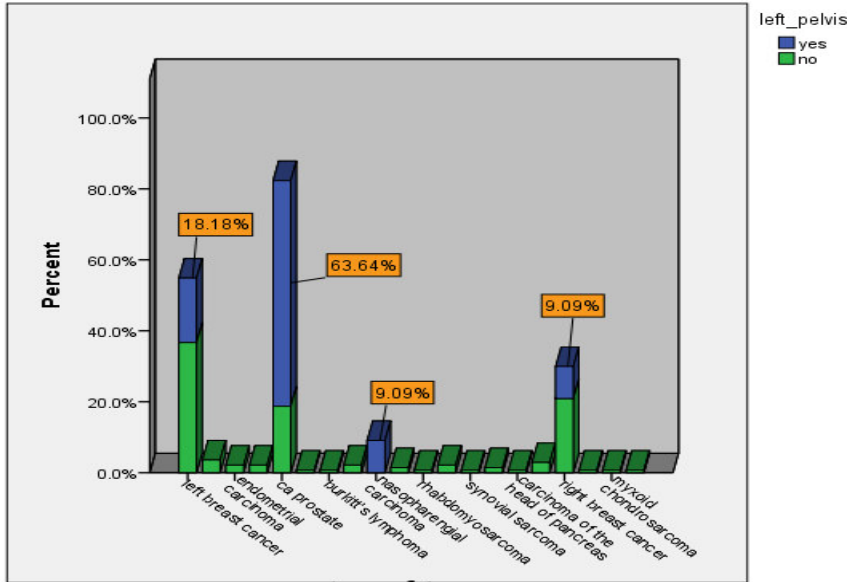


Fig 21: The left pelvis is involved in 18.3% of all bone metastasis (63.64%) from prostate cancer, 18.18% from left breast cancer, 9.09% from right breast cancer, and 9.09% from nasopharyngeal carcinoma .

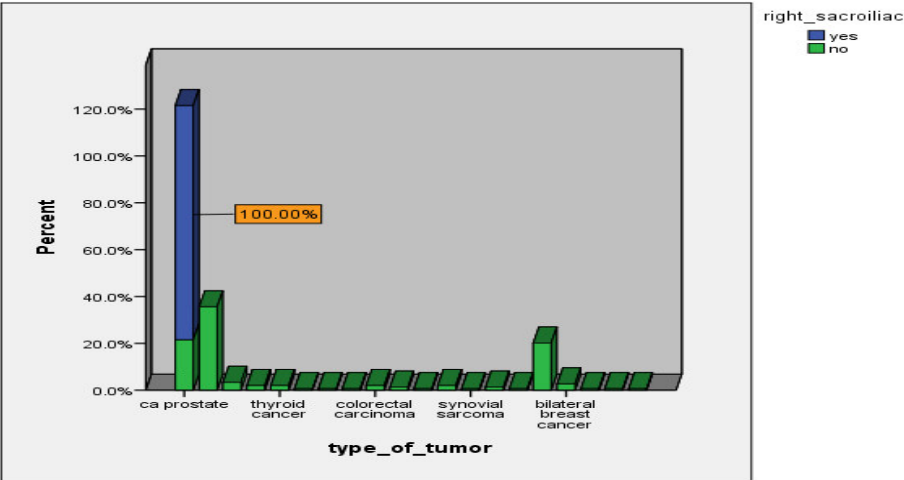


Fig 22: The right sacroiliac joint is involved in (1.7%) of all bone metastasis .(100%) from prostate cancer

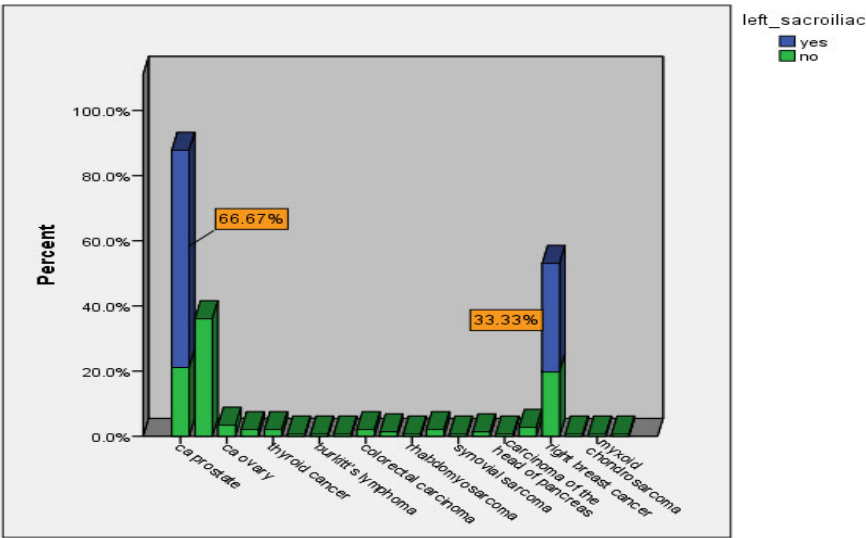


Fig 23: The left sacroiliac joint is involved in (5%) of all bone metastasis (66.67%) from prostate cancer and (33.33%) from right breast carcinoma

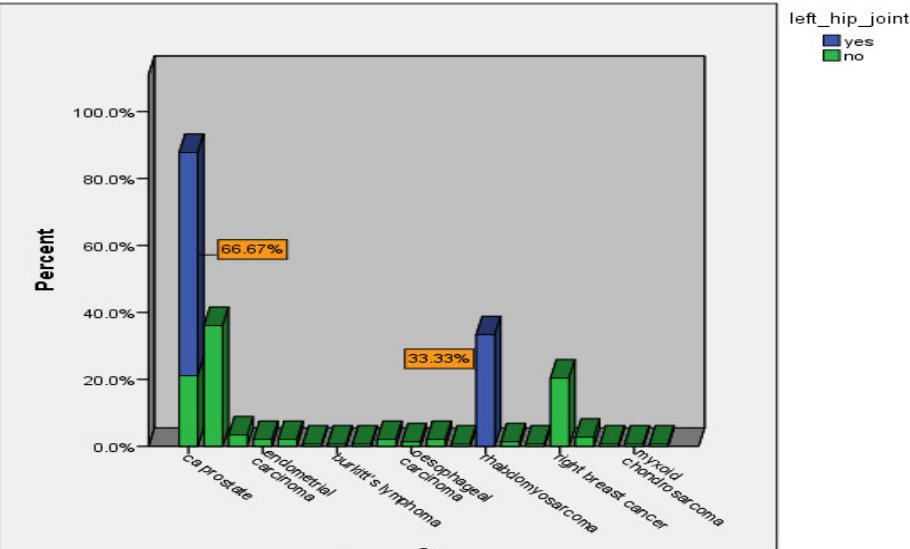


Fig 24: the left hip joint is involved in (5%) of all bone metastasis (66.67%) from prostate cancer and (33.33%) from rhabdomyosarcoma

