

ROLE OF PLEURAL BIOPSY IN THE EVALUATION OF THE PLEURAL EFFUSION

*¹Dr. RESHMA S BABU, ²Dr. ULHAS JADHAV, ³Dr. BABAJI GHEWADE

ABSTRACT--*In a hospital where thoracoscopy is not available, pleural biopsy can be used for the diagnosis of exudative effusion in conditions where malignancy is doubted and when results of pleural effusion are not conclusive. In a country with limited resources like India pleural biopsy can still be used for the diagnosis of pleural effusion. 1. To study use of pleural biopsy in evaluation of pleural effusion. 2. To correlate the pleural biopsy with biochemical and cytological findings of pleural fluid. 3. To find the complications of pleural biopsy. An observational cross sectional study will be conducted in Acharya Vinoba Bhave Rural Hospital among 70 patients admitted with exudative pleural effusion. Pleural biopsy will be done and pleural fluid investigations will be sent. Cause of pleural effusion can be obtained from the pleural biopsy. Malignancy and tuberculosis are the most frequent reasons of pleural effusion according to previous studies. Whether pleural biopsy gives definitive diagnosis as compared to pleural fluid cytology or pleural fluid CBNAAT. Pleural biopsy is associated with minimal post procedure complications. Pleural Biopsy helps in the identifying the etiology of undiagnosed pleural effusion. Complications associated with pleural biopsy is less. Closed pleural biopsy helps identifying cause of the exudative pleural effusion when thoracoscopy is not available.*

KEY WORDS-- *pleural biopsy, pleural effusion, thoracoscopy, tuberculosis, malignant pleural effusion*

I. INTRODUCTION

Pleural effusion is the collection of pleural fluid in the pleural space due to imbalance between rate of absorption and formation of pleural fluid. Pleural effusion itself not a disease but it's sign of an underlying disease. There are many causes of pleural effusions. Pleural effusions are generally classified into two broad categories- exudative and transudative pleural effusion (1-4).

When a patient with undiagnosed pleural effusion is evaluated, first question is to find whether the patient has transudate or exudate effusion, Transudative pleural effusion is due to systemic disease like heart failure, cirrhosis, nephrotic syndrome or hypoalbuminemia in which underlying pleura remains normal. In exudative effusion, the integrity of pleura abnormal due to tuberculosis, malignancy or other infections. The reason to make this differentiation is that additional diagnostic procedures are indicated for exudative pleural effusions to define cause of effusion (5-8).

Diagnosis of pleural effusion made by history, physical examination, chest X ray and analysis of pleural fluid. Pleural fluid obtained by the thoracentesis is sent for biochemical analysis, cytological analysis and

¹ *JUNIOR RESIDENT, RESPIRATORY MEDICINE, JNMC, DMIMS, reshmasbabu19@gmail.com, 8075708315

² PROFESSOR, RESPIRATORY MEDICINE, JNMC, DMIMS, drulhasjadhav@gmail.com 9970819099

³ HEAD OF DEPARTMENT, RESPIRATORY MEDICINE, JNMC, DMIMS, crownbabaji@gmail.com, 9822342770

microbiological investigations. In spite of all this, it is not possible to establish a diagnosis in all the cases. The common cause of undiagnosed pleural effusion is tuberculosis and malignancy. Invasive modalities like closed pleural biopsy or thoracoscopy is required **when** diagnosis cannot be obtained (3,9-12).

Thoracoscopy involves high instrument cost and procedure requires intercostal drainage, long hospital admission as well as huge hospital expenses. In comparison pleural biopsy is simple, cost effective procedure which can be easily done with minimal procedure related complications. Pleural biopsy is performed by Abrahams needle. Pleural biopsy can be used in the diagnosis of exudative pleural effusion, especially when malignancy is suspected and when the results of pleural effusion are inconclusive especially in a hospital where thoracoscopy is not available. In a country with limited resources like India, pleural biopsy can still be used to evaluate the causes of pleural effusion (12-15). With this background the study will be able to find out the usefulness of pleural biopsy to identify cause of pleural effusion.

