

**DESCRIPTIVE STUDY ON ANALYSIS OF
ABDOMINAL TUBERCULOSIS AND INCIDENCE
OF TUBERCULOSIS IN PERIANAL FISTULAE AT
MADRAS MEDICAL COLLEGE**

Dissertation Submitted for

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BONAFIDE CERTIFICATE

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I am fortunate to have my postgraduate colleagues, for their invaluable suggestions, relentless help for shouldering my responsibilities.

DR.BALAJI TIROUVINGADAME.B

DECLARATION

I, certainly declare that this dissertation titled, “**ANALYSIS OF ABDOMINAL TUBERCULOSIS AND INCIDENCE OF TUBERCULOSIS IN PERIANAL FISTULAE AT MADRAS MEDICAL COLLEGE** ”, represent a genuine work of mine . The contributions of any supervisors to the research are consistent with normal supervisory practice, and are acknowledged.

I, also affirm that this bonafide work or part of this work was not submitted by me or any others for any award, degree or diploma to any other university board, neither in India or abroad. This is submitted to The Tamil Nadu Dr.MGR Medical University, Chennai in partial fulfillment of the rules and regulation for the award of Master of Surgery Degree Branch 1 (General Surgery).

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INTRODUCTION

Tuberculosis is a disease caused by the bacterium *Mycobacterium* species which is one of the commonest disease known to mankind. It is a well known fact that it has its reputation of being one of the greatest killer diseases. There has been a trend of increased incidence of tuberculosis in human race in the last three decades which can be partly attributed to increase in population, social deprivation and HIVinfection.

The morbidity and mortality due to tuberculosis leads to a discovery of newer drugs and more emphasis on the disease. With improved socioeconomic conditions and disease diagnosis in the western countries, Tuberculosis in the western countries impose little clinical problem. But in most developing countries like India, it remains to be a major health hazard and major health related mortality, morbidity both physically and psychologically. As we know, Tuberculosis is one of the important gastroenterological problems inTropics.

Gastrointestinal Tuberculosis is one of the earliest known diseases which still remain the disease with diagnostic enigma due to its perplexing protean clinical manifestations. Importance of the condition lies in the diversity of the presentations and its wide spread effects on

the body. It may present as acute, subacute, chronic forms and give rise to malabsorption, more when associated with stricture formation.

The symptoms and signs often quite vague and laboratory investigations and radiological findings are sometimes non conclusive. There is no single feature which is diagnostic for abdominal Tuberculosis. In case of any localized involvement of the structures of the abdomen the presenting clinical picture will mimic the disease of the organ only. In various studies about abdominal tuberculosis worldwide, the results show wide range in its disparity in the nature course of disease, in its diagnosis and management. We planned a study on abdominal tuberculosis to understand the nature of disease process and to evaluate the various laboratory and radiological investigations and to study the management of the disease.

REVIEW OF LITERATURE

The diagnosis of abdominal tuberculosis is obscure. Joseph Walsh in 1909 remarked that “It is impossible to diagnose abdominal tuberculosis with any degree of certainty”²³. Unfortunately this remains even today in situations where it is relatively common. The signs and symptoms are often vague and laboratory investigation and radiological findings are non-specific. Many cases go unnoticed until a surgically removed specimen is examined Histo-pathologically. A majority of stenotic lesions of the bowel are of tuberculous in nature and has been found on the operating table has one of the common findings in cases of acute intestinal obstruction.¹

The term abdominal tuberculosis refers to tuberculous infection of the gastrointestinal tract. mesenteric lymph nodes, peritoneum and omentum, and of solid organs related to the gastrointestinal tract such as the liver and spleen.¹

PATHOLOGICAL PRESENTATION OF ABDOMINAL TUBERCULOSIS:

I Peritoneal tuberculosis—acute or chronic

A. Tuberculosis of the peritoneum

- (i) Wet or ascitic type: Generalized, Localized (loculated)
- (ii) Dry or fibrous type: Adhesive type, Plastic type, Miliary nodular type

B. *Tuberculosis of peritoneal folds and contents viz.* Mesenteric adenitis, Mesenteric cysts, Mesenteric abscesses, Bowel adhesions and rolled-up omentum. **II. Gastrointestinal**

1. Ulcerative
2. Hypertrophic or hyper plastic
3. Sclerotic or fibrotic

III. Tuberculosis of solid viscera (e.g. liver and spleen)

Epidemiology of abdominal tuberculosis

Starting from the mid-1980s a resurgence of tuberculosis occurred; due to a large extent to the epidemic of acquired immunodeficiency disease. Intestinal tuberculosis is seen more frequently in people of poor socioeconomic circumstances. Its incidence is higher in patients with caseo-pneumonic and advanced lung disease than in those with fibrotic lesions and early disease. The frequency of extra pulmonary tuberculosis has increased, largely an influence of human immunodeficiency virus (**HIV**) infection; over 50 per cent of individuals with HIV and tuberculosis develop extra pulmonary disease compared to 10 to 15 per cent in those without HIV². Finally, there is a marked increase in the emergence of multidrug-resistant *M. Tuberculosis* organisms, a cause of much concern for the global control of tuberculosis.

Microbiology

Mycobacterium tuberculosis is responsible for nearly all cases of abdominal tuberculosis. Other pathogenic organisms such as *M. bovis* have been largely eliminated by public-health measures and are rarely encountered today. Several other atypical or anonymous Mycobacteria whose pathogenicity is not yet established have been identified.^{3,4}

Routes of infection and pathogenesis

Mycobacterium tuberculosis spreads to the abdomen by several routes. Ingestion of contaminated food may cause primary intestinal tuberculosis. Secondary intestinal disease arises from swallowed sputum containing Tuberculous bacilli; it is influenced by the virulence and quantity of the bacilli, and by host resistance to the infection. The peritoneum, mesenteric nodes, and the intestine may become infected during the bacteraemic phase that may follow primary pulmonary tuberculosis. Mycobacteria may also spread from diseased adjacent organs, such as the fallopian tubes. When the intestines become infected by lymphatic spread from the mesenteric lymph nodes, the nodal disease is considered as the primary site and intestinal involvement is secondary. This conclusion is supported by the observation that the

earliest intestinal lesions are found in the submucosal layer, while the overlying mucosa is normal. In addition, more advanced abnormalities such as caseation necrosis are found in the mesenteric nodes rather than in the intestine. The bacteria may also be disseminated in the bile, since they are sequestered and excreted from granulomas in the liver.

Sites of intestinal involvement

The terminal ileum and ileo caecal junction are involved most frequently⁵. The other regions affected in order of decreasing frequency are; colon, jejunum, rectum and anal canal, duodenum, stomach, and esophagus. The site of predilection is dictated by factors such as the abundance of lymphoid tissue, the rate of absorption of the intestinal contents, prolonged stasis, which provides longer time of contact with the mucosa, and the digestive activity of the contents.

PATHOLOGY

INTESTINAL TUBERCULOSIS

Ulcerative lesions

Tuberculous intestinal ulcers are usually deep and are transversely placed in the direction of the lymphatics. Multiple ulcers may be seen, most often in the terminal ileum⁶. Disease progression is associated with the appearance of an inflammatory mass around the bowel. The diseased part of the gut becomes thickened and the serosal surface is studded with tubercles. There is often a marked increase in mesenteric fat with fat wrapping around the bowel loops. The regional nodes become enlarged and may caseate leading to mesenteric abscess formation. Bowel perforation is rare, and is usually confined by the perilesional inflammatory mass.

Hyperplastic lesions

In the hyperplastic form of intestinal tuberculosis, a fibroblastic reaction occurs in the submucosa and subserosa resulting in marked thickening of the bowel wall; this, together with involvement of the adjacent mesentery, lymph nodes, and the omentum, results in the formation of a mass lesion⁷. Hyperplastic lesions are believed to be the result of reduced bacterial virulence and increased host resistance.

Sclerotic lesions

The sclerotic variety is associated with strictures of the intestine, typically described as 'napkin-ring strictures', which may be single or multiple. When multiple, the strictures may occur in a short segment of the bowel or over the entire length of the intestine. Enteroliths can form proximal to the stricture. In some patients a combination of the different pathological forms may be seen.

PERITONEAL TUBERCULOSIS

Acute peritonitis

Acute tuberculous peritonitis is extremely rare and is encountered under the following circumstances: in the miliary phase of the disease, on perforation of intestinal disease, and with local dissemination from a ruptured, caseating mesenteric lymph node.

Chronic peritonitis

The chronic form of Tuberculous peritonitis is much more common and typically presents as ascites . The fluid is usually clear and straw colored, but may be sanguineous. Peritoneal adhesions that range from thin and flimsy to dense and thick may occur. In the presence of adhesions the ascitic fluid may become loculated presenting as a localized cyst. The characteristic lesions are miliary nodules. When the nodules increase in size and coalesce, plastic adhesions develop, which may completely obliterate the peritoneal cavity, forming an abdominal cocoon that may encase the intestines. The omentum thickens to form a transversely placed mass, the so-called rolled-up omentum.

CLINICAL FEATURES

Abdominal tuberculosis is seen most frequently in patients between the ages of 30 and 50 years. Females outnumber males by 2:1⁸. The onset of illness is usually insidious, and the initial symptoms are often vague and non-specific; this is particularly true of peritoneal tuberculosis. As the disease progresses the individual may develop fever, which is present in two-thirds of the patients, night sweats, malaise, weakness, anorexia, and weight loss⁹. The appearance of specific symptoms depends upon the predominant site of involvement

Peritoneal tuberculosis: Tuberculous peritonitis usually presents with ascites. Less frequently, fluid collection is mild but the fibrotic component is more prominent, resulting in thickening of the peritoneum with adhesions, fluid loculation, and the classic doughy feel of the abdomen. Patients with peritoneal tuberculosis, especially women, often have coexisting tuberculosis of the pelvic organs, since the genital tract is frequently the portal of entry of the tubercle bacillus. This point is worth remembering, since a good pelvic examination and endometrial biopsy may provide the simplest method of confirming the diagnosis.

Gastrointestinal tuberculosis: The most common presentation in patients with disease of the gastro-intestinal tract is abdominal pain. The

pain may be dull and vague, but when colicky it suggests intestinal obstruction. In such patients the pain is often exacerbated by eating and relieved by vomiting, but only transiently as it soon recurs. Diarrhea is another common symptom in patients with intestinal involvement. The stools may be watery, small in amount and mixed with blood when the disease affects predominantly the colon. In primary small-bowel disease the stools are large in amount, foul smelling, and resemble those seen in patients with mal absorption. Other symptoms include flatulence, nausea, altered bowel habit, and borborygmi. Abdominal distension suggests the presence of ascites or persistent sub acute intestinal obstruction.

Physical examination

Patients with chronic abdominal tuberculosis are often malnourished and anaemic. In some the abdomen may be completely normal on examination, but most demonstrate some abnormal findings. There may be visible peristalsis and the distended bowel loops can be palpated. The abdomen may show diffuse distension and tenderness, or the signs may be more localized, usually in the right lower quadrant. An ileo caecal mass may be felt in the Right iliac fossa or higher up in the right lumbar region. A doughy' abdomen suggesting peritoneal disease has become less common in recent years. A rolled-up omentum, when

present. is felt as a transversely placed mass in the epigastric region. Loculated ascites, mesenteric cysts, and mesenteric abscesses present as cystic masses.

Patients with ascites may have shifting dullness and a fluid thrill. In patients with large-bowel disease a diffusely thickened and tender colon may be felt. Other findings include hepato-splenomegaly, pelvic abnormalities mimicking gynaecological tumours, and features of gastric-outlet obstruction due to direct involvement of the stomach or extrinsic compression of the duodenum by enlarged mesenteric lymph nodes. Rectal examination may reveal anal fistulas, fissures, or stricture.

COMPLICATIONS

Intestinal obstruction and mal absorption are the most frequent complications^{10,11}. Much less common are bowel perforation and massive gastrointestinal hemorrhage. Perforation is usually confined, owing to the presence of surrounding adhesions; as a result, signs of free perforation of the bowel are rarely seen. Acute bleeding from the rectum or haematemesis are rare, but can occasionally be severe and life threatening. Fistulas, both internal between adjacent bowel loops or with other hollow organs (uterus, vagina), and external to the skin, are described but are rare.

DIAGNOSIS

Since the discovery of the tubercle bacillus it has been possible to make a precise diagnosis of tuberculosis. However, in patients with abdominal tuberculosis the causative organism is often difficult to identify and the diagnosis is generally made by indirect methods.

Laboratory tests

The most common abnormality on routine blood tests is an elevated erythrocyte sedimentation rate, found in over 90 per cent of patients. Other abnormalities include varying degrees of anaemia and leucopenia with relative lymphocytosis. A positive tuberculin skin test (Mantoux test) is an excellent screening test in non-endemic countries, but is of little use in endemic countries because of high rates of positivity in healthy individuals and in those who have received the bacillus Calmette-Guerin (BCG) inoculation.

Analysis of ascitic fluid

The ascitic fluid has a high white blood-cell count, with a predominantly lymphocytic response. A total white-cell count of 500/mm or more has a sensitivity of 81 per cent, a specificity of 48 per cent, and a diagnostic accuracy of 46 per cent for tuberculosis. If a high count is associated with a predominantly non-poly morpho nuclear

response, the specificity and accuracy improve to 82 percent and 78 per cent, respectively. The diagnostic usefulness of the white-cell count is reduced in the presence of coexisting conditions such as cirrhosis and HIV infection, both of which are associated with an increased incidence of abdominal tuberculosis. In patients with cirrhosis and AIDS the response of white cells is poor and the total white blood-cell count is often within the normal range. Another characteristic of Tuberculous ascites is high total protein, usually 2.5 g/dl or more, and the serum ascitic fluid albumin gradient (SAAG) is less than 1.1. However, high total protein is seen in several conditions, and the sensitivity, specificity, and diagnostic accuracy of this test for tuberculosis are only 65, 78, and 74 per cent, respectively. As with the white blood-cell count, the protein content of ascitic fluid in cirrhotic patients with peritoneal tuberculosis is significantly lower; the concentration is less than 2.5 g/dl in 30 to 50 per cent of patients, and the SAAG is highly variable. Other biochemical tests, such as lactate dehydrogenase (elevated to over 90 units/l), low pH, and ascitic fluid: blood glucose ratio of less than 0.96, are usually positive in Tuberculous ascites, but these tests are non-specific and of little diagnostic value¹².