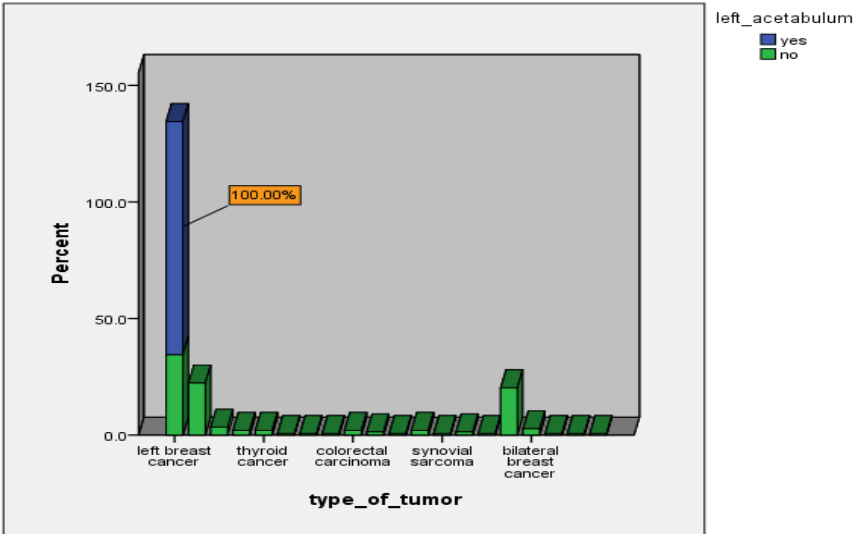


Fig 25: The left acetabulum is involved in (3.3%) of all bone metastasis (100%) from left breast cancer in both cases

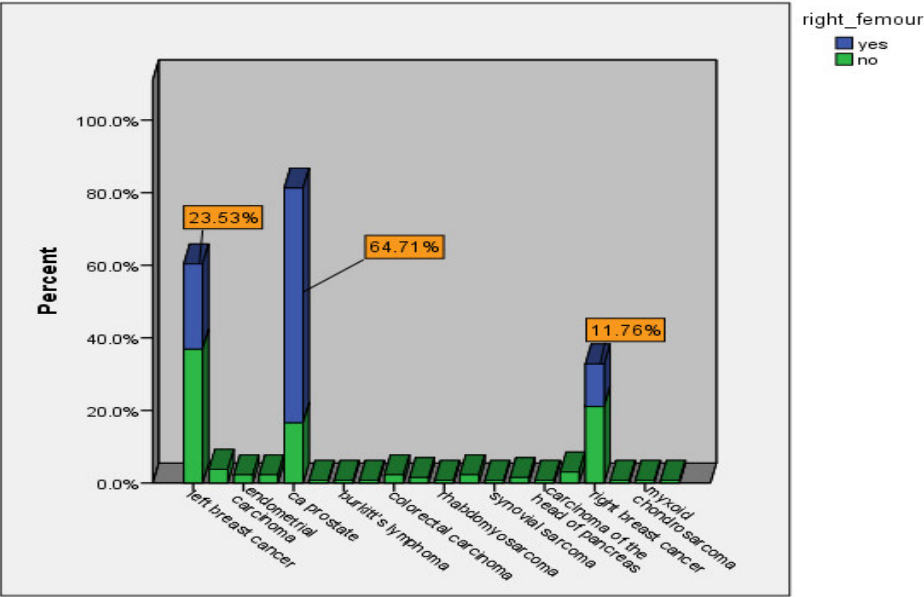


Fig 26: The right femur is involved in (28.3%) of all bone metastases (64.71%) of patient's metastases from prostate cancer, (23.53%) from left breast cancer, and (11.76%) from right breast cancer

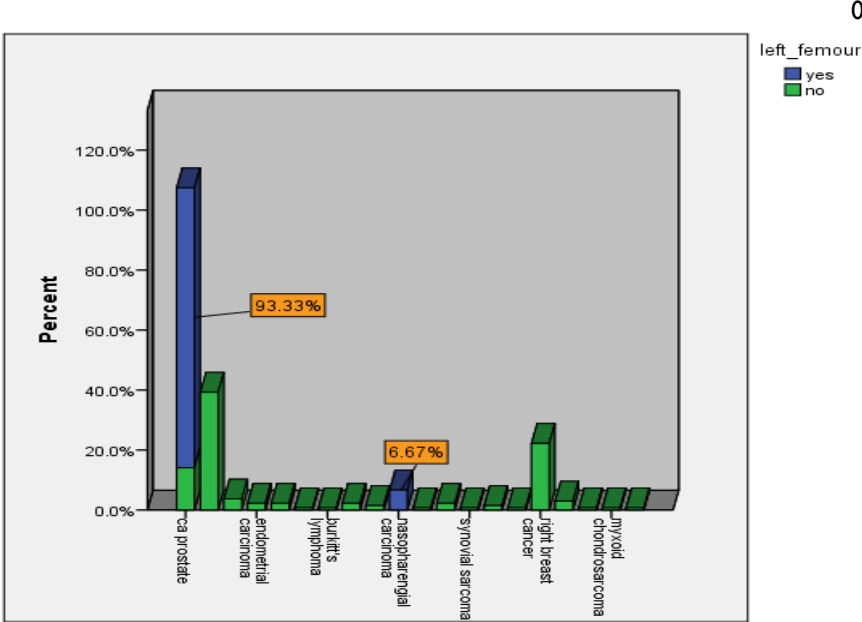


Fig 27: The left femur is involved in (25%) of all bone metastases (93.33%) of patients, the metastases from prostate cancer, and (6.67%) from nasopharyngeal carcinoma .