Thoracoscopy is investigation of choice where diagnosis of pleural effusion is inconclusive. But it is a costly procedure and requires the he back up of thoracic surgery. Closed pleural biopsy can be easily performed with minimal procedure related complication (16-19). Pleural biopsy can be used to obtain an etiological diagnosis in exudative pleural effusion, especially when malignancy is suspected and when the results of pleural effusion are not conclusive in a hospital where thoracoscopy is not available.

II. OBJECTIVES

- 1.To study use of pleural biopsy in evaluation of pleural effusion.
- 2.To correlate the pleural biopsy with biochemical and cytological parameters of pleural fluid.
3. To identify the complications of pleural biopsy.

III. METHODS

Study design: Observational cross sectional study.

Setting: Study will be conducted in the AVBRH, a tertiary care hospital attached to Jawaharlal Nehru Medical Collage (JNMC), situated in the rural area of Sawangi (Meghe) Wardha, in Central India from July 2019 to July 2021.

Participants:

Patient with pleural effusion will be selected on basis of inclusion and exclusion criteria among patients admitted in AVBRH.

INCLUSION CRITERIA:

- 1.Patient more than 18 of years.
- 2.Patient with exudative pleural effusion as per Lights criteria
- 3.Chest X-ray showing signs of pleural effusion

EXCLUSION CRITERIA:

1. Bleeding and coagulation disorders
2. Patient who are not cooperative and not giving consent.

Variables:

Exudative pleural effusion based on light's criteria
Pleural biopsy (histopathology report) for diagnosing the cause of the pleural effusion.
Pleural fluid CBNAAT and ADA for diagnosing tuberculosis
Pleural fluid cytology for diagnosing the malignancy.

IV. DATA SOURCES/ MEASUREMENT

Pleural fluid sample obtained from the patient will be evaluated for pleural fluid ADA

Pleural fluid sample will be sent for pleural fluid CBNAAT.

Pleural fluid cytology will be obtained from pathology department, AVBRH

Pleural Biopsy specimen will send to histopathology department, AVBRH

Bias: There will be bias in selection of patients and procedure and technique related bias.

Study size:

Sample size = $Z^2(SN(1-SN)/W^2)$

Z = Z VALUE (1.96 for 95% confidence interval)

SN = SENSITIVITY (90%)

W = ABSOLUTE PRECISION (5%)

By considering sensitivity of pleural biopsy in diagnosis of pleural effusion as 90% and confidence interval of 95% i.e. 1.96 and absolute precession of 5 % the calculated sample size is 70

Statistical methods:

70 patients will be selected for the study. Pleural biopsy will be done using the Abram's needle. Specimen will be send for histopathological analysis. Histopathological results will be collected and will be analysed using SPSS 10.

V. EXPECTED OUTCOMES/RESULTS

Participants: 70 cases of exudative pleural effusion admitted in AVBRH will be selected for the study after obtaining written informed consent for pleural biopsy.

Descriptive data: Patients of pleural effusion will be classified in to exudative and transudative. Exudative effusion patients will undergo pleural biopsy Outcome data: Pleural biopsy report

Main results: Pleural biopsy report which revealed the cause of pleural effusion. Most conditions which will cause pleural effusion are malignancy and tuberculosis. Pleural biopsy can be used instead of thoracoscopy for diagnosis of pleural effusion in places where thoracoscopy is not available.

VI. DISCUSSION

Pleural biopsy will help in diagnosis of the exudative pleural effusion in places where thoracoscopy is not available. Main cause of the exudative effusion could be tuberculosis and malignancy. Pleural biopsy can be used in places where thoracoscopy is not available. It is low cost procedure associated with minimal procedure related complications. A couple of studies were considered and reviewed for understanding the scenario of related factors and conditions in this region (20-75).