Adenosine Deaminase:

Lately, adenosine deaminase (**ADA**) in ascitic fluid has received much attention. ADA is an enzyme present in several cell types including macrophages, lymphocytes, and erythrocytes. Its concentration in body fluids correlates with the number and degree of stimulation of lymphocytes, and is a marker of host immune response. Several studies have shown that ADA activity is an extremely useful diagnostic test for Tuberculous ascites, with specificity and sensitivity of over 95 per cent. However, in the presence of cirrhosis, seen in more than 50 per cent of patients with peritoneal tuberculosis in the West, the sensitivity of ADA drops to 30 per cent; this is related to the abnormal T-lymphocyte activation and proliferation in those with cirrhosis. For similar reasons, ADA activity is likely to be less useful in HIV-positive patients, limiting the effectiveness of this test in subgroups of patients who are highly susceptible to tuberculosis. However, in those without these predisposing conditions, ADA is extremely useful and should be a first-line investigation, obviating the need for more invasive and expensive diagnostic tests^{13,14,15}. Another test that may prove useful is assay of interferon (produced by T-lymphocytes) in the ascitic fluid¹⁶.

Bacterial isolation and culture:

Positive identification of *M. tuberculosis* provides the definitive diagnosis of tuberculosis. Acid-fast smears prepared from ascitic fluid have an extremely low yield, with positive results in fewer than 5 per cent of patients. Culture of ascitic fluid for *M. tuberculosis* is more useful and is positive in 20 to 45 per cent of patients. An important disadvantage of culture is that the results take 4 to 6 weeks¹⁷, which severely limits the clinical usefulness of the test. While awaiting the results of culture, anti-tuberculosis treatment should be started, which can be modified later based on drug-sensitivity findings.

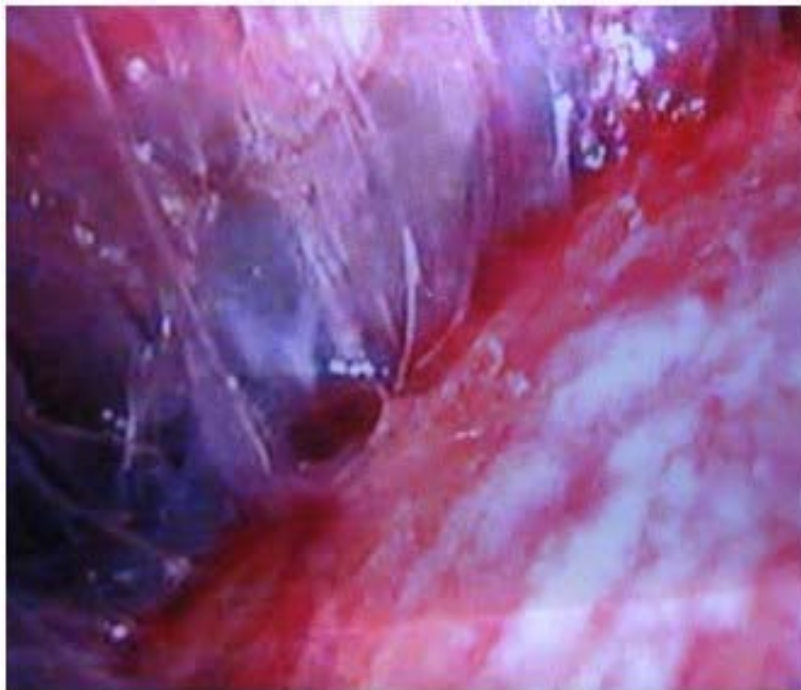
Laparoscopy and peritoneal biopsy:

Blind needle biopsy of the peritoneum can be performed in the presence of ascites using special needles (Abrams' or Cope's). Most workers note a low complication rate, but fatality has been reported. The main risk of blind biopsy is bowel perforation, particularly in patients with adhesions between bowel loops and the anterior abdominal wall. Open biopsy of parietal peritoneum under local anaesthesia is much safer. A less traumatic and more useful approach is to obtain targeted biopsy specimens from diseased areas such as

enlarged lymph nodes under ultra sonographic or computed tomographic (CT) guidance. The single most sensitive diagnostic test is laparoscopic examination of the peritoneum¹⁸. The peritoneal lining loses its smooth, glistening appearance and becomes rough, irregular and dull. The characteristic finding, as described above, is the presence of miliary tubercles. In addition, fibrous adhesions ranging from tiny filaments to thick bands may be seen. Ascitic fluid, if present, should be sent for biochemical analysis and culture. Targeted biopsy specimens should be obtained, preferably.



LAPAROSCOPY – PERITONEAL TUBERCLES & ASCITES



LAPAROSCOPY – ADHESIVETUBERCULOSIS

from the miliary tubercles. Peritoneal biopsy should be performed even if peritoneal nodules are not seen; histological examination shows caseating granulomas in nearly 90 per cent of patients. The combination of the distinctive visual and histological appearances accurately identifies nearly all patients. However, positive identification of *M. tuberculosis* histologically or by culture is made in only one-half of the patients. The complication rate of laparoscopy is low (less than 5 percent) but care should be taken in the presence of adhesions. This procedure is especially rewarding in patients with AIDS and cirrhosis, because of a much broader differential diagnosis of ascites and the poor sensitivity of the other tests.

Imaging studies

Plain radiographs:

Plain X-ray may show calcification of the mesenteric lymph nodes and calcified granulomas in the spleen, liver or pancreas, but these findings do not imply active disease. Patients in intestinal obstruction may show dilated bowel loops and air-fluid levels. An abnormal chest radiograph indicating pulmonary tuberculosis helps in the differential diagnosis but is seen in fewer than 50 percent of patients with gastrointestinal disease^{19,20}.

Ultrasonography:

Several ultrasonographic findings have been described. The peritoneum assumes a thickened, irregular, echo-poor, sheet-like or nodular appearance. Ultrasound is very sensitive in detecting small quantities of fluid. Because of their small size, peritoneal nodules are rarely detected, but are better visualized in the presence of ascites and appear as tiny, echo-poor deposits²¹. A characteristic finding is the presence of alternating echogenic and echo-free layers produced by the bowel wall, the serosa, and the adjacent bowel loop with interloop fluid collections; this is termed the 'club sandwich' appearance. In addition, isolated or matted enlarged lymph nodes, with hypoechoic or anechoic centres caused by caseation necrosis, may be seen.

Computed tomography:

Similar findings as noted on ultrasound, but with better definition and resolution, are seen with the CT scan¹⁷. The ascitic fluid has high-density appearances because of its elevated protein content. The thickening and nodularity of the peritoneum and mesentery can be more easily identified with CT than ultrasonography. In patients with intestinal disease there is thickening of the bowel wall and of the ileocaecal valve. The ultrasound and CT findings, although fairly

characteristic, are not pathognomonic of abdominal tuberculosis. Similar appearances can be seen in other diseases such as lymphoma, metastatic carcinoma, peritoneal mesothelioma, and pseudomyxoma peritonei.



CT ABDOMEN – ILEOCAECAL TUBERCULOSIS



CT ABDOMEN – MESENTRIC ADENITIS

Barium studies:

Barium studies provide useful information on the extent and severity of the intestinal disease. The earliest abnormalities include altered bowel motility, and irregularity and thickening of the mucosal folds. Barium swallow may show compression of the oesophagus by a mediastinal node, with or without mucosal ulceration. In patients with more advanced disease, mucosal ulcers, deformity of the bowel lumen, and stricture formation are noted. Rarely, sinus tracts and fistulas may be seen. Thickening of the ileocaecal valve, a wide-open valve accompanied by narrowing of the terminal ileum (Fleischner sign)¹⁹, and a fibrotic terminal ileum opening into a contracted caecum (Sterlin sign) are characteristic of intestinal tuberculosis.

Enteroclysis (small-bowel enema) is

generally considered the ideal method of assessing the small bowel. This technique involves passing a tube into the distal duodenum, which greatly reduces enthusiasm for the test, both by the radiologist and the patient. Most radiologists therefore prefer the use of small-bowel series. The regular 20 per cent wt/vol barium mixture has a high density, which often prevents proper examination of overlapping bowel loops. Low-density barium preparations (13 per cent wt/vol) containing methylcellulose have been introduced in the belief that

they permit better dispersion of a barium that is less prone to flocculation and has greater trans radiancy, allowing a 'see-through' effect with good visualization of overlying loops. Whether these preparations are indeed better remains to be confirmed. For the large bowel the study of choice is air-contrast barium enema.



BARIUM MEAL STUDY - ILEAL STRICTURE



BARIUM ENEMA - PULLED UP CAECUM

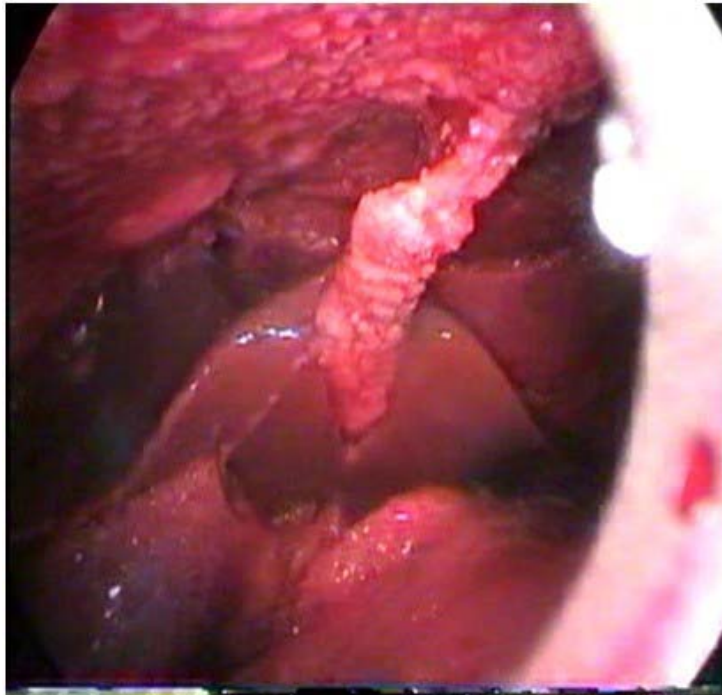
Overall, barium studies have a low sensitivity and are abnormal in only about 60 per cent of patients with endoscopically proven disease. A rapid transit and lack of barium retention may occur in inflamed segments of the small bowel, resulting in false-negative results. Moreover, the barium abnormalities are non-specific and can be mimicked by other conditions such as Crohn's disease and lymphoma.

Endoscopy with biopsy

Fibreoptic endoscopes have made it possible to visualize directly the gastrointestinal tract. The ileocaecal region can be accessed easily by colonoscopy, and the use of enteroscopes allows examination of the proximal small bowel. The usual endoscopic findings are mucosal ulcerations, nodularity, deformity, narrowing, and stricture of the bowel. Rarely, there is diffuse disease of the colon with hyperaemia and friability of the mucosa, mimicking ulcerative colitis. The endoscopic abnormalities in tuberculosis can be mistaken for Crohn's disease. Ulcers seen in tuberculosis are usually transversely located and have sharply defined margins with the surrounding mucosa showing erythema, while in



COLONOSCOPY – ULCERATIVE TUBERCULOSIS



LAPAROSCOPY – PERITONEAL TUBERCLES

Crohn's disease the ulcers are serpiginous, often longitudinal, with a relatively normal surrounding mucosa²²⁻²⁴.

Biopsy specimens obtained at endoscopy show granulomas and epithelioid cells in 40 to 74 per cent of patients with intestinal tuberculosis. However, confirmatory findings of caseation necrosis are present in only 8 to 21 per cent, acid-fast bacilli are observed infrequently, and positive cultures are obtained in only 6 to 40 per cent of patients. Although endoscopy provides a definitive diagnosis in only about one-third of the patients, it is very useful in excluding other conditions such as lymphoma, carcinoma, and caecal amoeboma.

Serological studies

Because of the difficulty in identifying *M. tuberculosis* in tissue material, a sensitive and specific serological test should be very helpful. Infection with *M. tuberculosis* stimulates a cell-mediated as well as a humoral immune response. The tuberculin skin test is a manifestation of the cell-mediated immune response. Serological tests based on the detection of specific antibodies to *M. tuberculosis* have been under investigation for several years. *Mycobacterium tuberculosis* has a

complex structure and contains numerous antigens that cross-react with each other as well as with antigens of other mycobacterial species. Enzyme-linked immunosorbent assays based on the use of one of several bacterial antigenic preparations are commercially available, but they provide sensitivity, specificity, and diagnostic accuracy of only 80 per cent, and moreover are unable to identify clearly²⁵ (a) active disease from dormant infection, (b) *M. tuberculosis* infection from other mycobacterial infections, and (c) individuals with previous BCG inoculation.

Polymerase chain reaction

The low positive identification rate for *M. tuberculosis* in intestinal tuberculosis is related to the presence of small numbers of organisms in the tissues. By amplifying tiny quantities of DNA and RNA, the polymerase chain reaction is thus ideally suited for this condition. The technique has been used in a variety of clinical specimens including sputum, cerebrospinal fluid, pleural and peritoneal fluids, and biopsy tissues, with sensitivity, specificity and positive predictivity of 85, 99, and 95 per cent, respectively. Based on studies with pulmonary secretions, the polymerase chain reaction can detect as few as 50 organisms per reaction, which is at least fivefold below the

lower limit of detection by culture. The technique has also been successfully used on endoscopically obtained biopsy specimens but much more work is required before it can be recommended for routine use.^{26,27}

TREATMENT

Medical treatment

The treatment of abdominal tuberculosis is primarily medical. Since acid-fast bacilli and caseation necrosis are seen in a minority of patients, and culture results take several weeks, chemotherapy plays very important role in the treatment of abdominal tuberculosis.²⁸

Short-term therapy:

The current guidelines put forward by the World Health Organization and the United States Centers for Disease Control recommends a much shorter course of therapy. The initial treatment consists of a three-drug regimen of isoniazid (300 mg), Rifampicin (450 mg), and Pyrazinamide (1.5 g) in countries where the resistance rate to Isoniazid is below 4 per cent. In countries with higher drug- resistance rates, a fourth drug is added, Ethambutol (25 mg/kg) or streptomycin (1 g). All drugs are administered daily for 2 months, after which the patient is switched to two drugs; Isoniazid 300mg/day and Rifampicin 450 mg/day for 4 months. Facilities for directly observed therapy are available, initial treatment is the same as in short-term therapy, but after 2 months patients are started on a twice-a-week regimen consisting of

Isoniazid (600 mg/day) and Rifampicin (600 mg/day) for 4 months
Thus, the total length of treatment for both short-term and directly
observed therapies is 6 months.