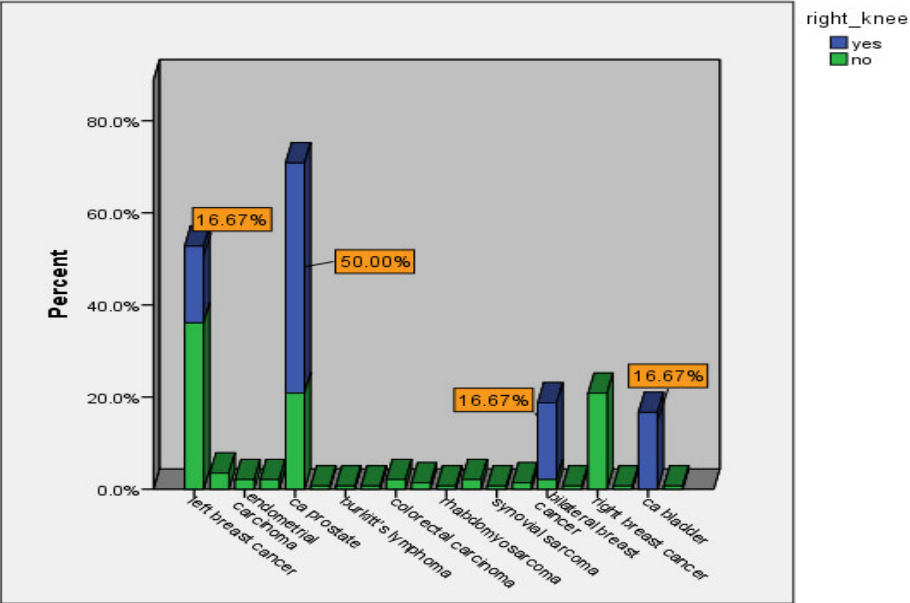


Fig 28: The right knee joint is involved in 10% of all bone metastases (50%) from prostate cancer, 16.67% from left breast cancer, 16.67% from bilateral breast cancer, and 16.67% from the bladder .

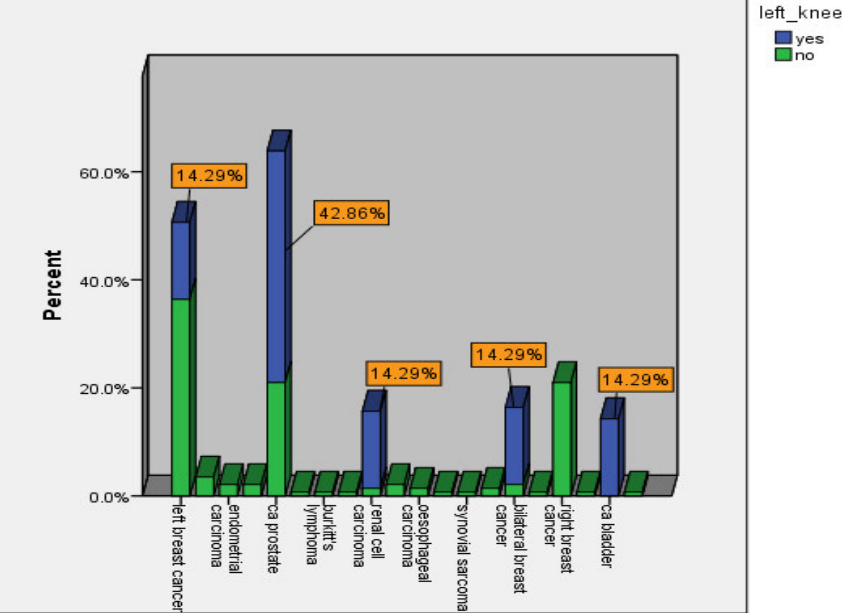


Fig 29: The left knee joint is involved in 11.7% of all bone metastases (42.86%) from prostate cancer, 14.29% from renal cell carcinoma, 14.29% from right breast cancer, 14.29% from bilateral breast cancer, and 14.29% from urinary bladder cancer

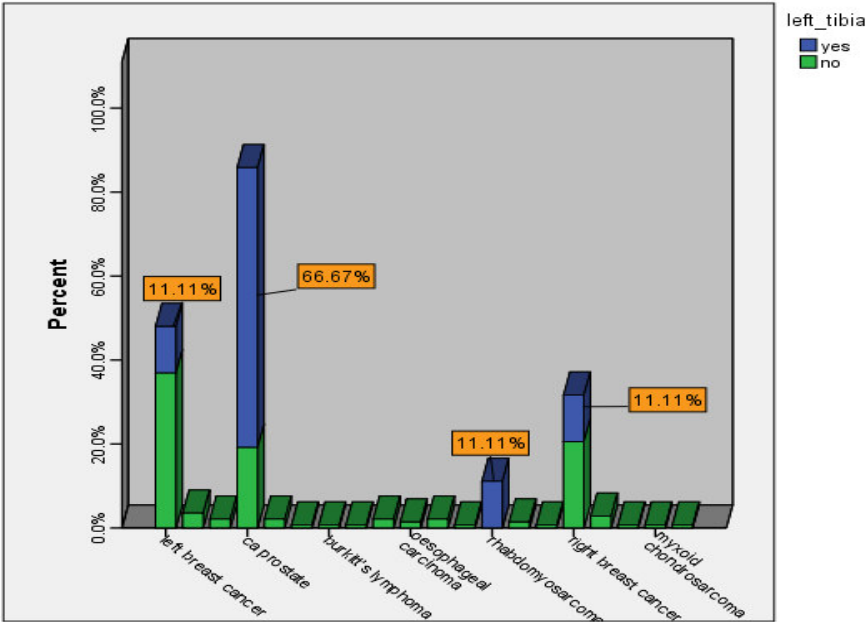
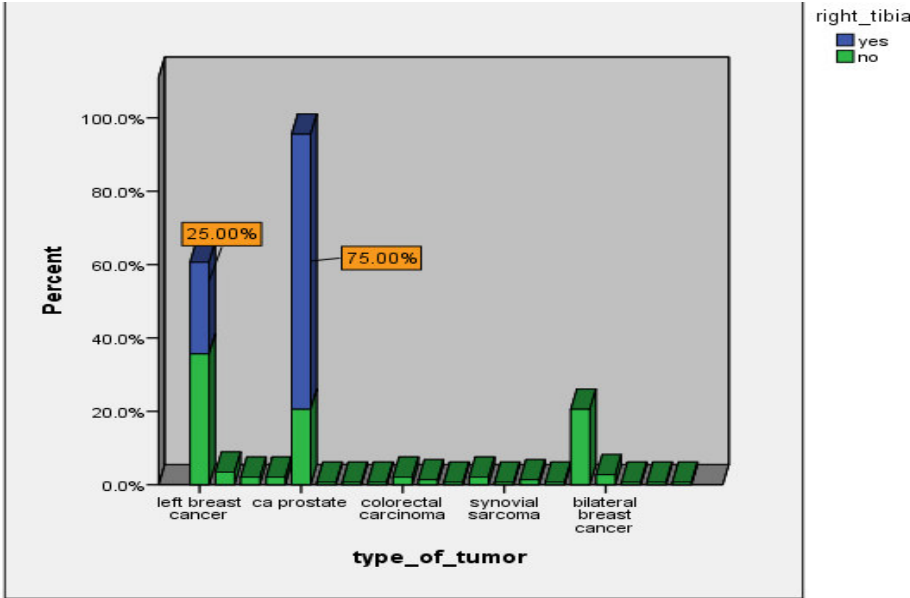