Limitations: The main limitation of the study is that there will be no thoracoscopy to compare with. Second limitation of the study will be the technique used for the pleural biopsy which can vary with persons. One attempt of biopsy was done in the current study but other studies had 2/3 biopsies done.

Interpretation: Pleural biopsy will be used in diagnosis of the exudative pleural effusion. Histopathology report from the biopsy specimen gives diagnosis of exudative pleural effusion which will be compared with pleural fluid cytology, pleural fluid ADA and CBNAAT. Cause of pleural effusion will be confirmed by pleural fluid cytology and Pleural fluid ADA and CBNAAT. Pleural biopsy can be used instead of thoracoscopy in places where thoracoscopy is not available. Complication associated with pleural biopsy is minimal.

Generalisability: According to study pleural biopsy can be used in the diagnosis of exudative pleural effusion. In the current setting, tuberculosis and malignancy was most common cause. The complication associated with procedure is less and is low cost procedure.

REFERENCES

1. Light RW, Diagnostic principles in pleural disease. *Eur Respir J* 1997;10;476-81
2. Saguil A, Wyrick K, Hallgren J. Diagnostic Approach to pleural effusion. *Am Fam physician* 2014;90(2): 99-104
3. Hooper C, Lee Y C, Maskell N. Investigation of a unilateral pleural effusion in adults. *British Thoracic Society Pleural disease guideline 2010*. *Thorax* 2010;65 suppl 2: ii 4-17
4. Sudipta Pandit, Arunabha Datta. Role of pleural biopsy in etiological diagnosis of pleural effusion. *Lung India* 2010 ;27 (4):202-204.
5. Somnath Bhattacharya, Tapan D Bairagya. Closed pleural biopsy is still useful in the evaluation of malignant pleural effusion. *Journal of laboratory physicians* 2012;4(1):35-38.
6. Govind Singh Rajawat, Supreet Batra. Diagnostic yield and safety of closed needle pleural biopsy in exudative pleural effusion. *Avicenna journal of medicine* 2017;7(3):121-12
7. Robert H.Poe, Robert.H.Israel; Mark J Utell. Sensitivity, Specificity and predictive values of closed pleural biopsy. *Arch Intern med*.1984;144(2):325-328.
8. Gouda A M, Dalati T A, Al -Shareef NS. A comparison between Cope and Abrams needle in the diagnosis of pleural effusion. *Ann Thorac Med* 2006; 1:12-15. 53.
9. Dutt N, Agarwal D. Closed pleural biopsy: A victim of Western advancement. *Lung India* 2011:28
10. . Sharma, S. K., J. Chaubey, B. K. Singh, R. Sharma, A. Mittal, and A. Sharma. "Drug Resistance Patterns among Extra-Pulmonary Tuberculosis Cases in a Tertiary Care Centre in North India."