Although these guidelines are based on studies in pulmonary tuberculosis, there is no evidence that patients with abdominal tuberculosis require longer therapy. Patients should be monitored every 4 to 6 weeks for drug-related side-effects. Isoniazid should be discontinued if the liver enzymes (alanine and aspartate aminotransferases) increase threefold over baseline. It is advisable to administer pyridoxine hydrochloride (5-10 mg/day) to all patients to prevent isoniazid-induced peripheral neuropathy. If a coexisting open pulmonary lesion is present, the patient should be isolated for 2 weeks. If not, the patient may be treated at home.

The clinical response to treatment is excellent. Systemic symptoms such as fever, malaise, and weight loss subside in a few weeks. Mucosal abnormalities take longer, but follow-up barium studies and endoscopy demonstrate regression of the lesions in most individuals. The majority of patients (70%) with symptoms of subacute bowel obstruction and evidence of intestinal stricture show complete resolution of the radiological abnormality.²⁸

Anti tuberculosis-drug resistance

Multidrug-resistant *M tuberculosis* received little attention until the early 1990s when several outbreaks were reported in patients with HIV. In 1994, the World Health Organization initiated global surveillance for antituberculosis-drug resistance in 35 countries and results of the first 4 years (1994-1997) of this project were recently published. Among patients with no prior treatment, a median of 9.9 per cent (range 2-41 per cent) were resistant to at least one drug and multidrug resistance was seen in 1.4 per cent (0-14.4 per cent). The resistance rates to individual drugs were: Isoniazid 7.3 per cent, streptomycin 6.5 per cent, Rifampicin 1.8 per cent, and Ethambutol 1 per cent. Much higher rates of drug resistance were observed in patients with history of previous anti tuberculous treatment.

Drug resistance is a major threat to tuberculosis control programmes worldwide. Patients with resistant strains are less likely to be cured, and the treatment is more expensive and more toxic than in those with susceptible organisms. Multidrug resistance occurs more frequently in non-compliant patients, in the presence of HIV infection, and in malnourished individuals. The current emphasis is to create more

facilities for directly observed therapy. Studies show that unsupervised treatment has a drug-completion rate of only 25 to 50 per cent, while directly observed therapy results in 85 to 90 per cent cure, with relapse rates of only 5 percent.

ROLE OF SURGERY

Indications:

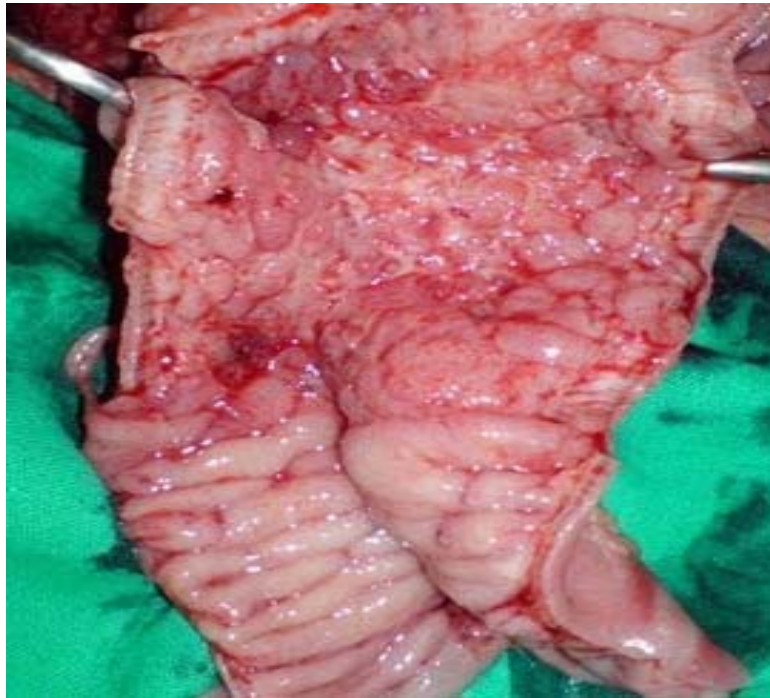
Surgery is reserved for mechanical complications of tuberculosis or when medical therapy fails. Emergency surgery is indicated in the presence of acute complications such as free perforation of the bowel and severe intestinal haemorrhage. The most common indication for surgery is intestinal obstruction secondary to stricture formation. In these patients, unless there is complete obstruction, conservative management is advocated. If symptoms persist, elective surgery is performed at a later stage, with significantly less morbidity. Predictors for surgical intervention are long strictures (12 cm or more in length) and multiple areas of involvement. Other indications for surgery are bowel adhesions, intra- abdominal abscess due to a confined perforation, mesenteric abscess, and internal or external fistulas. Surgery is also appropriate if the diagnosis is in doubt and when malignancy cannot be ruled out with reasonable accuracy.^{15, 29,30}

Surgical procedures:

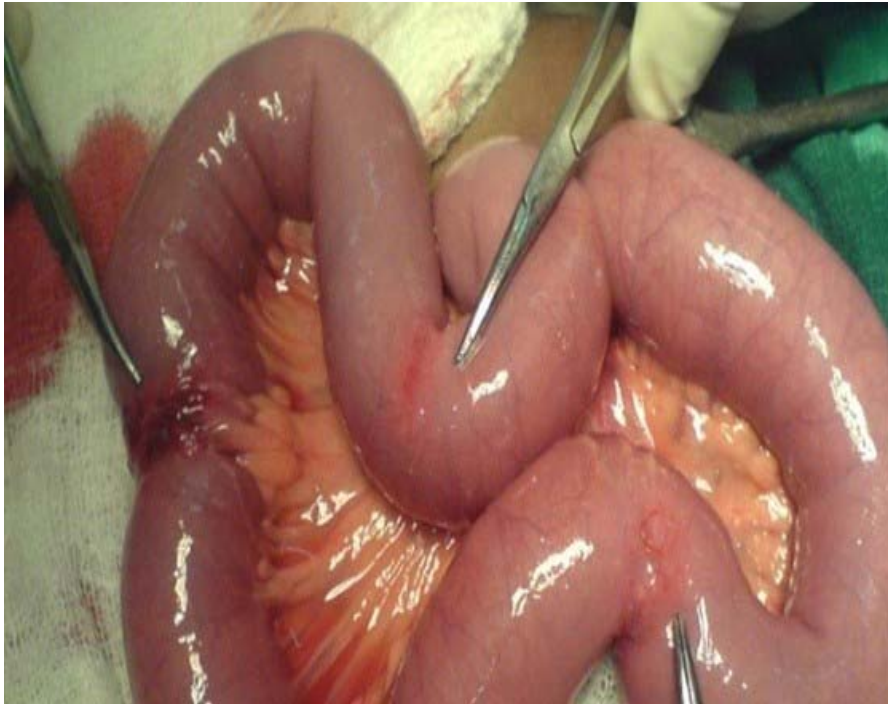
Patients presenting with acute bowel perforation should be treated by resection of the involved segment and primary anastomosis. Simple closure of perforation is followed by a high incidence of re perforation and fistula formation. Drainage tubes are not recommended and, if used, should be removed early



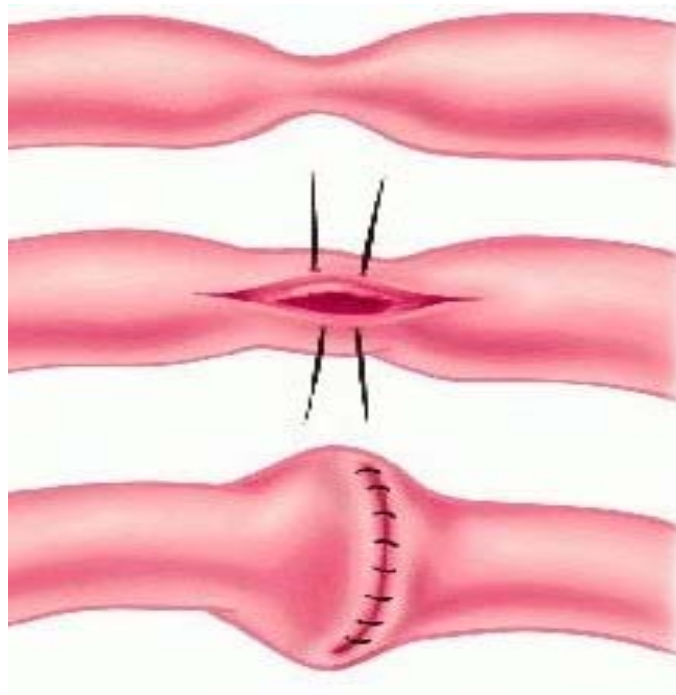
MESENTRIC ADENITIS



ULCERATIVE TUBERCULOSIS



TUBERCULOUS ILEAL STRICTURE



STRICTUROPLASTY

because of increased risk of abdomino cutaneous fistulas if they are left in place for more than 7 days.

Surgical techniques for patients with bowel obstruction have evolved over time. At one time, bypass procedures such as ileotransverse colostomy and enteroenterostomy were practiced commonly, but have now been abandoned because of complications such as blind-loop syndrome, mal absorption, and perforation. Similarly, radical bowel resection and extensive dissection of mesenteric lymph nodes have fallen out of favour because of complications. including the development of short-bowel syndrome. The order of the day is minimal surgery. The procedure of choice for an ileocaecal mass is limited resection with 5-cm margins from visibly abnormal tissue in place of the standard right hemicolectomy. Segmental bowel resection has become rare and is only performed if multiple strictures are located in close proximity. Most surgeons prefer stricturoplasty for lesions involving the small bowel. Other procedures advocated are ileocaecoplasty and coloplasty for ileocaecal and colonic strictures. respectively.³¹⁻³³

Approach to intestinal adhesions:

Intestinal adhesions require a meticulous surgical approach. In addition to releasing the adhesions, enteric stenting has been found useful in preventing adhesive bowel obstruction. Patients with florid sepsis may be treated by laparostomy, followed by secondary abdominal closure once the sepsis is controlled; this prevents recurrent adhesions and reduces postoperative morbidity and mortality. Patients with enterocutaneous fistulas are treated by resection of the fistulous tract, drainage of any intra-abdominal abscess, and intestinal anastomosis. A multicentre, prospective, randomized trial has shown that the interposition of a barrier film for a sufficient length of time between diseased areas of intestine contributes to adhesion-free healing. Seprafilm and Interceed are a few of the commercially available barrier films proved to be effective. Some of these patients may need stenting in addition to barrier-film placement.

SPECIFIC FORMS OF TUBERCULOSIS

Appendiceal:

Tuberculosis of the appendix is reported in 0.1 to 3 per cent of patients with tuberculosis. Isolated tuberculosis of the appendix is

rare. Appendectomy followed by antituberculosis chemotherapy is the treatment of choice.

Anal canal:

Patients with tuberculosis of the anal canal may present with multiple fistulas, fissures, or perianal abscesses. Treatment consists of drainage of the abscess combined with antituberculous chemotherapy.³⁴⁻³⁶

Gastric:

Involvement of the stomach is rare, occurring in 0.3 to 2.3 per cent of patients with pulmonary tuberculosis. The rarity of this location is related to several factors including the resistance of the gastric mucosa to infection, the lack of lymphoid tissue in the stomach, and the constant flow of contents through the stomach. Tuberculosis of the stomach presents as an ulcerative, granulomatous, or fibrosing lesion; the last two types may result in gastric-outlet obstruction. At endoscopy, the lesion may be mistaken for peptic ulcer disease, gastric carcinoma, or gastric sarcoidosis. Similar appearances are also seen in syphilis of the stomach. The diagnosis may be confirmed on endoscopic biopsy specimens.

Antituberculous chemotherapy should be initiated in all patients; this is curative in most, especially those with ulcerative lesions, which can be confirmed on repeat endoscopy and biopsy. Surgical intervention is required if gastric-outlet obstruction persists despite treatment. The usual surgical approach is a partial gastric resection such as a Billroth gastrectomy or a sleeve resection. Gastric cancer presents in a similar fashion, and both tuberculosis and cancer may coexist in the same patient. Therefore, if there is any doubt about the diagnosis, it is best to proceed with frozen-section biopsy followed by surgical resection if indicated.³⁷⁻⁴³

Liver and spleen:

Hepatic tuberculosis has become exceedingly rare these days. The diagnosis is usually made accidentally during exploratory laparotomy or at autopsy in immunocompromised patients. The lesions typically are granulomas, with or without central caseating necrosis, calcified masses, and biliary strictures. Tuberculous periportal lymph nodes may cause obstructive jaundice by compressing the bile duct. Patients with hepatic tuberculosis usually have hepatomegaly, with or without jaundice. Symptoms related to abdominal tuberculosis often overshadow those due to liver disease. Liver enzymes, in particular

serum alkaline phosphatase, are usually elevated. Tuberculosis should be differentiated from other conditions associated with hepatic granulomas such as leprosy, sarcoidosis, Hodgkin disease, brucellosis, infectious mononucleosis, inflammatory bowel disease, drug-induced liver damage, and syphilis. Chronic active hepatitis may also mimic tuberculosis of the liver. The treatment of hepatic tuberculosis is chemotherapy. It should be remembered that most antituberculous drugs (except ethambutol) are hepatotoxic, and may aggravate the liver damage and worsen the jaundice. These patients therefore should be kept under close observation during anti tuberculous chemotherapy.

Splenic tuberculosis is also rare and may present as a splenic abscess or with hypersplenism. The presence of multiple hypoechoic lesions on ultrasonography of the spleen in a HIV-positive patient is highly suggestive of disseminated tuberculosis. The diagnosis is usually made following surgical resection of the diseased spleen.

'Silent' tuberculosis:

Subclinical or 'silent abdominal tuberculosis is seen mainly in patients belonging to upper socioeconomic groupings in the Third World countries. The onset of illness is often precipitated by stressful

conditions. The patients present with atypical symptoms, resulting in frequent misdiagnosis. The response to short- term antituberculous drug therapy is excellent.