Fig 30: The right tibia is involved in (6.7%) of all bone metastasis (75%) from prostate cancer and (25%) from left breast cancer .

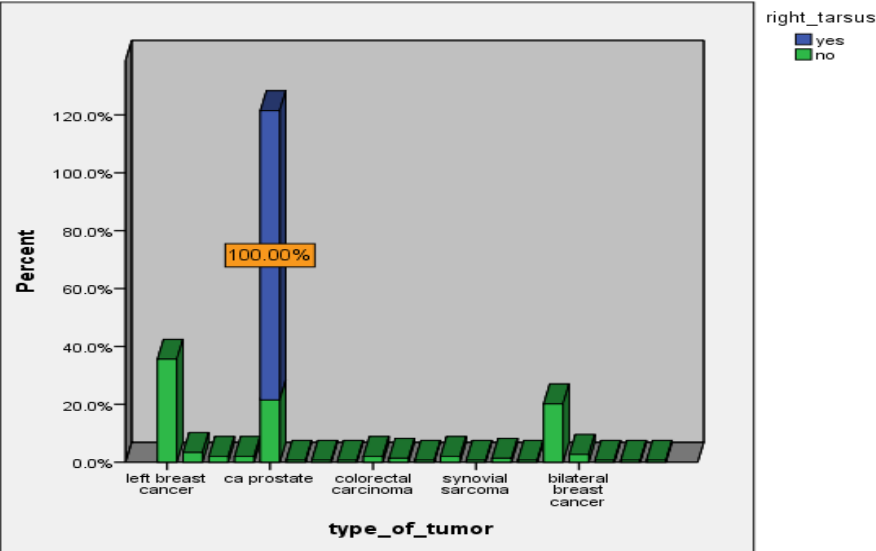


Fig 32: The right tarsus is involved in (1.7%) of all bone metastasis (100%) from prostate cancer .

Fig 31: The left tibia was involved in (15%) of all bone metastases (66.67%) from prostate cancer (11.11%) from left breast cancer, (11.11%) 87.5 from right breast cancer, and (11.11%) from rhabdomyosarcoma.

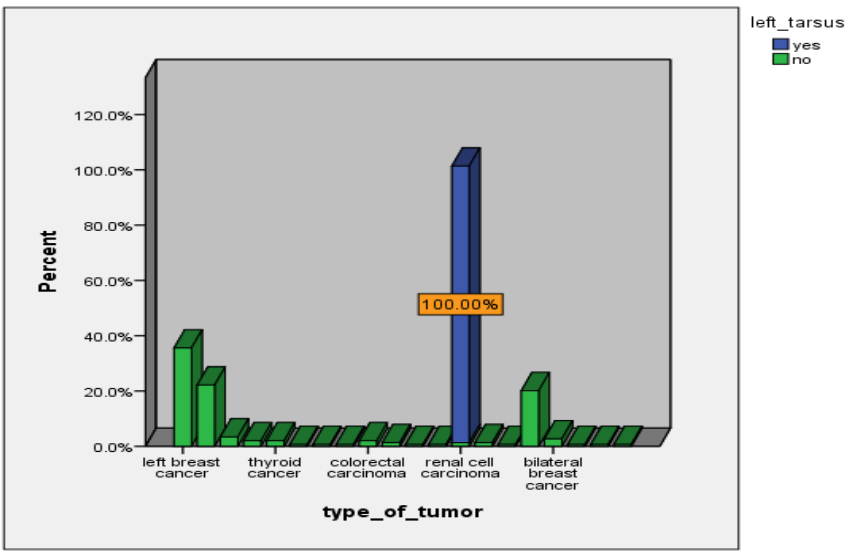


Fig 33: The left tarsus is involved in (1.7%) of all bone metastasis (100%) from renal cell carcinoma.