- INTERNATIONAL JOURNAL OF TUBERCULOSIS AND LUNG DISEASE 21, no. 10 (October 2017): 1112–17. <https://doi.org/10.5588/ijtld.16.0939>. 24.
11. Gupta, Vivek, and Arvind Bhake. “Assessment of Clinically Suspected Tubercular Lymphadenopathy by Real-Time PCR Compared to Non-Molecular Methods on Lymph Node Aspirates.” *ACTA CYTOLOGICA* 62, no. 1 (February 2018): 4–11. <https://doi.org/10.1159/000480064>.
 12. Ghorpade, Deesha, Sheetu Singh, Deepak Talwar, Sagar Chandrashekariah, Surya Kant, Rajesh Swarnakar, Srinivas Rajagopala, et al. “Post-Tuberculosis Bronchiectasis in India: Outcomes of the Indian EMBARC Registry.” *EUROPEAN RESPIRATORY JOURNAL* 52, no. 62 (September 15, 2018). <https://doi.org/10.1183/13993003.congress-2018.PA2748>
 13. Gupta, Vivek, and Arvind Bhake. “Assessment of Clinically Suspected Tubercular Lymphadenopathy by Real-Time PCR Compared to Non-Molecular Methods on Lymph Node Aspirates.” *ACTA CYTOLOGICA* 62, no. 1 (February 2018): 4–11. <https://doi.org/10.1159/000480064>.
 14. “Reactive Lymphoid Hyperplasia or Tubercular Lymphadenitis: Can Real-Time PCR on Fine-Needle Aspirates Help Physicians in Concluding the Diagnosis?” *ACTA CYTOLOGICA* 62, no. 3 (2018): 204–8. <https://doi.org/10.1159/000488871>.
 15. Sharma, Surendra Kumar, Alladi Mohan, Achintya Dinesh Singh, Hridesh Mishra, Sonali Jhanjee, Ravindra Mohan Pandey, Binit Kumar Singh, et al. “Impact of Nicotine Replacement Therapy as an Adjunct to Anti-Tuberculosis Treatment and Behaviour Change Counselling in Newly Diagnosed Pulmonary Tuberculosis Patients: An Open-Label, Randomised Controlled Trial.” *SCIENTIFIC REPORTS* 8 (June 11, 2018). <https://doi.org/10.1038/s41598-018-26990-5>.
 16. Sharma, S. K., R. Sharma, B. K. Singh, V. Upadhyay, and I. Mani. “A Study of Non-Tuberculous Mycobacterial (NTM) Disease Among Tuberculosis Suspects at a Tertiary Care Center in North India.” *AMERICAN JOURNAL OF RESPIRATORY AND CRITICAL CARE MEDICINE* 199 (2019).
 17. Sharma, Surendra K., and Keertan Dheda. “What Is New in the WHO Consolidated Guidelines on Drug-Resistant Tuberculosis Treatment.” *INDIAN JOURNAL OF MEDICAL RESEARCH* 149, no. 3 (March 2019): 309–12. https://doi.org/10.4103/ijmr.IJMR_579_19.
 18. Sharma, Surendra K., and Keertan Dheda. “What Is New in the WHO Consolidated Guidelines on Drug-Resistant Tuberculosis Treatment.” *INDIAN JOURNAL OF MEDICAL RESEARCH* 149, no. 3 (March 2019): 309–12. https://doi.org/10.4103/ijmr.IJMR_579_19.
 19. Dhar, Raja, Sheetu Singh, Deepak Talwar, Sagar Chandrashekariah, Surya Kant, Rajesh Swarnakar, Srinivas Rajagopala, et al. “Phenotypes in Bronchiectasis from the EMBARC India Registry.” *EUROPEAN RESPIRATORY JOURNAL* 52, no. 62 (September 15, 2018). <https://doi.org/10.1183/13993003.congress-2018.OA4952>
 20. Modi S, Agrawal A, Bhake A, Agrawal V. Role of adenosine deaminase in pleural fluid in tubercular pleural effusion. *J Datta Meghe Inst Med Sci Univ* 2018;13(4):163-167.
 21. Gotarkar S, Ingole A. Knowledge of Anganwadi worker with respect to early childhood development. *J Datta Meghe Inst Med Sci Univ* 2018;13(4):168-170.
 22. Goje K, Phatak S. Testicular torsion causing infarction of testis, ultrasonography and color Doppler imaging. *J Datta Meghe Inst Med Sci Univ* 2018;13(4):215-216.
 23. Sarode RD, Tendolkar VD. Psychological pain as predictor of impulse control among BAMS new entrants: A correlation study. *J Datta Meghe Inst Med Sci Univ* 2018;13(4):171-174.