Abdominal Tuberculosis in childhood: Abdominal tuberculosis in children is seen most frequently in Immunosuppressed individuals and in those who have not been vaccinated with BCG. Clinically. weight loss and malaise occur in 95 per cent of these children, followed by abdominal distension (83 per cent), abdominal pain (79 per cent), and anaemia (76 per cent). Laboratory tests show leucocytosis and an altered albuminglobulin ratio; pulmonary tuberculosis is seen on the chest radiograph. Imaging studies, including plain radiographs, ultrasonography, CT and gastrointestinal contrast studies, are helpful in localizing the defect. The diagnosis is confirmed on tissue obtained by endoscopic biopsy or with echo- or CT-guided fine-needle aspiration; samples may be stained by the Ziehl-Nielsen method for acid-fast bacilli, subjected to microbiological culture and drug- sensitivity testing, and examined histopathologically. A short course of chemotherapy is an effective and economical treatment, and is associated with minimal side-effects in children.

HIV and tuberculosis:

Abdominal tuberculosis in HIV-infected patients is invariably a manifestation of disseminated disease and results in significant mortality. Extrapulmonary tuberculosis is seen in over 50 per cent of patients. Fever, weight loss, lymphadenopathy, and splenic abscesses are seen more commonly in HIV- infected patients than in those without HIV infection. The diagnostic techniques are the same as in uninfected individuals and include bacteriological testing of body fluids, abdominal ultrasound and CT scanning combined with guided-needle aspiration biopsies, barium examination, fibreoptic endoscopy, and laparoscopy. Serological tests such as enzyme-linked immunosorbent assay lack sensitivity due to a poor humoral immune response. Most patients respond well to conventional antituberculous drugs used in standard doses. Some experts recommend a longer course of therapy (9 months). Despite treatment, a few patients experience a downhill course, which may be due to drug resistance or the presence of overwhelming infection.

PROGNOSIS

The prognosis of uncomplicated abdominal tuberculosis is good. Most patients respond well to medical therapy and the outcome of surgical management of complications is generally good. Bowel obstruction or perforation associated with plastic adhesions and the development of enterocutaneous fistulas with intra-abdominal abscesses are associated with increased morbidity and poor prognosis.

GENERAL RECOMMENDATIONS

The main advance in the diagnostic approach to patients with suspected abdominal tuberculosis is the use of endoscopy. Most patients with peritoneal tuberculosis can be diagnosed by laparoscopy, since the peritoneal appearances are fairly distinctive and the histological yield is high. If this facility is not available, adenosine deaminase in ascitic fluid should be assessed because of its high positive predictive value for tuberculosis, especially in patients without coexisting cirrhosis or HIV infection. In contrast to peritoneal tuberculosis, the diagnosis of gastrointestinal tuberculosis is more difficult. All endoscopically accessible lesions should be biopsied and the specimens used for histology, staining for acid-fast

bacilli, and culture. Even if a positive diagnosis of tuberculosis is not made, diseases such as carcinoma and lymphoma can be excluded. The next step in endemic countries is a therapeutic trial with antituberculous drugs. The majority of patients, even those with strictures, show a good response. In the West, where tuberculosis has made a major comeback, the main differential diagnosis is with Crohn's disease, and the clinical, radiological, and histological features of the two are often indistinguishable. Here, a specific diagnostic test for *M. tuberculosis* such as polymerase chain reaction, may prove extremely helpful.

AIMS & OBJECTIVES OF THE STUDY

1. To find out demographic profile of abdominal tuberculosis
2. To study the clinical presentations of abdominal tuberculosis.
3. To evaluate the diagnostic modalities of abdominal tuberculosis.
4. To discuss the medical and surgical modalities of treatment of abdominal tuberculosis

MATERIALS AND METHODS

STUDY AREA:

INSTITUTE OF GENERAL SURGERY, MADRAS MEDICAL
COLLEGE, CHENNAI.

SAMPLE SIZE:

50 CASES OF ABDOMINAL TUBERCULOSIS

STUDY DESIGN:

DESCRIPTIVE STUDY

STUDY PERIOD:

FEB 2017 TO SEP 2017

METHOD OF SAMPLING:

CONVENIENT SAMPLING

INCLUSION CRITERIA:

ALL PATIENTS WHO WERE DIAGNOSED ABDOMINAL
TUBERCULOSIS BASED ON RADIOLOGICAL/LAPAROSCOPIC
EVIDENCE WITH OR WITHOUT HISTOPATHOLOGICAL
EVIDENCE...

EXCLUSION CRITERIA:

ALL PATIENTS IN PAEDIATRIC AGE GROUP.

METHODOLOGY

This descriptive study was done for a period of 6 Months from Feb 2017 to Sep 2017 in Government General Hospital, Madras Medical College, Chennai , India. The Proforma (Annexure) contains the following details demography , presentation, lab reports, radiological reports and management of these patients ,which were recorded after the consent of the patients. Collected datas were analysed.

RESULTS

The total number of cases who were diagnosed to be of abdominal tuberculosis in our study period was 50 patients. The male and females in our study group include 33 and 17 respectively. The sex ratio was 1.9:1 between males and females. (Table 1, Fig 1). There was a higher incidence of abdominal tuberculosis in the age group between 31-40 years of age in both the genders followed by the patients between 21-30 years of age. (Table 2, Fig 2). The clinical manifestations varying from Acute intestinal obstruction to non specific illness, with acute intestinal obstruction was a initial clinical presentation in 12 of the patients, abdominal distension in 15 patients, mass in the abdomen in 12 patients, abdominal pain and non specific symptoms in 11 patients. (Table 3, Fig 3).

All the patients underwent routine laboratory investigations like total blood count and Erythrocyte sedimentation rate (ESR). Lymphocytosis was found in 60% of patients and raised ESR was found in 90 % of the patients in the study group. The ascitic fluid analysis which was done in 15 patients who presented with ascites showed the biochemical and microbiological parameters of tuberculosis in 10

patients. (Table 4, Fig 4). Three patients showed positive serology for HIV.

All patients underwent Roentgenogram of chest which showed features of active and healed pulmonary lesions, pleural effusion in one sixth of the patients with sensitivity of 17% for diagnosis of tuberculosis. X-ray of abdomen done in all patients who showed features of intestinal obstruction, perforative peritonitis, and calcified lymph nodes. Ultrasonographic evaluation of abdomen which was done in all patients revealed features suggestive of tuberculosis in 23 patients (47.2%). The computed tomography (CT) of abdomen was done in 7 patients suspected to be having features of abdominal tuberculosis clinically and ultrasonography did not provide a conclusive diagnosis. Also CT of abdomen was done in 28 other patients who underwent ultrasonographic examination which did not provide a conclusive diagnosis or other diagnosis for further diagnostic evaluation. CT done in total of 38 patients concluded to be of abdominal tuberculosis in 29 patients. Barium meal and Barium enema done in 10 and 7 patients respectively for small bowel and large bowel lesions when indicated showed a sensitivity of 78% and 70% respectively. (Table 5, Fig 5). Colonoscopic examination done in 8 patients when large bowel lesions were suspected revealed tuberculous lesions in 2 patients. Diagnostic

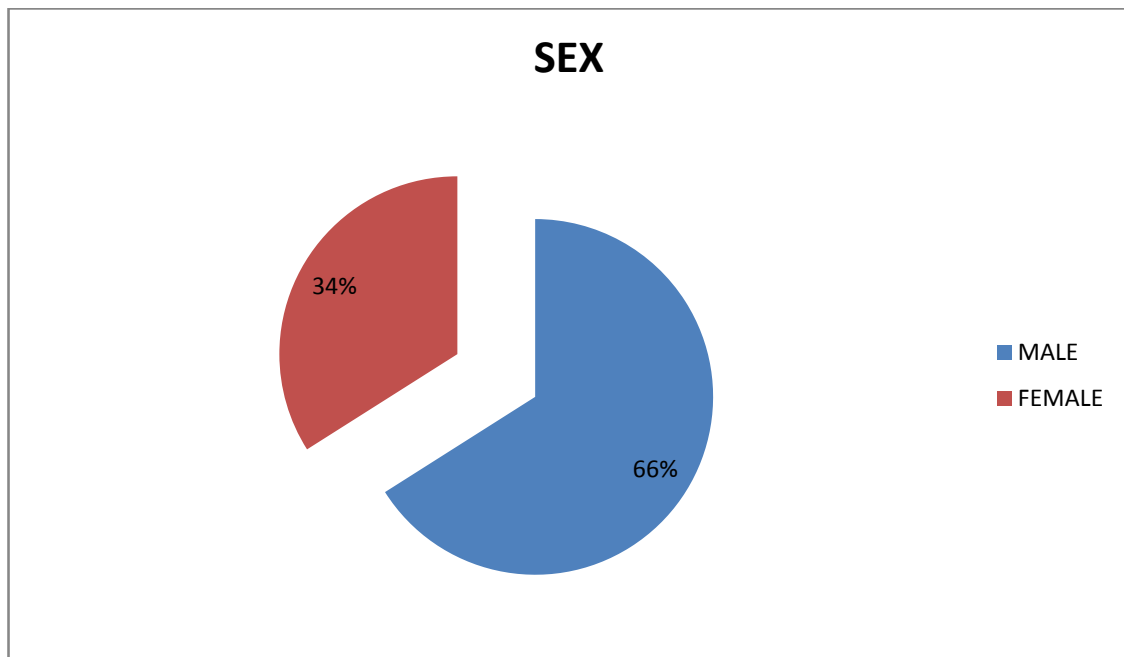
laparoscopy was done in 24 patients of our study group which included the patients who did not have conclusive diagnosis after radiological diagnosis and also patients with radiological investigations supporting abdominal tuberculosis for further confirmatory evidence and tissue diagnosis. It showed a sensitivity of 87 % in our study group.

Of the 50 patients in the study group, only 23 patients needed surgical interventions whereas 27 patients were managed with chemotherapeutic drugs. Emergency surgical management was done in total of 8 patients which include laparotomy and adhesiolysis in 2 patients, stricturoplasty in 2 patients, limited resection in 2 patients, and perforation closure and covering ileostomy in the other 2 patients. 15 patients had undergone management electively after complete investigatory evaluation, the surgical procedures include right hemicolectomy in 6 patients, adhesiolysis in 2 patients, stricturoplasty in 3 patients, diversion ileostomy in 1 patient, adhesiolysis and abscess drainage through laparoscopy in 3 patients, sealed perforation closure which presented with interloop abscess in a case.. Of the 50 patients only 37 had histological evidence of abdominal tuberculosis. Only 1 patient out of 50 having intra abdominal tuberculosis had associated perianal fistulae with a incidence of 2%..

TABLE 1: SEX DISTRIBUTION OF ABDOMINAL TUBERCULOSIS IN OUR STUDY GROUP

Sex	Frequency	Percent
Male	33	66.0
Female	17	34.0
Total	50	100.0

Fig 1: Sex Distribution of abdominal tuberculosis in the study group

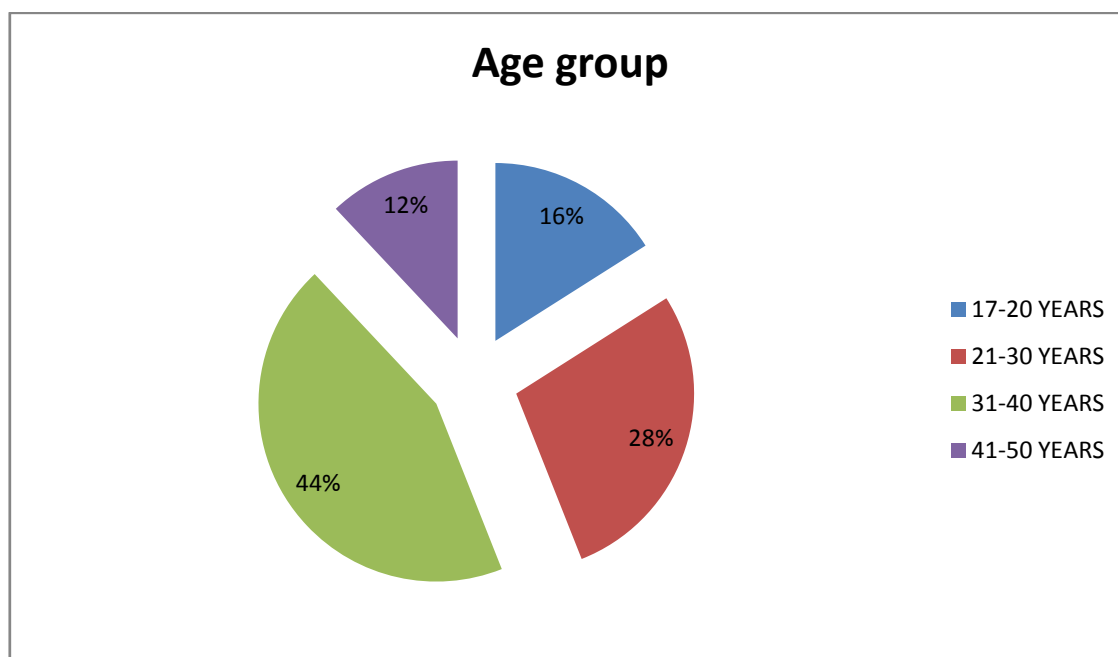


Male: Female ratio 1.9:1

TABLE 2: AGE DISTRIBUTION AMONG THE GENDERS IN OUR STUDY GROUP:

Age group	Frequency	Percent
17-20 Years	8	16.0
21-30 Years	14	28.0
31-40 Years	22	44.0
41-50 Years	6	12.0
Total	50	100.0

Fig 2: AGE DISTRIBUTION AMONG THE GENDERS IN OUR STUDY GROUP



**TABLE3: CLINICAL PRESENTATION OF
ABDOMINAL TUBERCULOSIS - OUR STUDY**

Presentation	Frequency	Percent
Abdominal mass	12	24.0
Abdominal pain	6	12.0
Ascites	15	30.0
Intestinal Obstruction	12	24.0
Non specific symptoms	5	10.0
Total	50	100.0