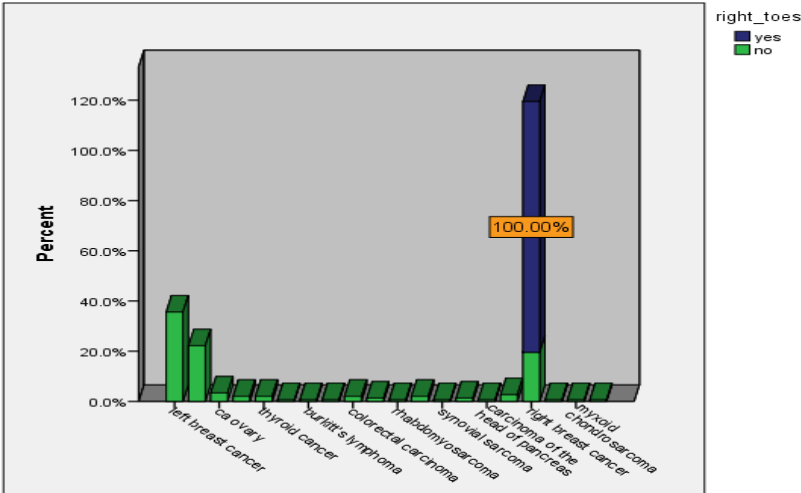


Fig 34: The right toes are involved in (1.7%) of all bone metastases (100%) from right breast carcinoma.

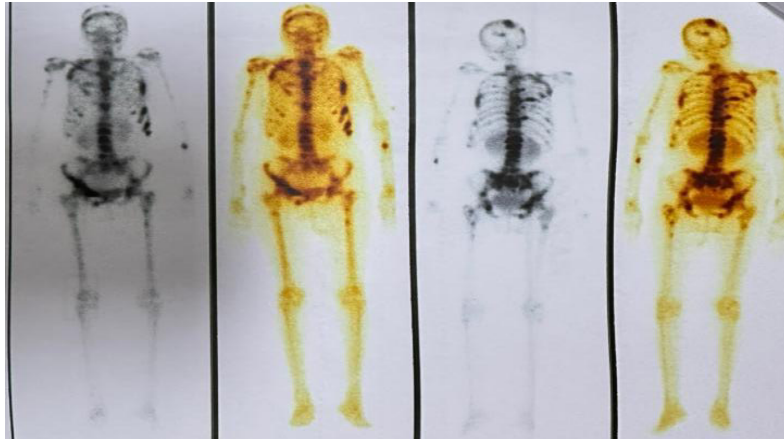


Fig 35: patient with Ca breast: Anterior and posterior views obtained 2 hours after 20 mCi of TC ^{99m}. MDP: Revealed multiples metastasis with active trace uptake is seen involving: skull, sternum, multiple bilateral ribs anteriorly and posteriorly, right humerus, right and left hemipelvis, both femori, and multilevel on vertebral column.

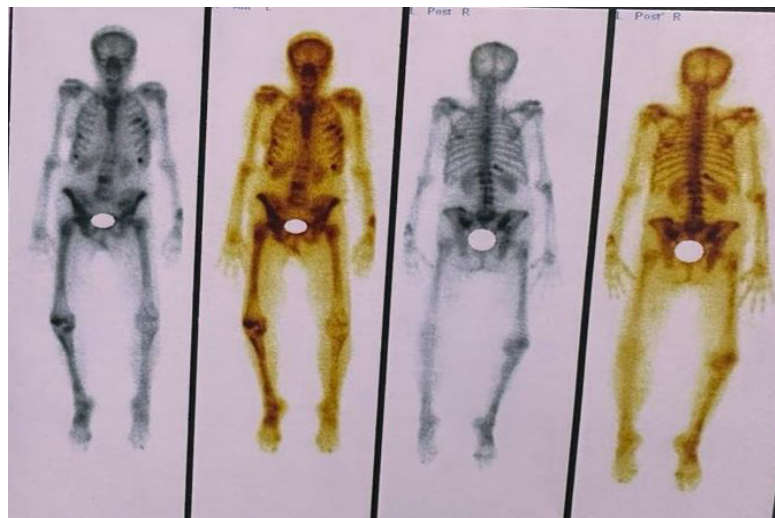


Fig 36: Patient with uterine leiomyosarcoma cancer. Anterior and posterior views obtained 2 hours after 20 mCi of TC ^{99m}. MDP Revealed multiples metastasis with active trace uptake is seen involving: multiple ribs, left iliac bone posteriorly, and multilevel on vertebral column

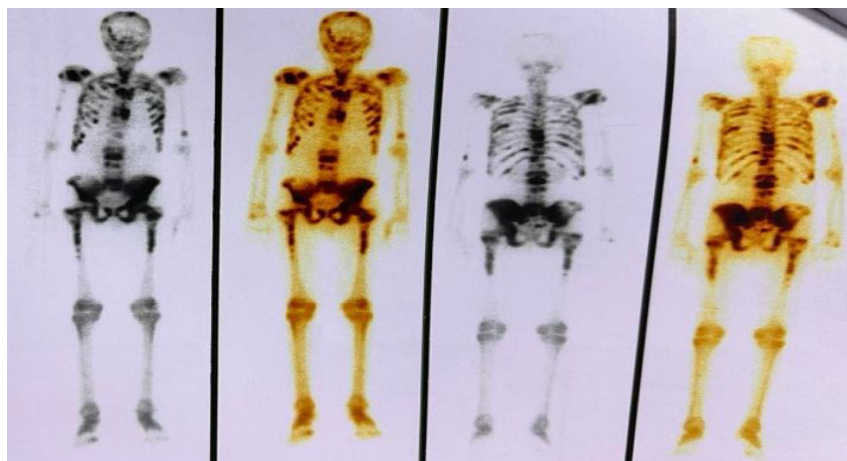


Fig 37: Patient with Ca prostate: Anterior and posterior views obtained 2 hours after 20 mCi of TC ^{99m}. MDP Revealed multiples metastasis with active trace uptake is seen involving: skull, sternum, multiple bilateral ribs anteriorly and posteriorly, both humeri, right and left hemipelvis, both femori, and multilevel on vertebral column.