24. Mishra KK, Kelkar P, Kumar K. An interesting case of trichotillomania in a pre-school child. *J Indian Assoc Child Adolesc Ment Health* 2018;14(4):131-135.
25. Sthapak E, Gajbe U, Singh BR. Study of communication between musculocutaneous and median nerves in man. *J Anat Soc India* 2018;67: S37-S44.
26. Tripathi A, Avasthi A, Grover S, Sharma E, Lakdawala BM, Thirunavukarasu M, et al. Gender differences in obsessive-compulsive disorder: Findings from a multicentric study from northern India. *Asian J Psychiatry* 2018; 37:3-9.
27. Yeola ME, Gode D, Bora AK. Evaluation of abdominal malignancies by minimal access surgery: Our experience in a rural setup in central India. *World J Laparoscopic Surg* 2018;11(3):115-120.
28. Srivastava TK, Mishra V, Waghmare LS. Formative assessment classroom techniques (FACTs) for better learning in pre-clinical medical education: A controlled trial. *J Clin Diagn Res* 2018;12(9):JC01-JC08.
29. Balwani M, Bawankule C, Ramteke V, Pasari A. Hepatitis C virus, directly acting antivirals and Guillain-Barré syndrome. *Saudi J Kidney Dis Transpl* 2018;29(5):1237-1239.
30. Balwani MR, Pasari A, Meshram A, Jawahirani A, Tolani P, Laharwani H, et al. An initial evaluation of hypokalemia turned out distal renal tubular acidosis secondary to parathyroid adenoma. *Saudi J Kidney Dis Transpl* 2018;29(5):1216-1219.
31. Goyal RC, Choudhari SG, Tankhiwale SR. Assessment of competency based medical internship training with 'cumulative grade points average system'-An innovative step towards meeting 'vision 2015' of medical council of india. *Indian J Public Health Res Dev* 2018;9(8):155-162.
32. Yeola ME, Gode D, Bora AK. Diagnostic laparoscopy as an effective tool in evaluation of intra-abdominal malignancies. *World J Laparoscopic Surg* 2018;11(2):68-75.
33. Sharma S, Singh AD, Sharma SK, Tripathi M, Das CJ, Kumar R. Gallium-68 DOTA-NOC PET/CT as an alternate predictor of disease activity in sarcoidosis. *Nucl Med Commun* 2018;39(8):768-778.
34. Daigavane S, Prasad M. To observe the proportion of amblyopia among children presenting in a rural hospital in Central India. *J Datta Meghe Inst Med Sci Univ* 2018;13(3):119-121.
35. Gadge A, Acharya N, Shukla S, Phatak S. Comparative study of transvaginal sonography and hysteroscopy for the detection of endometrial lesions in women with abnormal uterine bleeding in perimenopausal age group. *J SAFOG* 2018;10(3):155-160.
36. Anjankar SD. Urethral protrusion of the distal end of shunt. *J Pediatr Neurosci* 2018;13(3):371-372.
37. Swarnkar M, Pandey P. Heterotopic subserosal pancreatic tissue in jejunum. *Formosan J Surg* 2018;51(4):167-170.
38. Choudhari MS, Sonkusale MI, Deshpande RA. Sudden cardiac arrest on 5 th day after coronary artery bypass graft surgery: Diagnostic dilemma. *Ann Card Anaesth* 2018;21(3):341-342.
39. Kirnake V, Arora A, Sharma P, Goyal M, Chawlani R, Toshniwal J, et al. Non-invasive aspartate aminotransferase to platelet ratio index correlates well with invasive hepatic venous pressure gradient in cirrhosis. *Indian J Gastroenterol* 2018;37(4):335-341.
40. Kürhade G, Nayak BS, Kurhade A, Unakal C, Kurhade K. Effect of martial arts training on IL-6 and other immunological parameters among Trinidadian subjects. *J Sports Med Phys Fitness* 2018;58(7-8):1110-1115.