Fig 3: Clinical presentation of abdominal tuberculosis in our study

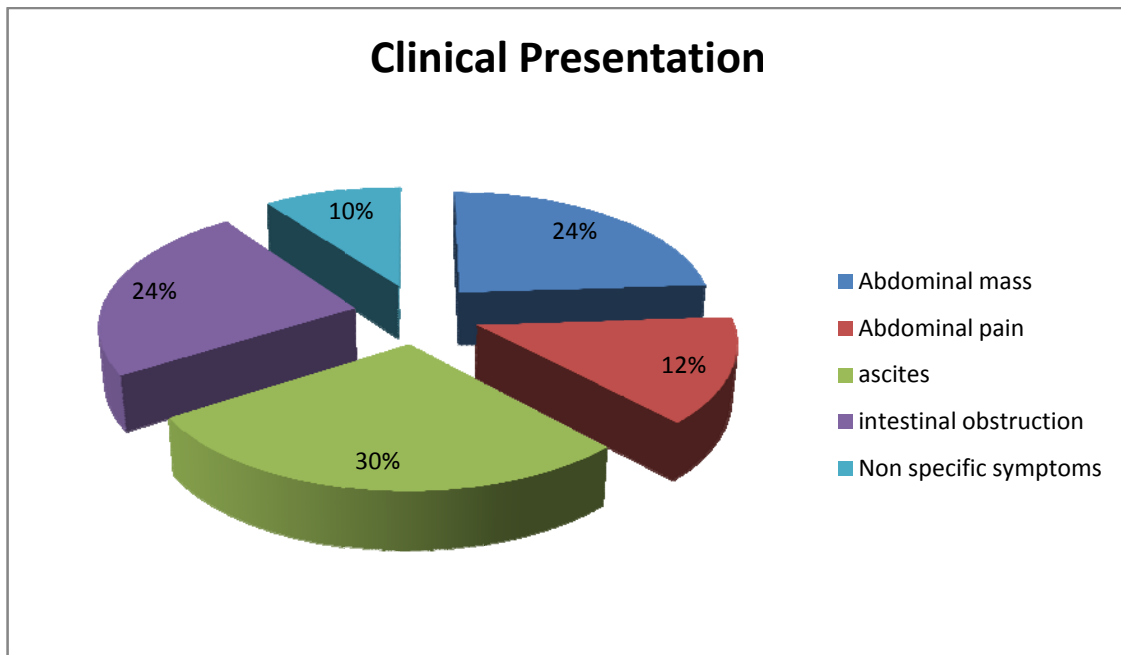


TABLE 4: LABORATORY INVESTIGATIONS DONE IN OUR PATIENTS AND ITS SENSITIVITY

Lymphocytes	Frequency	Percent
Absent	20	40.0
Present	30	60.0
Total	50	100.0

Fig 4: Sensitivity of Laboratory investigations.

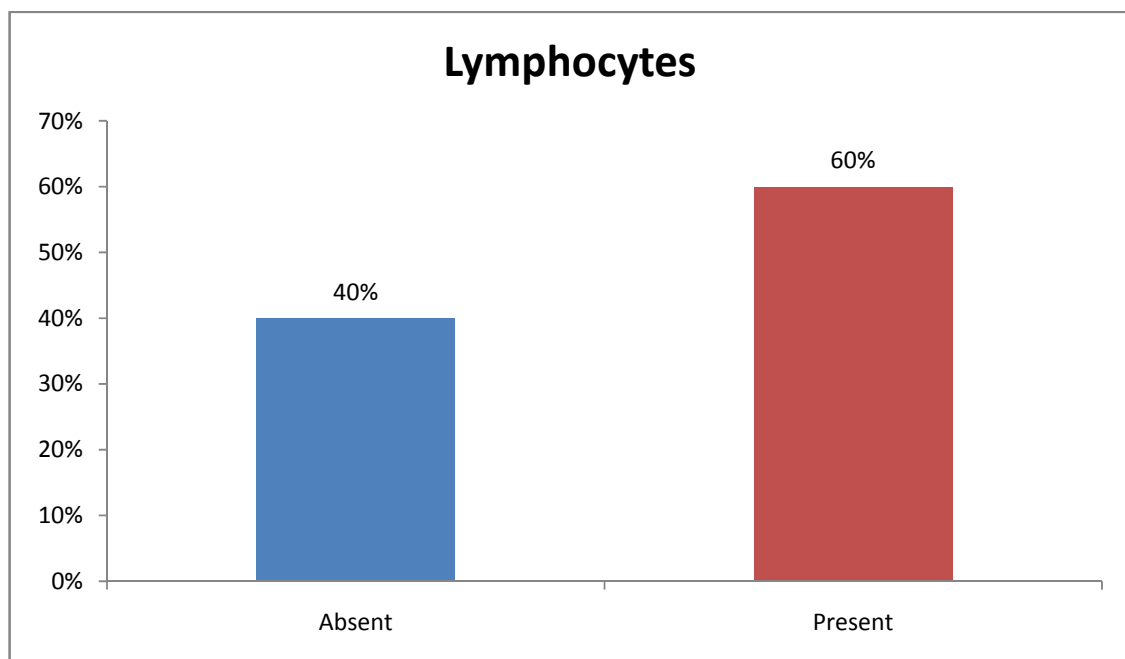


TABLE 5: ESR

ESR	Frequency	Percent
Normal	5	10.0
Raised	45	90.0
Total	50	100.0

Fig 5: Sensitivity of ESR

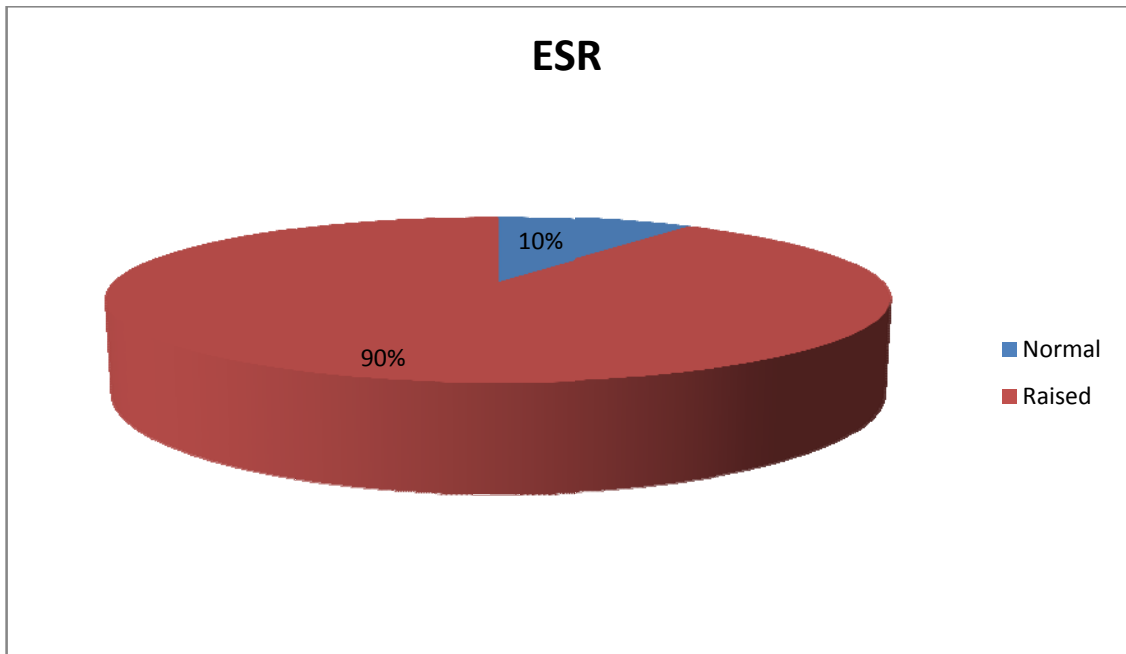
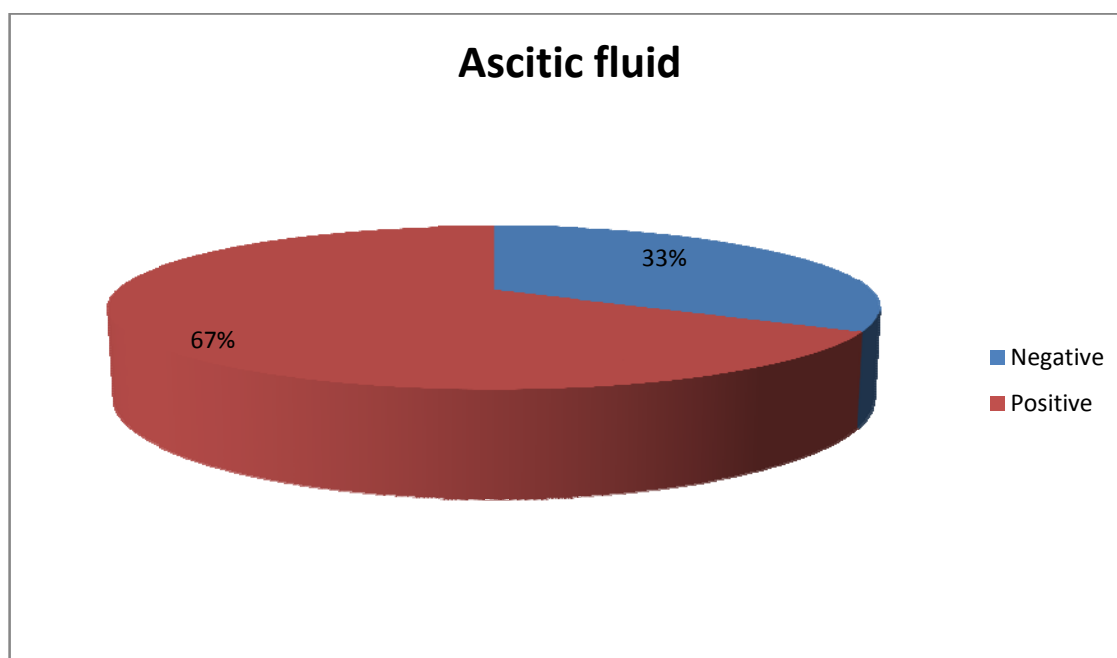