3. DISCUSSION

Bone metastases represent a frequent and significant manifestation of distant metastasis in malignancies, with a variety of primary cancers demonstrating a propensity to spread to bone tissue. In the present study, 53.3% of patients presented between the ages of 41 and 60, a demographic peak consistent with the broader clinical profile of metastatic bone disease. A higher incidence of bone metastases was observed in women (68.7%) compared to men (31.3%), aligning with findings from an international study by Vahid et al.⁶, where 72.5% of patients were women and 27.5% were men, which has been attributed to the higher incidence of breast cancer in females. This gender disparity in bone metastasis is well-documented, as certain malignancies, particularly breast cancer, have a more frequent predilection for skeletal involvement. In terms of specific malignancies, the prevalence of breast cancer with bone metastasis in this cohort was 43.4%, followed by prostate cancer at 41.7%. This distribution mirrors the findings of Vahid et al. (2016), where breast cancer accounted for 69.1% of cases and prostate cancer for 59.4%. The high frequency of bone metastases in breast and prostate cancers is due in part to their hematogenous spread, which is influenced by the tumor's biology and vascular characteristics. For instance, breast cancer metastasizes predominantly through the venous system, utilizing the posterior intercostal veins and paravertebral venous plexus, facilitating its spread to the spine, ribs, pelvis, and femur. The study found that, among patients with breast cancer, 50.9% had left-sided breast cancer, 46.1% had right-sided breast cancer, and 3% had bilateral breast cancer. This finding, indicating a higher incidence of left-sided breast cancer, is consistent with studies such as that by Magid H. Amer⁷, which also noted a higher incidence of left-sided breast cancer. This pattern may be related to anatomical and hormonal factors, although the exact mechanism remains an area of ongoing investigation. Regarding the distribution of bone metastases, lumbar lesions were the most common, representing 51.7% of cases, followed by thoracic lesions at 50%, and cervical lesions at 16.7%. These findings align with the results from Vahid et al.,⁶ who reported lumbar lesions in 39.6%, thoracic lesions in 34.4%, and cervical lesions in 5.1%. The preferential involvement of lumbar and thoracic vertebrae in metastasis can be attributed to the Batson venous plexus, a network of veins that provides a route for cancer cells to bypass the lungs and directly enter the vertebral column. This phenomenon is particularly relevant in prostate cancer, where the spread to the spine is early and subsequent metastases occur to other skeletal sites, including the ribs and femur. For prostate cancer, the pattern of metastasis observed in this study closely mirrored that of Vahid et al.⁶ with the spine being the most common site, followed by the femur, ribs, and pelvis. This is consistent with the hypothesis that prostate cancer cells are early directed into the spine via the Batson venous plexus, before spreading to other parts of the skeleton. Similarly, the spread of breast cancer to the spine, ribs, and pelvis is well explained by the free communication between the posterior intercostal veins and the paravertebral venous plexus, facilitating these skeletal metastases. This study underscores the significant patterns of bone metastasis in breast and prostate cancers, highlighting the need for vigilant monitoring of skeletal involvement in these malignancies. The findings align closely with international studies, offering further evidence of the shared characteristics of metastatic spread in these common cancers. The Batson venous plexus plays a

crucial role in this process, explaining the preferential metastasis to the spine and other skeletal sites in both breast and prostate cancer.

4. CONCLUSION

Bone is a common site of distant metastasis in malignancies, with a particular predilection for certain primary cancers, such as breast and prostate cancer. Studies show that the age range for bone metastases typically spans from 41 to 60 years, with a higher prevalence in women. Notably, breast cancer, particularly left-sided breast cancer, is more commonly associated with bony metastasis than right or bilateral breast cancer. In our study, 40% of patients with metastatic breast cancer presented with bony metastases, the most frequent of which was located in the spine. This finding aligns with existing literature, which identifies the spine as the primary site for bone metastases in breast cancer patients. Prostate cancer, which is another malignancy that frequently metastasizes to the bones, shares many of the same sites of metastasis. In addition to the spine, the femur, ribs, skull, shoulders, sternum, pelvis, tibia, humerus, knees, hip joints, scapula, sternoclavicular joints, and tarsus are commonly affected in prostate cancer patients. This pattern of metastasis is critical for clinicians to recognize, as it aids in the timely diagnosis and management of metastatic spread. To advance scientific research and improve cancer care in Sudan, it is essential to address the issue of patient data archiving, which remains a significant barrier. We urge the relevant authorities to prioritize the establishment of more nuclear medicine departments across the nation to facilitate early detection and monitoring of metastases. Furthermore, we recommend that the government implement strategies aimed at enhancing early detection and educating the public on cancer risks and available treatment options. A comprehensive evaluation of cancer patients upon their first visit, coupled with regular follow-ups, is vital to manage and potentially slow the progression of malignancies, thus improving patient outcomes and reducing the burden of cancer in Sudan.

5. AUTHORS' CONTRIBUTION STATEMENT

The idea and design of the study were contributed to by all authors. They took care of the material preparation and data collection. The analysis and final draft were finished by Taha Ali Mohammed Taha, Babekir Hassan Mohammed, and Abdalla Atta Abukleawa Mahmoud. The manuscript was written by Motwakil Imam Awadelkareim in its initial draft. Every author offered feedback on earlier drafts of the work. All authors have read and approved the final manuscript.

6. ACKNOWLEDGMENT

Deep appreciation to all of the students at the fascinating Shendi University

7. FUNDING

This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

8. CONFLICT OF INTEREST

Conflict of interest declared none.