41. Balwani MR, Bawankule C, Khetan P, Ramteke V, Tolani P, Kute V. An uncommon cause of rapidly progressive renal failure in a lupus patient: Pauci-immune crescentic glomerulonephritis. *Saudi J Kidney Dis Transpl* 2018;29(4):989-992.
42. Mohite D, Hande A, Gupta R, Chaudhary M, Mohite P, Patil S, et al. Immunohistochemical evaluation of expression pattern of p53, p63, and p73 in epithelial dysplasia. *J Datta Meghe Inst Med Sci Univ* 2018;13(3):122-129.
43. Rathi N, Chandak M, Mude G. Comparative evaluation of dentinal caries in restored cavity prepared by galvanic and sintered burs. *Contemp Clin Dent* 2018;9(5): S23-S27.
44. Gupta V, Bhake A. Reactive Lymphoid Hyperplasia or Tubercular Lymphadenitis: Can Real-Time PCR on Fine-Needle Aspirates Help Physicians in Concluding the Diagnosis? *Acta Cytol* 2018;62(3):204-208.
45. Zodpey S, Sharma A, Zahiruddin QS, Gaidhane A, Shrikhande S. Allopathic Doctors in India: Estimates, Norms and Projections. *J Health Manage* 2018;20(2):151-163.
46. Yadav S, Agrawal M, Hariharan C, Dewani D, Vadera K, Krishna N. A comparative study of serum lipid profile of women with preeclampsia and normotensive pregnancy. *J Datta Meghe Inst Med Sci Univ* 2018;13(2):83-86.
47. Bhinder HHPS, Kamble TK. The study of carotid intima-media thickness in prediabetes and its correlation with cardiovascular risk factors. *J Datta Meghe Inst Med Sci Univ* 2018;13(2):79-82.
48. Munjal R, Mudey G. Nasal carriage of *Staphylococcus aureus* among undergraduate medical students: Prevalence and antibiogram including methicillin resistance, inducible clindamycin resistance, and high-level mupirocin resistance. *J Datta Meghe Inst Med Sci Univ* 2018;13(2):91-94.
49. Mittal V, Jagzape T, Sachdeva P. Care seeking behaviour of families for their sick infants and factors impeding to their early care seeking in rural part of central India. *J Clin Diagn Res* 2018;12(4):SC08-SC12.
50. Choudhary S, Tarafdar P, Jawade S, Singh A. A point to note in pili torti. *Int J Trichology* 2018;10(2):95-97.
51. Madke B, Gardner JM. Enhanced worldwide dermatology-pathology interaction via Facebook, Twitter, and other social media platforms. *Am J Dermatopathol* 2018;40(3):168-172.
52. Girish M, Rawekar A, Jose S, Chaudhari U, Nanoti G. Utility of Low Fidelity Manikins for Learning High Quality Chest Compressions. *Indian J Pediatr* 2018;85(3):184-188.
53. Goswami J, Balwani MR, Kute V, Gumber M, Patel M, Godhani U. Scoring systems and outcome of chronic kidney disease patients admitted in intensive care units. *Saudi J Kidney Dis Transpl* 2018;29(2):310-317.
54. Mohite PM, Anjankar AJ, Patnod S. Organo PHOSPHORUS pOISONING: Prognostic value of GCS score and other clinical indicators in assessing the final outcome. *J Indian Acad Forensic Med* 2018;40(2):197-205.
55. Mathur K, Ninave S, Patond S, Ninave S, Wankhade P. A comparative study of estimation of stature by Bertillon's system among individuals of different regions of India. *J Indian Acad Forensic Med* 2018;40(3):301-306.
56. Kumar S, Bhayani P, Hathi D, Bhagwati J. Hyponatremia initial presenting feature of normal pressure hydrocephalus in elderly patient: A rare case report. *J Gerontology Geriatrics* 2018;66(3):156-157.