TABLE: 6 – ASCITIC FLUID ANALYSIS

Ascitic fluid	Frequency	Percent
Negative	5	33%
Positive	10	67%
Total	15	100.0

Fig 6: Sensitivity of ascitic fluid



**TABLE 7: RADIOLOGICAL INVESTIGATIONS
DONE IN OUR PATIENTS.**

Chest x ray	Frequency	Percent
Normal	41	82.0
Positive	9	18.0
Total	50	100.0

Fig 7: Frequency of x-ray changes in lung

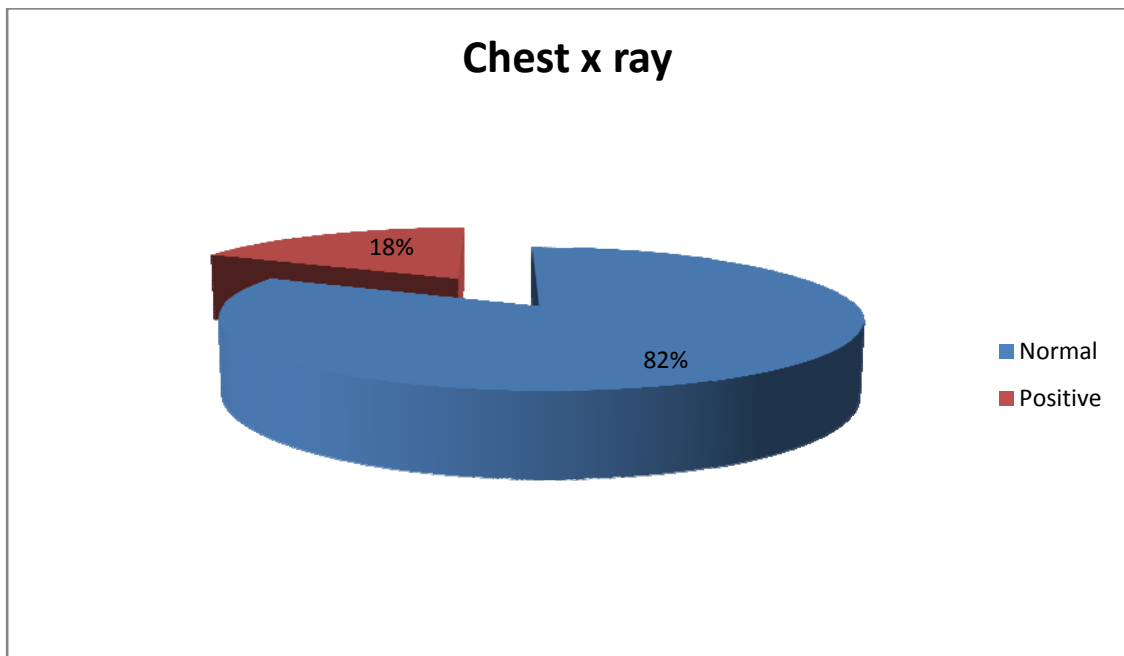


TABLE 8: BARIUM MEAL STUDY

Barium meal	Frequency	Percent
Negative	3	30.0
Positive	7	70.0
Total	10	100.0

Fig 8: Sensitivity of Barium Meal Study

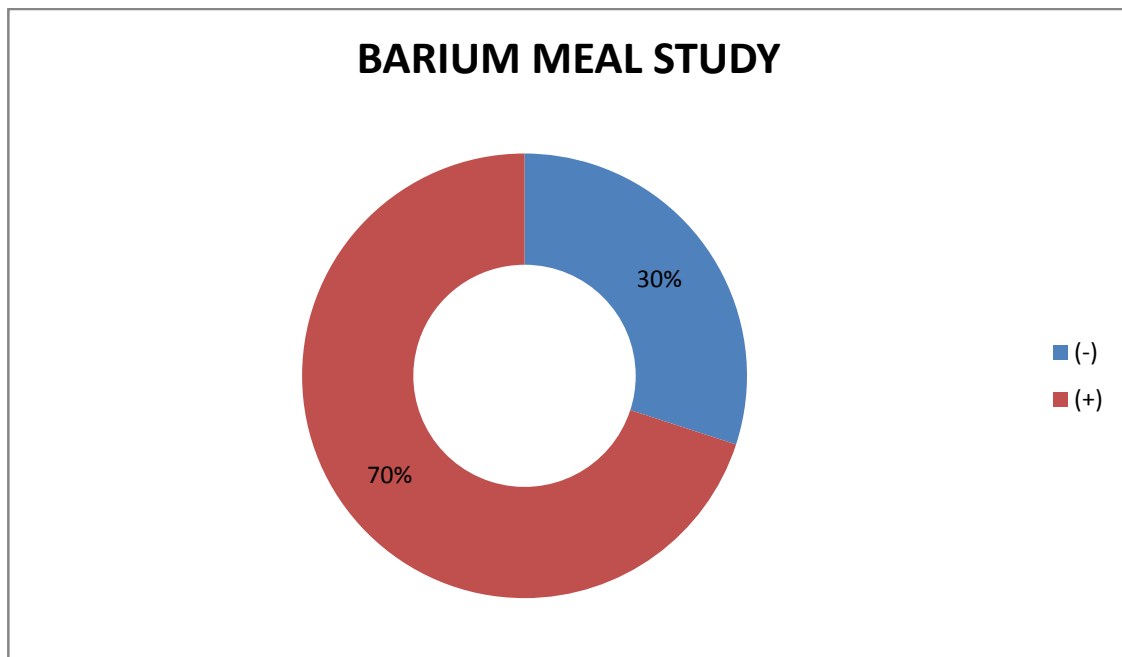


TABLE 9: BARIUM ENEMA STUDY

Barium enema	Frequency	Percent
Negative	2	40.0
Positive	3	60.0
Total	5	100.0

Fig 9: Sensitivity of Barium Enema Study

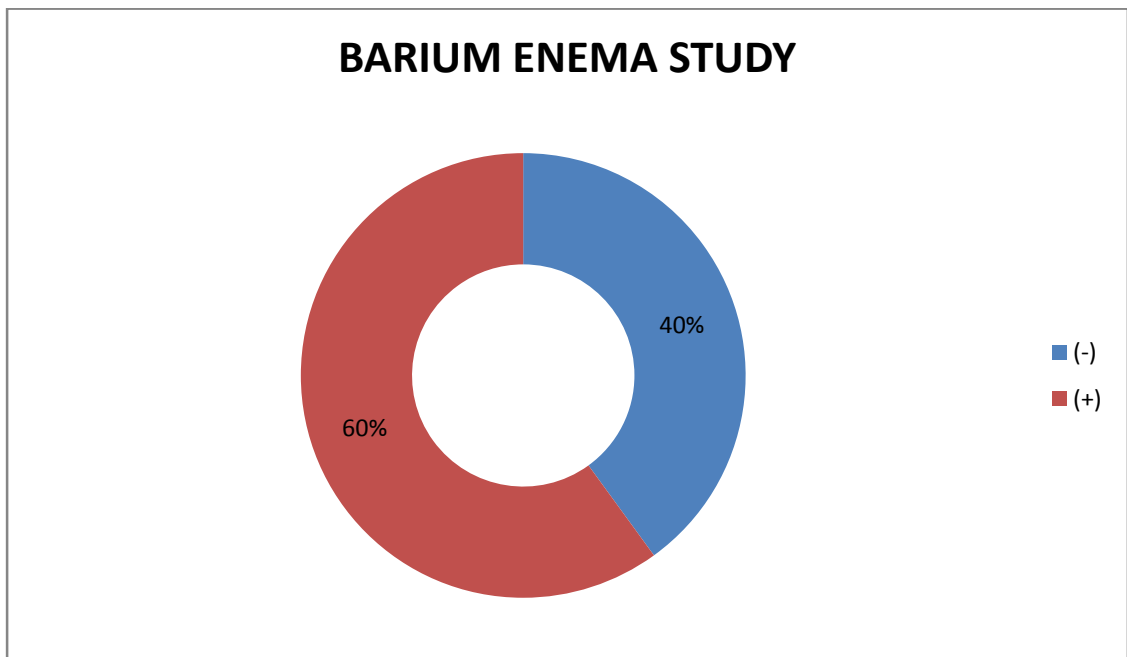


TABLE 10: CT ABDOMEN

CT ABD	FREQUENCY	PERCENT
Negative	9	22.0
Positive	29	78.00
Total	38	100.0

Fig 10 : Sensitivity of CT abdomen

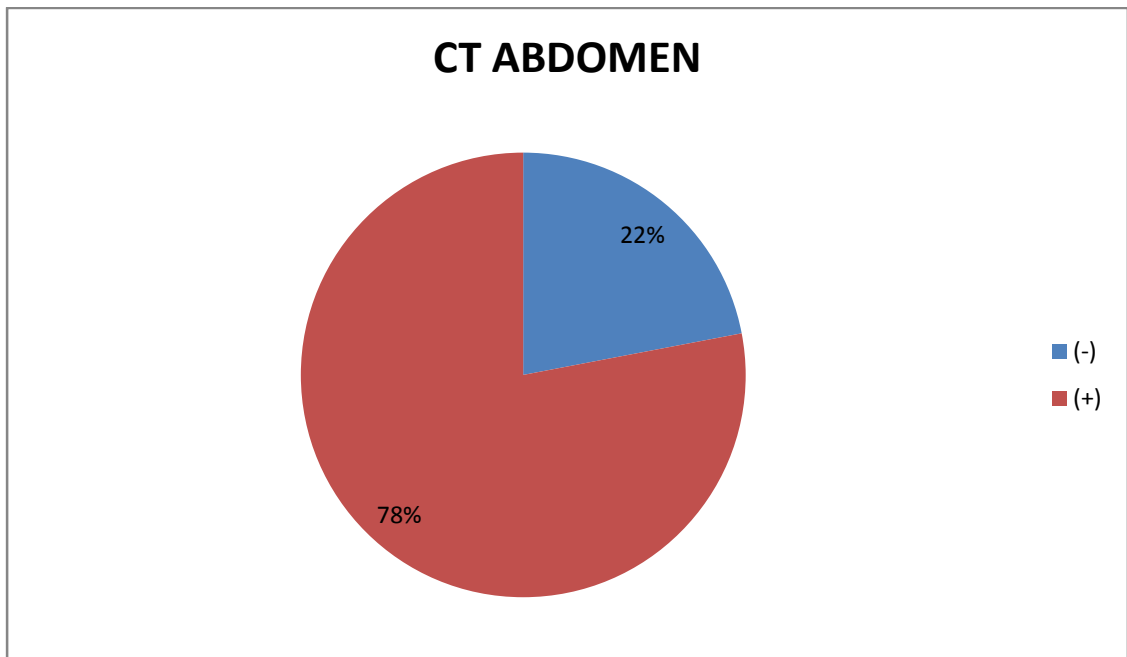


TABLE 11 : DIAGNOSTIC LAPAROSCOPY

Diagnostic laparoscopy	Frequency	Percent
Negative	3	13.0
Positive	21	87.0
Total	24	100.0

Fig: 11 : Sensitivity of Diagnostic Laparoscopy

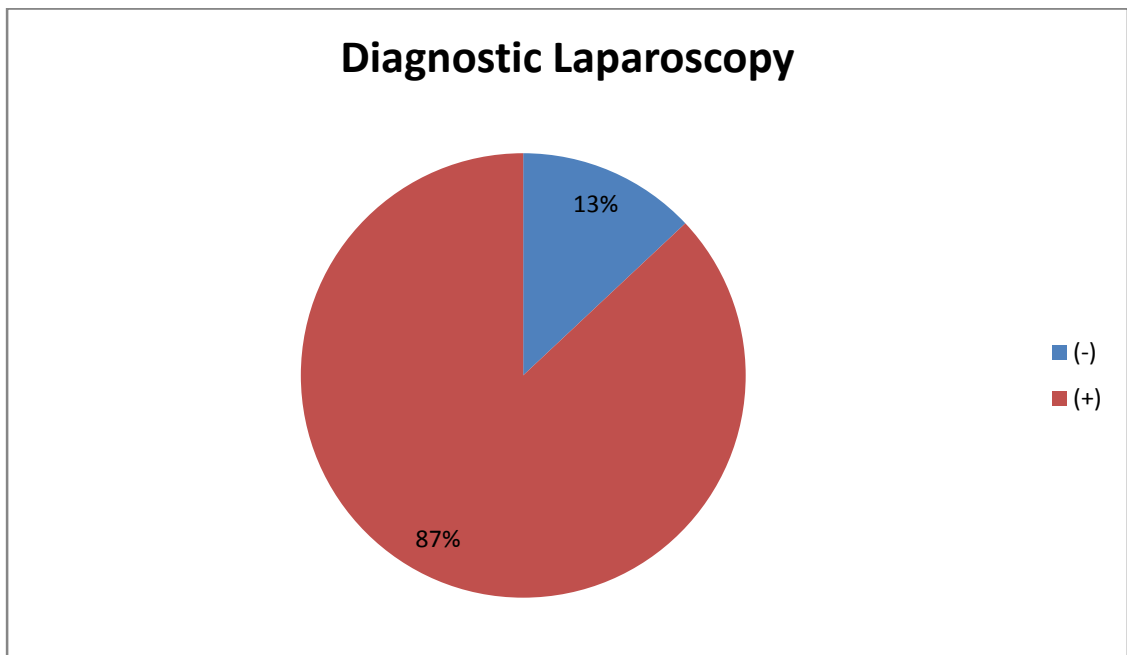


TABLE 12 : SENSITIVITY OF COLONOSCOPY AND DIAGNOSTIC LAPAROSCOPY

S. No.	Investigations	Total No. of patients underwent	Positive	Sensitivity
1.	Colonoscopy	8	2	25%
2	Laparoscopy	24	21	87%

Fig 12: Sensitivity of colonoscopy and D-laparoscopy

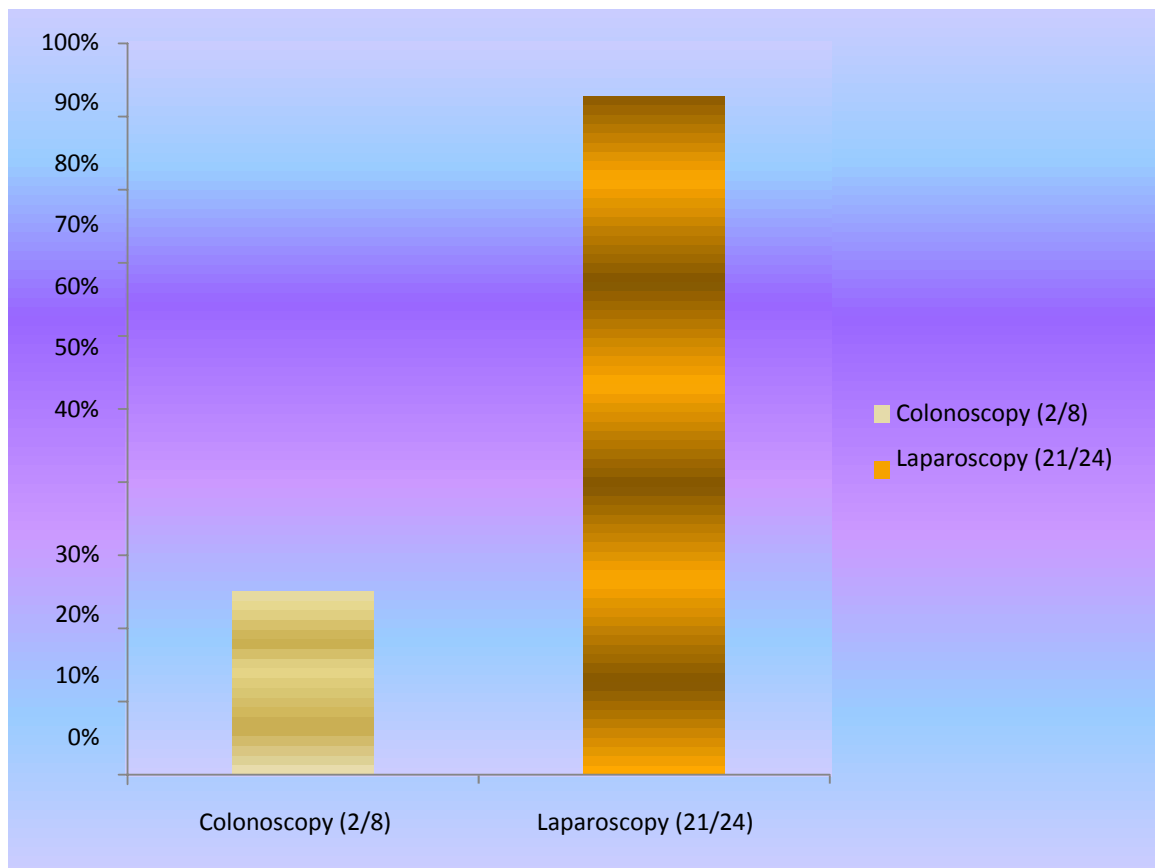


TABLE 13: DIFFERENT DIAGNOSIS IN OUR STUDY

Diagnosis	Frequency	Percent
ILEALPERFORATION	2	4.0
ILEALSTRICTURE	7	14.0
ILEOCAECALT.B	6	12.0
INTERLOOPADHESIONS	8	16.0
MESENTRICLYMPADENITIS	6	12.0
T.B.ABDOMEN	10	20.0
TUBERCULOUSASCITES	11	22.0
Total	50	100.0

Fig 13: Frequency of different diagnosis

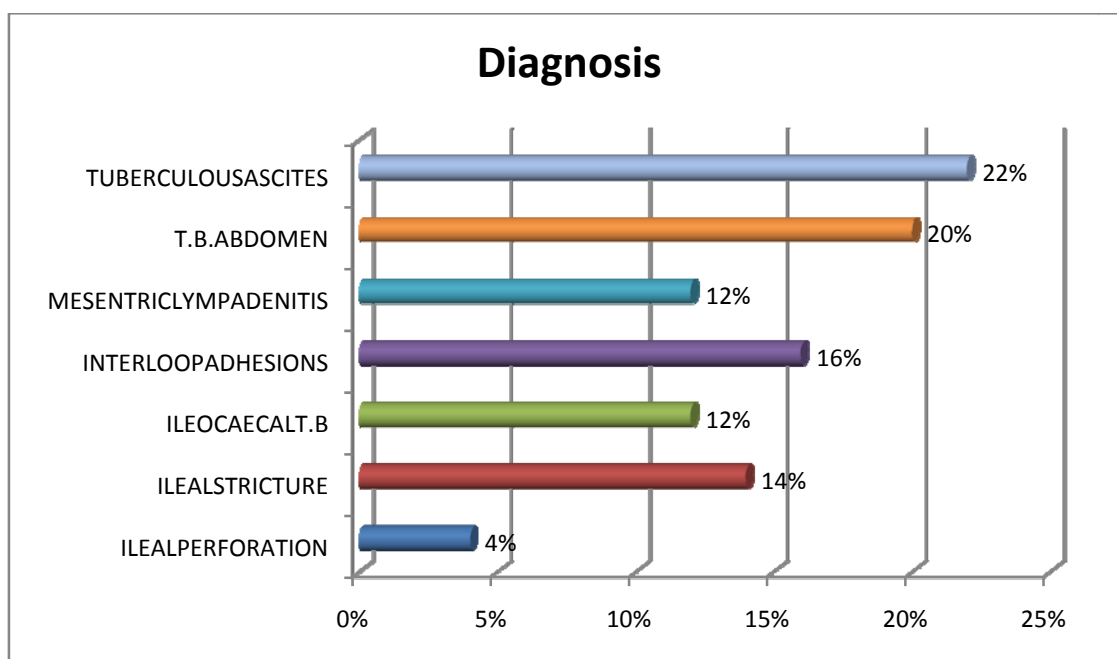


TABLE 14: COMPILATION OF RADIOLOGICAL INVESTIGATIONS IN OUR STUDY

INVESTIGATION	NO.OF PATIENTS UNDERWENT	NUMBER OF POSITIVE STUDY	SENSITIVITY
BARIUM MEAL	10	07	70%
BARIUM ENEMA	05	03	60%
CT ABDOMEN	38	29	77%
DIAGNOSTIC LAPAROSCOPY	24	21	87%

TABLE 15: MANAGEMENT OF ABDOMINAL TUBERCULOSIS IN OUR PATIENTS.