9. REFERENCES

1. Jiang W, Rixiati Y, Zhao B, Li Y, Tang C, Liu J. Incidence, prevalence, and outcomes of systemic malignancy with bone metastases. *Journal of Orthopedic Surgery*. May 2020. doi:10.1177/2309499020915989
2. Murali Poduval, Skeletal System Anatomy in Adults, www.emedicine.medscape.com, Jul 28, 2015
3. Allen Gabriel, Breast Anatomy, www.emedicine.medscape.com, Jun 29, 2016
4. Nicolas A Muruve, Prostate Anatomy, www.emedicine.medscape.com, Sep 13, 2017 Page 768, 772, 781-782, 713-714, 668 in Robbins and Cotran pathologic basis of disease (Ninth edition), Kumar, V., Abbas, A. K., & Aster, J. C. (2015). (Mechanisms of Bone Metastasis by G. David Roodman, New England Journal of Medicine, April 15, 2004. Mechanisms of bone metastasis, Gregory R. Mundy, ACS journals, Nov. 20, 2000.
5. Arnold I. Brenner DO, June Koshy, Jose Morey, Cheryl, and Jason DiPoce, The Bone Scan, Science Direct, November 22, 2011.
6. Valid Reza DabbaghKakhki I, Pattern and distribution of bone metastases in common malignant tumors, Pubmed.gov, 2013, Kazem Anvari, Ramin Sadeghi, Anooshe-Sadat Mahmoudian, and Maryam Torabian-Kakhki.
7. Magid H. Amer, Genetic factors and breast cancer laterality, ncbi.nlm.nih.gov, 2014 Apr 16.
8. Keene JS, Sellinger DS, McBeath AA, Engber WD. Metastatic breast cancer in the femur. A search for the lesion at risk of fracture. *Clin Orthop Relat Res*. (203): 282-8; February 1986.
9. Santini D, Vincenzi B, Tonini G, Sammarra M, Quattrocchi CC, Piciocchi S, et al. Osteosclerotic lesions are more common in breast cancer bone metastases. Oct. 2007, *Radiol Med*. 112 (7): 1049-1059.
10. Alarmo EL, Kallioniemi A. Bone morphogenetic proteins: twofold function in carcinogenesis in breast cancer? *Endocrine-related cancer*. June 17, 2010 (2): R123-39...
11. Doot RK, Gralow JR, Specht JM, Peterson LM, Muzi M, Schubert EK, et al. Kinetic analysis of 18F-fluoride PET images of breast cancer bone metastases. *J Nucl Med*. 2010 Apr. 51 (4):521-7.
12. Hung JJ, Jeng WJ, Hsu WH, Wu KJ, Chou TY, Hsieh CC, et al. Prognostic factors of post-recurrence survival in completely resected stage I non-small cell lung cancer with distant metastasis. *Thorax*. 2010 Mar. 65 (3):241-5.
13. Edwards J. Src kinase inhibitors: an emerging treatment option for prostate cancer. *Expert Opin Investing Drugs*. 2010 May. 19 (5):605-14.
14. Bittner M, Wedin R, Bauer H, Tsagozis P. Outcome of surgical treatment for bone metastases caused by colorectal cancer. *J Gastrointest Oncol*. 2021 Oct. 12 (5):2150-2156.
15. British Orthopaedic Oncology Society and British Orthopaedic Association. Metastatic bone disease: a guide to good practice (2015 revision). British Association for Surgical Oncology. Available at https://baso.org.uk/media/61543/boos_mbd_2016_boa.pdf. 2015; accessed: March 13, 2024.
16. Zeng L, Chow E, Bedard G, Zhang L, Fairchild A, Vassiliou V, et al. *Int J Radiat Oncol Biol Phys*. 2012 Nov 1. 84 (3):e337-42. Quality of life following palliative radiation therapy for patients with painful bone metastases: findings of an international study validating the EORTC QLQ-BM22. *Int J Radiat Oncol Biol Phys*. 2012 Nov 1. 84 (3):e337-42. Quality of life following palliative radiation therapy for patients with painful bone metastases: findings of an international study validating the EORTC QLQ-BM22.
17. Frankel BM, Jones T, Wang C. A clinical assessment of segmental polymethylmethacrylate-augmented pedicle screw fixation in patients with osteoporosis-related bone softening and metastatic tumor involvement. *Neurosurgery*. 2007 Sep. 61 (3):531-7; discussion 537-8.
18. Orthopedic management of pelvic lesions and extremities, Harrington KD. *Clin Orthop Relat Res*. 312:136-47, Mar. 1995. *Clin Orthop Relat Res*. 312:136-47, Mar. 1995.
19. Mundy GR, Yoneda T. Facilitation and suppression of bone metastasis. *Clin Orthop Relat Res*. 1995 Mar. (312):34-44.
20. Elaasser B, Arakil N, Mohammad KS. Bridging the Gap in Understanding Bone Metastasis: A Multifaceted Perspective. *Int J Mol Sci*. 2024 Feb 29. 25 (5): (Guise TA, Yin JJ, Taylor SD, Kumagai Y, Dallas M, Boyce BF, et al. Evidence linking parathyroid hormone-related protein to the pathophysiology of osteolysis-mediated human breast cancer. *J Clin Invest*. 1996 Oct 1. 98 (7):1544-9.
21. Cancer statistics. National Cancer Institute. Available at <https://www.cancer.gov/aboutcancer/understanding/statistics>. September 25, 2020; accessed: March 13, 2024.
22. Turpin A, Duterque-Coquillaud M, Vieillard MH. Bone Metastasis: Current State of Play. *Transl Oncol*. 2019 Dec 23. 13 (2):308-320.
23. Key statistics about bone cancer. American Cancer Society. Available at <https://www.cancer.org/cancer/bone-cancer/about/key-statistics.html>. January 12, 2023; accessed: March 13, 2024.
24. Jawad MU, Pollock BH, Wise BL, Zeitlinger LN, O'Donnell EF, Carr-Ascher JR, et al. Sex, racial/ethnic, and socioeconomic disparities in patients with metastatic bone disease. *J Surg Oncol*. 2022 Mar. 125 (4):766-774.
25. Harrington KD. Orthopedic surgical management of skeletal complications of malignancy. *Cancer*. 1997 Oct 15. 80 (8 Suppl):1614-27.
26. Yazawa Y, Frassica FJ, Chao EY, Pritchard DJ, Sim FH, Shives TC. Metastatic bone disease. A study of the surgical treatment of 166 pathologic humeral and femoral fractures. *Clin Orthop Relat Res*. 1990 Feb. (251):213-9.
27. Hirbe AC, Morgan EA, Weilbaecher KN. The CXCR4/SDF-1 chemokine axis: a potential therapeutic target for bone metastases? *Curr Pharm Des*. 2010. 16 (11):1284-90.
28. Kirkinis MN, Spelman T, May D, Choong PFM. Metastatic bone disease of the pelvis and extremities: rationalizing orthopedic treatment. *ANZ J Surg*. 2017 Nov. 87 (11):940-944.
29. Baumber R, Gerrand C, Cooper M, Aston W. Development of a scoring system for survival following