57. Jaiswal S, Banait S, Daigavane S. A comparative study on peripapillary retinal nerve fiber layer thickness in patients with iron-deficiency anemia to normal population. *J Datta Meghe Inst Med Sci Univ* 2018;13(1):9-11.
58. Deshpande P, Gupta V, Bhake A. Methylation pattern of retrotransposons: Biomarker for human cancer. *J Datta Meghe Inst Med Sci Univ* 2018;13(1):66-70.
59. Phatak S, Marfani G. Galactoceles ultrasonography and elastography imaging with pathological correlation. *J Datta Meghe Inst Med Sci Univ* 2018;13(1):1-3.
60. Swarnkar M, Agrawal A. Kimura's disease. *Formosan J Surg* 2018;51(1):26-28.
61. Chiwhane A, Pradeep. Study of rhythm disturbances in acute myocardial infarction. *J Assoc Phys India* 2018;66(January):54-58.
62. Gupta V, Bhake A. Assessment of Clinically Suspected Tubercular Lymphadenopathy by Real-Time PCR Compared to Non-Molecular Methods on Lymph Node Aspirates. *Acta Cytol* 2018;62(1):4-11.
63. Anjankar S. Askin's tumor in adult: A rare clinical entity. *J Datta Meghe Inst Med Sci Univ* 2018;13(1):54-57.
64. Jain J, Banait S, Tiewsoh I, Choudhari M. Kikuchi's disease (histiocytic necrotizing lymphadenitis): A rare presentation with acute kidney injury, peripheral neuropathy, and aseptic meningitis with cutaneous involvement. *Indian J Pathol Microbiol* 2018;61(1):113-115.
65. Jain V, Waghmare L, Shrivastav T, Mahakalkar C. SNAPPS facilitates clinical reasoning in outpatient settings. *Educ Health* 2018;31(1):59-60.
66. Bains SK, John P, Nair D, Acharya S, Shukla S, Acharya N. Aptitude of medical research in undergraduate students of a medical university - Miles to go before we sow. *J Clin Diagn Res* 2017;11(12):JC07-JC11.
67. Taksande A, Meshram R, Yadav P, Lohakare A. Rare presentation of cerebral venous sinus thrombosis in a child. *J Pediatr Neurosci* 2017;12(4):389-392.
68. Choudhari MS, Charan N, Sonkusale MI, Deshpande RA. Inadvertent diversion of inferior vena cava to left atrium after repair of atrial septal defect - Early diagnosis and correction of error: Role of intraoperative transesophageal echocardiography. *Ann Card Anaesth* 2017;20(4):481-482.
69. Swarnkar M, Jain SC. Heterotopic subserosal pancreatic tissue in Jejunum-an incidental rare finding. *J Krishna Inst Med Sci Univ* 2017;6(4):105-108.
70. Taksande A, Meshram R, Yadav P, Borkar S, Lohkare A, Banode P. A rare case of Budd Chiari syndrome in a child. *Int J Pediatr* 2017;5(10):5809-5812.
71. Gupta V, Bhake A. Diagnosis of clinically suspected and unsuspected tubercular lymphadenopathy by cytology, culture, and smear microscopy. *Indian J Tuberc* 2017;64(4):314-317.
72. Gupta V, Bhake A. Clinical and cytological features in diagnosis of peripheral tubercular lymphadenitis – A hospital-based study from central India. *Indian J Tuberc* 2017;64(4):309-313.
73. Sharma SK, Chaubey J, Singh BK, Sharma R, Mittal A, Sharma A. Drug resistance patterns among extra-pulmonary tuberculosis cases in a tertiary care centre in North India. *Int J Tuberc Lung Dis* 2017;21(10):1112-1117.
74. Jyoti J, Nitin V, Shashank B, Pradeep D. Gamma glutamyl transferase levels in patients with acute coronary syndrome: A cross-sectional study. *J Cardiovasc Dis Res* 2017;8(4):121-125.
75. Ali S, Ghewade B, Jadhav U, Cladius S. Study of serum interferon gamma in tubercular pleural effusions. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):93-98.