TREATMENT	Frequency	Percent
MEDICAL MANAGEMENT	27	54.0
SURGICAL MANAGEMENT	23	46.0
Total	50	100.0

Fig 15: Management of abdominal tuberculosis in our patients.

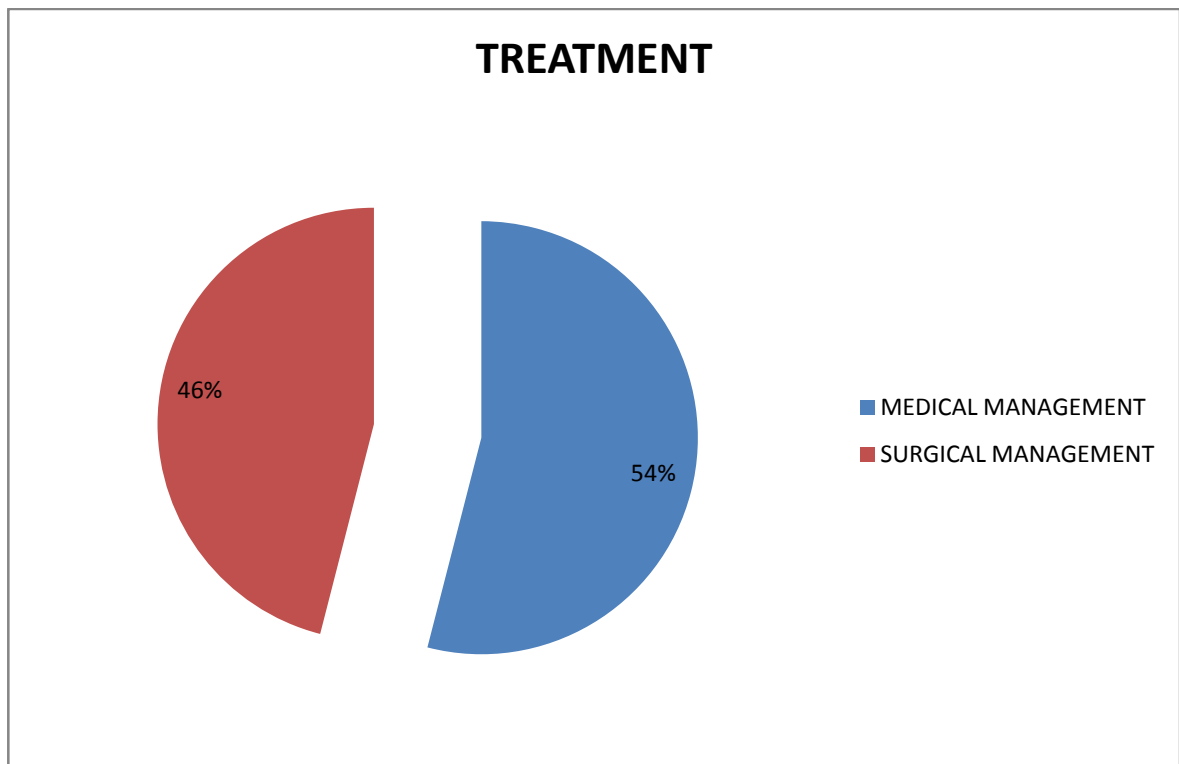


TABLE 16: TIME OF SURGICAL MANAGEMENT

Timing	Frequency	Percent
Elective	15	65.0
Emergency	8	35.0
Total	23	100.0

Fig:16 Time of surgical management in our patients

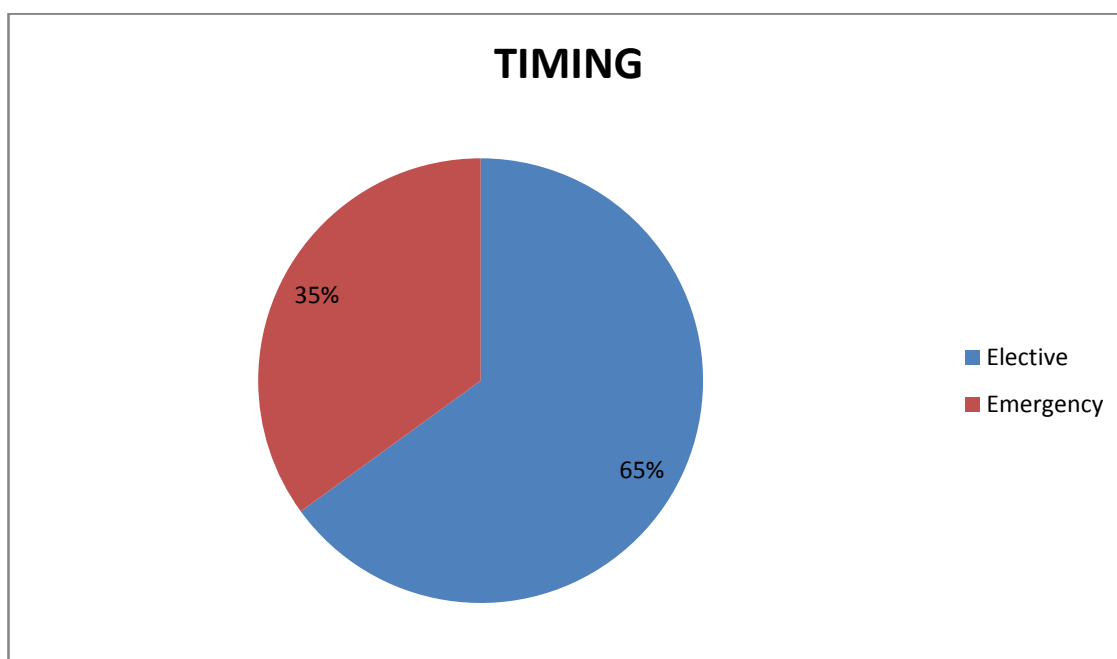
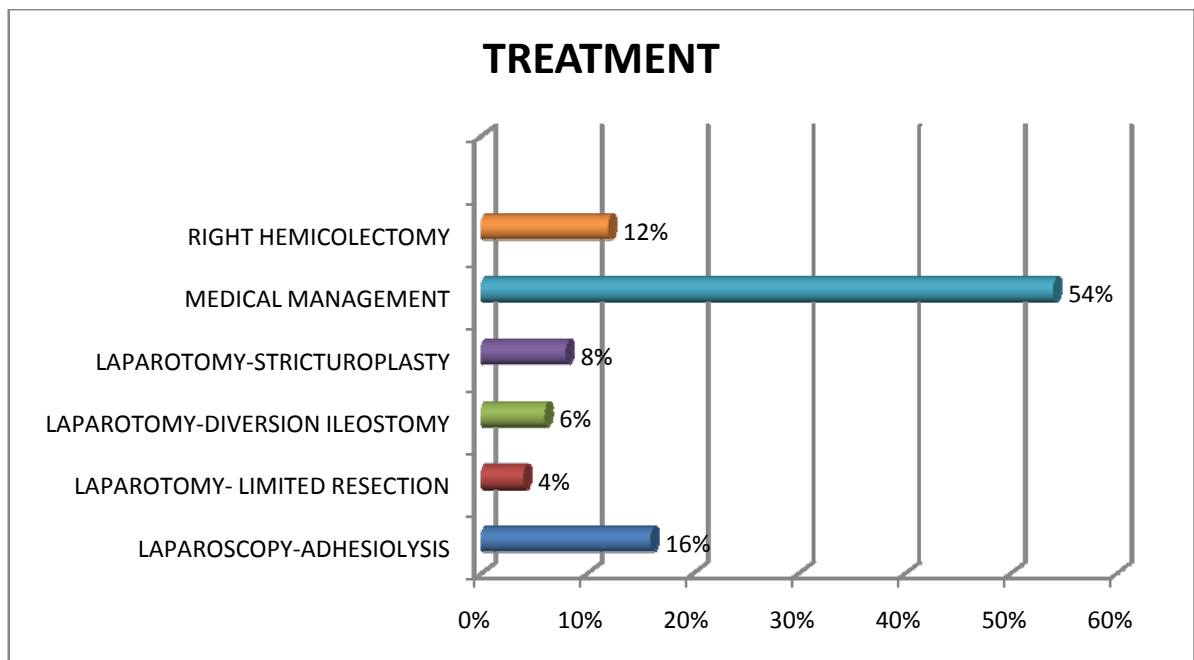


TABLE 17: DETAILS OF SURGICAL PROCEDURE DONE FOR ABDOMINAL TUBERCULOSIS IN OUR PATIENTS

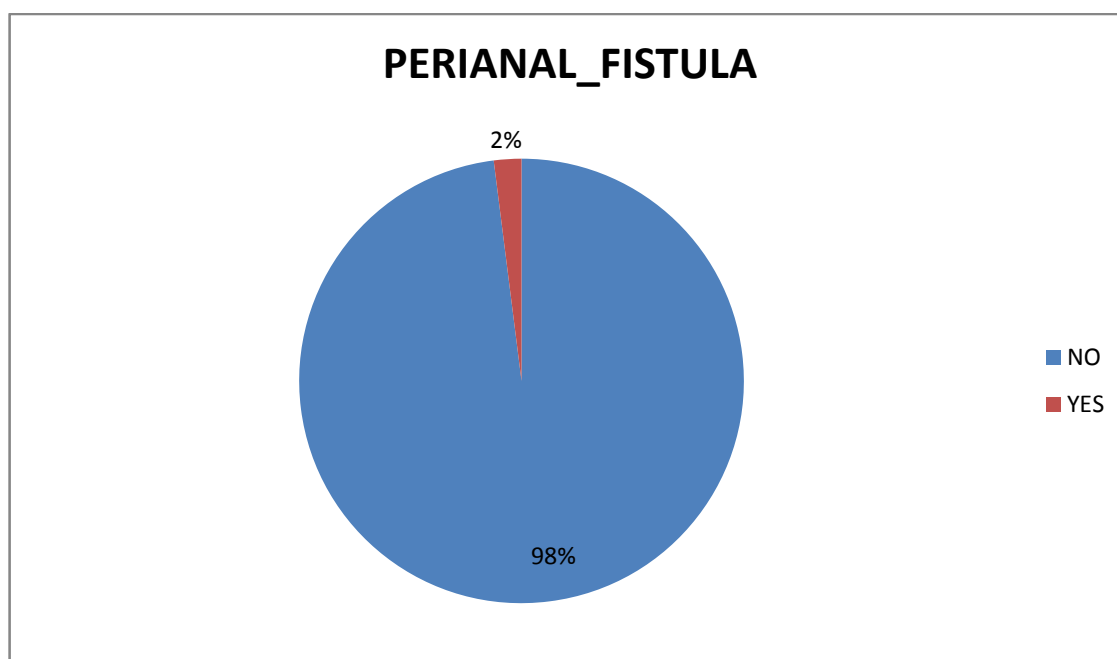
Treatment	Frequency	Percent
Laparoscopy-Adhesiolysis	8	16.0
Laparotomy- LimitedResection	2	4.0
Laparotomy-DiversionIleostomy	3	6.0
Laparotomy-Strictureplasty	4	8.0
Medical Management	27	54.0
Right Hemicolectomy	6	12.0
Total	50	100.0



**TABLE 18: INCIDENCE OF PERIANAL FISTULA IN
ABDOMINAL TUBERCULOSIS**

Perianal Fistula	Frequency	Percent
No	49	98.0
Yes	1	2.0
Total	50	100.0

**Figure 18: Incidence of Perianal Fistulae in Abdominal
Tuberculosis patients**



DISCUSSION

In our study commonest age group affected was 2nd to 4th decade of life. Age incidence of present series is similar to reported by other workers [Adams and Miller, 1946, Dutta Guptha 1950, Sharma et al ,1972 B.K Bhansali,1968]. Pritam Das and Shukla .et.al 1978 , also reported same age group. The male and females in our study group include 33 and 17 respectively. The sex ratio was 1.9:1 between males and females Addison et al 1981 reported high incidence in males.

The clinical manifestations varying from Acute intestinal obstruction to non specific illness with acute intestinal obstruction and abdominal pain was a initial clinical presentation in twelve of the patients, abdominal distension in 15 patients, mass in the abdomen in 12 patients, and non specific symptoms in 11 patients. Badaoui E , BerneyT , Kaiser L et al 2000,also reported similar presentation.