- surgery for metastatic bone disease. *Bone Joint J.* 2021 Nov. 103-B (11):1725-1730 ...
30. Nakanishi K, Hijikata Y, Uchino K, Sugimoto Y, Iba H, Watanabe S, et al. Predicting Skeletal-related Events Using SINS. *Spine (Phila Pa 1976)*. 2024 Mar 13 .
31. Mirels H. Metastatic disease in long bones. A proposed scoring system for diagnosing impending pathologic fractures. *Clin Orthop Relate Res.* 1989 Dec. (249):256-64 .
32. Hipp JA, Springfield DS, Hayes WC. Predicting pathologic fracture risk in the management of metastatic bone defects. *Clin Orthop Relat Res.* 1995 Mar. (312):120-35 .
33. F-fluoride uptake in bone metastasis: morphologic and metabolic analysis on integrated PET/CT, Kawaguchi M, Tateishi U, Shizukuishi K, Suzuki A, Inoue T. May 2010, *Ann Nucl Med.* 24 (4):241-7.
34. Richhariya A, Shore ND, Wang H, Chung K, and SJ all had freed land. Treatment regimens in US community-based urology group practices for patients with bone metastases and prostate cancer. Aug. 2012; 80 (2):293-8; *Urology*.
35. Callstrom MR, Dupuy DE, Solomon SB, Beres RA, Littrup PJ, Davis KW, et al. Percutaneous image-guided cryoablation of painful metastases involving bone: multicenter trial. *Cancer.* 2013 Mar 1. 119 (5):1033-41 .
36. Haider MT, Smit DJ, Taipaleenmäki H. MicroRNAs: Emerging Regulators of Metastatic Bone Disease in Breast Cancer. *Cancers (Basel)*. 2022 Jan 30. 14 (3):
37. Raje N, Terpos E, Willenbacher W, Shimizu K, García-Sanz R, Durie B, et al. An international phase 3 trial that was randomized, double-blind, double-dummy, controlled, and randomized compared denosumab and zoledronic acid to treat bone disease in patients with recently diagnosed multiple myeloma. Mar. 19, 2018; *Lancet Oncol.* 19 (3): 370–381.
38. Bakhshandeh M, Mofid B, Sahinbas H, Faeghi F, Mirzaei H, et al., Faghihi Moghaddam F. The clinical effectiveness of external beam radiation therapy (EBRT) alone versus EBRT plus whole-body hyperthermia was investigated in patients with painful bony metastases in a phase III clinical trial. 2024 Feb 23; *J Therm Biol* 120:103804, 2024.
39. Sacino AN, Rhines LD, Chen H, Sahgal A, Bettegowda C, Maralani P, and others. A new standard of care for spinal metastases is stereotactic body radiation therapy. 2024 Mar 4. 26 (Supplement_1): S76–S87. *Neuro Oncol.*
40. Barrett-Lee P, Simmonds P, Hood K, Coleman R, Abraham J, Casbard A, et al. This phase 3 study compared oral ibandronic acid with intravenous zoledronic acid for the treatment of bone metastases from breast cancer. It was non-inferiority, open-label, and randomized. 2014 Jan 15 (1):114–22; *Lancet Oncol.* Bisphosphonates and additional bone agents as treatments for breast cancer: Wong MH, Stockler MR, Pavlakis N. *Cochrane Database Syst Rev.* 2012; 2:CD003474, February 15.
41. Irelli A, Zugaro L, Di Staso M, Cannita K, Cocciolone V, Lanfiuti Baldi P, et al. Bone-targeted therapy to avoid skeletal-related complications in patients with breast cancer that has spread. *Bone.* 2016 Jun. 87:169-75 .
42. Yu X, Zhu L. Nanoparticles for the Treatment of Bone Metastasis in Breast Cancer: Recent Advances and Challenges. *Int J Nanomedicine.* 2024. 19:1867-1886. Zaikova O, Fosså SD, Bruland OS, Giercksky KE, Sandstad B, Skjeldal S. Radiotherapy or surgery for spine metastases? *Acta Orthop.* 2011 Jun. 82 (3):365-71 .
44. Errani C, Bazzocchi A, Spinnato P, Facchini G, Campanacci L, Rossi G, et al. What's new in the management of bone metastases? *Eur J Orthop Surg Traumatol.* 2019 Oct. 29 (7): 1367-1375 .
45. Orita Y, Sugitani I, Matsuura M, Ushijima M, Tsukahara K, Fujimoto Y, et al. Prognostic factors and the therapeutic strategy for patients with bone metastasis from differentiated thyroid carcinoma. *Surgery.* 2010 Mar. 147 (3):424-31. Sebghati J, Khalili P, Tsagkozis P. Surgical treatment of metastatic bone disease of the distal extremities. *World J Orthop.* 2021 Oct 18. 12 (10):743-750 .
46. Redmond KJ, Lo SS, Soltys SG, Yamada Y, Barani IJ, Brown PD, et al. Consensus guidelines for postoperative stereotactic body radiation therapy for spinal metastases: results of an international survey. *The Journal of Neurosurg Spine.* Mar. 26, 2017, 396–307.
47. Clayer MT, Tang X. Low risk of cardiac events during intramedullary instrumentation of lung cancer metastases. *Acta Orthop.* 2007 Aug. 78 (4):547-50 .
48. Camnasio F, Scotti C, Peretti GM, Fontana F, Fraschini G. Prosthetic joint replacement for long bone metastases: analysis of 154 cases. *Arch Orthop Trauma Surg.* 2008 Aug. 128 (8):787-93. Forauer AR, Kent E, Cwikiel W, Esper P, Redman B. Selective palliative transcatheter embolization of bone metastases from renal cell carcinoma. *Acta Oncol.* 2007. 46 (7):1012-8 .
49. Chow E, van der Linden YM, Roos D, Hartsell WF, Hoskin P, Wu JS, et al. Single versus multiple fractions of repeat radiation for painful bone metastases: a randomized, controlled, non-inferiority trial. *Lancet Oncol.* 2014 Feb. 15 (2):164-71 .
50. Nieder C. Repeat palliative radiotherapy for painful bone metastases. *Lancet Oncol.* 2014 Feb. 15 (2):126-8 .
51. Lutz S, Balboni T, Jones J, Lo S, Petit J, Rich SE, et al. Palliative radiation therapy for bone metastases: Update of an ASTRO Evidence-Based Guideline. *Pract Radiat Oncol.* 2017 Jan.-Feb. 7 (1):4-12.