M P Sharma and Vikram Bhatia stated in their review article about the clinical presentation dominated by constitutional symptoms. 40-70% of patients present with fever, 80-95% with pain, 11-20% with diarrhea, 40 -90% with weight loss. N.Rangabashyam.et.al cited vague symptoms to be the predominant presenting symptoms. The abdominal pain accounts for in 77-94% of patients, followed by

vomiting and abdominal distension. Majority of patients in the present study had present with chronic symptoms. 8 out of 50 patients had presented with an acute abdomen and underwent emergency surgery for the same. J.M. Findly 1981 found that acute presentation was present in 13.5% patients. Commonest symptom is vague abdominal pain .Other common symptoms were vomiting, fever, loss of appetite and loss of weight. Pritam Das and Sukula had reported abdominal pain in 94% of cases Bockus et al 1964 emphasized abdominal pain , anorexia ,loss of weight as common presenting symptoms. Shukula 1970 and L.E Hugas also reported these symptoms as common presentation

Any patient present with vague abdominal pain, vomiting , loss of appetite , loss of weight and not fitting with any other clinical diagnosis, diagnosis of abdominal tuberculosis should be considered.

Routine laboratory tests have limited value in the diagnosis of abdominal TB. Although the mantoux test is freely available, its value in the diagnosis active TB remains uncertain. Lymphocytosis was found in 60% of patients and raised ESR was found in 90 % of the patients in the study group.

RADIOLOGICAL STUDIES

Chest X-ray:

Evidence of tuberculosis in a chest Roentgenogram supports the diagnosis of abdominal tuberculosis but a normal chest X-ray does not rule it out. Sharma *et al* studied 70 cases of abdominal tuberculosis and found evidence of active or healed lesions on chest X-ray in 22 (46%). X-rays were more likely to be positive in patients with acute complications (80%). In Prakash's series of 300 patients, none had active pulmonary tuberculosis but 39 per cent had evidence of healed tuberculosis. Tandon *et al* found chest X-ray to be positive in only 25 per cent of their patients. Hence, about 75 per cent cases do not have evidence of concomitant pulmonary disease. Plain X-ray abdomen may show enteroliths, features of obstruction *i.e.*, dilated bowel loops with multiple air fluid levels, evidence of ascites, perforation or intussusception. In addition, there may be calcified lymph nodes, calcified granulomas and hepatosplenomegaly.

In our study group, out of all 50 patients, of chest x ray in 9 patients (18%) ,6 patients showed features of healed pulmonary tuberculosis whereas active pulmonary lesions found in 3 patients in contrast to the above literatures. 82% of patients in our study group did

not have any evidence of concomitant pulmonary tuberculosis. Chest X-ray gives evidence of old or active pulmonary tuberculosis but rather they provide the indirect suspicion of abdominal tuberculosis.

12 patients in our study group out of 50, had evidence of features of intestinal obstruction, perforation, and ascites. None of the features in abdominal x-ray gave diagnostic evidence of abdominal tuberculosis but rather they provide the features of various presentations of abdominal tuberculosis. X ray abdomen though plays an important role in evaluation, but do not provide the diagnostic conclusion of abdominal tuberculosis.

Small bowel barium meal and large bowel barium enema: The features which may be seen: Accelerated intestinal transit; hyper segmentation of the barium column (“chicken intestine”), precipitation, flocculation and dilution of the barium; stiffened and thickened folds; luminal stenosis with smooth but stiff contours (“hour glass stenosis”), multiple strictures with segmental dilatation of bowel loops, may also be found; and fixity and matting of bowel loops. Barium meal was done in 10 of our patients which revealed the features of abdominal tuberculosis in 7 patients. Barium meal studies do have limitations in its diagnosis

and application for the patients and cannot be used in all cases but has its significance in cases of strictures with proximal dilation. Barium enema with radiological diagnostic features as explained earlier was found in 2 of our patients but the features were nonspecific. Barium enema gives supportive evidence to the diagnosis. Barium studies though accurate for intrinsic bowel abnormalities, do not detect lesions in the peritoneum. Enteroclysis followed by a barium enema may be the best protocol for evaluation of intestinal tuberculosis and that is to be evaluated further.

Ultrasonography

Ultrasonography is a cheap and valuable investigation in the evaluation of abdominal tuberculosis when the clinical features have suspicion of abdominal tuberculosis. They are particularly useful in cases of abdominal mass (Cocoon), lymph nodes, ascites and dilated bowel loops. USG was found to be 47.2% of patients in our study group whereas other study groups from Uvqur et al showed sensitivity of 57% and Dobok G et al viewed a sensitivity of 54.2%. Ultrasonography is a non expensive diagnostic tool which can be used as initial radiological diagnostic modality.

Computed tomography:

CT of abdomen has high sensitivity than USG for diagnosis of cases with diffuse lymphadenopathy, low density lymph node with multi locular appearance following intravenous contrast, High density ascites, and mottled low density masses in the omentum thickening of the Bowel wall adjacent to mesentery and also with hepatic pseudo tumors. CT abdomen has a sensitivity of 71% in 35 patients in which CT was done which was marginally comparable to the observations of Dobak G et al who experienced a sensitivity of 69%.

Colonoscopy

Colonoscopy is an excellent tool to diagnose colonic and terminal ileal involvement but is still often underutilized in many of the institutions. Mucosal nodules of variable sizes (2 to 6 mm) were found in 2 of the 8 patients in which the colonoscopy was performed. Colonoscopy has its limitations in its indications and observer variant.

Immunological tests

Chawla *et al* reported a high sensitivity of ELISA for soluble antigen and Bhargava *et al* used competitive ELISA with monoclonal antibody against 38 Kd protein and found a sensitivity of 81 per cent, specificity of 88 per cent and diagnostic accuracy of 84 per cent. These above tests were not evaluated in our study. The values of

immunological tests are not clearly understood in our clinical evaluation.

Ascitic fluid examination:

The ascitic fluid in tuberculosis is straw coloured with protein >3g/dl, and total cell count of 150-4000/ μ l, consisting predominantly of lymphocytes (>70%). The ascites to blood glucose ratio is less than 0.9650 and serum ascites albumin gradient is less than 1.1 g/dl. In our study group laboratory and microbiological evidence done in 22 patients revealed features of tuberculosis in 10 patients. Ascitic fluid microbiological evidence needs 6-8 weeks of period for organism growth and it has its diagnostic limitation.

ADA levels were determined in the ascitic fluid of 49 patients by Dwivedi *et al.* The levels in tuberculous ascites were significantly higher than those in cirrhotic or malignant ascitis. In co infection with HIV the ADA values can be normal or low. Falsely high values can occur in malignant ascitis. High interferon- γ level in tubercular ascites have been reported to be useful diagnostically. Combining both ADA and interferon estimations may further increase sensitivity and specificity.

Laparoscopic findings:

Bhargava *et al* studied 87 patients with high protein ascites, of which 38 were diagnosed as having tuberculosis. They found visual appearances to be more helpful (95% accurate) than histology, culture or guinea pig inoculation (82% and 37.5% sensitivity respectively). Caseating granulomas may be found in 85-90 per cent of the biopsies.

In our study group where 24 patients underwent diagnostic laparoscopy, 21 patients had features diagnosis of tuberculosis based on visual appearances. Tissue diagnosis taken for confirmation showed caseating granulomas in 18 patients.

Laparoscopy plays a very important role in the diagnosis of abdominal tuberculosis and it needs to be evaluated as initial diagnostic tool. Compared to other diagnostic modalities it has more sensitivity

Management:

On discussing about the surgical management of abdominal tuberculosis, Surgery is reserved for mechanical complications of tuberculosis or when medical therapy fails. Emergency surgery is indicated in the presence of acute complications such as free perforation of the bowel and severe intestinal haemorrhage. The most common indication for surgery is intestinal obstruction secondary to stricture formation. Predictors for surgical intervention are long

strictures (12 cm or more in length) and multiple areas of involvement. Other indications for surgery are bowel adhesions, intra-abdominal abscess due to a confined perforation, mesenteric abscess, and internal or external fistulas. Surgery is also appropriate if the diagnosis is in doubt and when malignancy cannot be ruled out with reasonable accuracy.^{15, 29,30}

Surgical procedures:

In our study group, only 23 patients needed surgical interventions whereas 27 patients were managed with chemotherapeutic drugs. Most frequent interventions were Right Hemicolectomy (6 patients), Strictureplasty (5 patients), Laparotomy and adhesiolysis (4 patients), Laparoscopy intervention (3 patients) and Laparotomy and Diversion Ileostomy (3 patients). Similar observations were made by Balasubramaniam *et al.*

Surgical techniques for patients with bowel obstruction have evolved over time. At one time, bypass procedures such as ileotransverse colostomy and enteroenterostomy were practiced commonly, but have now been abandoned because of complications such as blind-loop syndrome, malabsorption, and perforation. Similarly,

radical bowel resection and extensive dissection of mesenteric lymph nodes have fallen out of favour because of complications, including the development of short-bowel syndrome. The procedure preferred for an ileo-caecal mass is limited resection in place of the standard right hemicolectomy. Other procedures advocated are ileocaecoplasty and coloplasty for ileocaecal and colonic strictures. respectively.³¹⁻³³ Drainage tubes are not recommended and, if used, should be removed early because of increased risk of abdomino cutaneous fistulas if they are left in place for more than 7 days. All patients who underwent surgery were given anti tuberculous drugs post operatively.

CONCLUSION

Abdominal Tuberculosis is a diagnostic enigma due to its vague clinical manifestations. The diagnosis can remain inconclusive even after laboratory and radiological evaluation. To diagnose tuberculosis a high degree of clinical suspicion is the prime need.

Laparoscopy has a very significant role in establishing diagnosis in doubtful cases. Most patients with peritoneal tuberculosis can be diagnosed by laparoscopy, since the peritoneal appearances are fairly distinctive for tuberculosis.

The recommended surgical procedures today are conservative. Post operative conventional anti tubercular therapy for at least 6 months is recommended, but preoperative anti tubercular therapy is still controversial.

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PROFORMA

Name:

Age

Sex: IP

No:

Presenting Complaints

- Abdominal pain
- Vomiting
- Abdominal distension
- Loss of appetite
- Loss of weight
- Low grade fever, night sweats
- Diarrhea /Constipation
- Bleeding per rectum

Past History: DM / HTN / TB/ Epileptic /Surgery

Personal History: Diet/ Alcoholic /smoker

General Physical examination

- Consciousness & Orientation
- Built &Nourishment
- Pallor & Icterus

- Hydration
- Generalised edema & lymphadenopathy
- Vitals

Abdominal Examination:

- Distension
- Soft /rigidity
- Visible intestinal peristalsis
- Tenderness –yes/no
- Abdominal mass - size/ site/ consistency/mobility.
- Presence of free fluid
- Presence of gurgling
- Bowel sounds –present / absent.

Respiratory System :

- Chest –for any abnormality
 - Breath sounds –normal /abnormal **Cardiovascular system**

for heart status Central Nervous System:

- Higher mental functions
- Spinal tenderness.

- Any focal neurological deficit

Investigations.

- Haemoglobin
- total and differential white cell count
- ESR
- Blood sugar, Urea, Creatinine.
- Serum Electrolytes.
- Blood Grouping
- X-ray Chest & Abdomen
- Mantoux test.
- Ultrasound abdomen
- CT Scan Abdomen
- Barium studies
- Colonoscopy

Provisional diagnosis:

Final diagnosis:

Management:

Medical/Surgical

Surgical Post operative events:

INFORMATION SHEET

We are conducting a study on “ANALYSIS OF ABDOMINAL TUBERCULOSIS AND INCIDENCE OF TUBERCULOSIS IN PERIANAL FISTULA” among patients attending Rajiv Gandhi Government General Hospital, Chennai

The purpose of this study is to assess “ANALYSIS OF ABDOMINAL TUBERCULOSIS AND INCIDENCE OF TUBERCULOSIS IN PERIANAL FISTULA”

We are selecting certain cases and if you are found eligible, we may be using clinical profile, lab test reports and radiological reports for study purposes which does not affect your final report or management.

The privacy of the patients in the research will be maintained throughout the study. In the event of any publication or presentation resulting from the research, no personally identifiable information will be shared.

Taking part in this study is voluntary. You are free to decide whether to participate in this study or to withdraw at any time; your decision will not result in any loss of benefits to which you are otherwise entitled.

The results of the special study may be intimated to you at the end of the study period or during the study if anything is found abnormal which may aid in the management or treatment.

Signature of Investigator

Signature of Participant / Guardian

Place:

Date :

ANNEXURE 4 -ஆ

ஆராய்ச்சிதகவல் தாள்

ஆராய்ச்சிதலைப்பு: வயிற்று பகுதியில் காசநோயினால் ஏற்படும் பாதிப்பு, கண்டறிதல், வெவ்வேறு விதமான தீர்வுகள் சம்பந்தமான ஓர் ஆய்வு.

ஆய்வாளர் : பா.பாலாஜி திருவேங்கடம்
பொது அறுவை சிகிச்சைதுறை
சென்னைமருத்துவக்கல்லூரி,
சென்னை-600 003.

இராஜிவ்காந்தி அரசு பொதுமருத்துவமனைக்கு வரும் நோயாளிகளில் வயிற்று பகுதியில் காசநோய் அறிகுறிகள் உள்ளோர்களை தேர்ந்தெடுத்து அவர்களின் வெவ்வேறு அறிகுறிகள், கண்டறியும் முறைகள் அதற்கான தீர்வு முறைகள் பற்றிய ஆய்வு இங்கு நடைபெற்று வருகிறது.

நீங்களும் இந்த ஆராய்ச்சியில் பங்கேற்க நாங்கள் விரும்புகிறோம். இந்த ஆராய்ச்சியில் உங்களுடைய வயிற்று பகுதி காசநோய் சம்பந்தமாக சில சிறப்புப் பரிசோதனைக்கு உட்படுத்தி தகவல்களை ஆராய்வோம். இதனால் தங்களது நோய்யின் ஆய்வரிக்கையும் அல்லது சிகிச்சையோ பாதிப்புக்குள்ளாகாது என்பதையும் தெரிவித்துக் கொள்கிறேன்.

முடிவுகளை அல்லது கருத்துக்களை வெளியிடும் போதும் அல்லது ஆராய்ச்சியின் போதும் தங்களது பெயரோ அல்லது அடையாளங்களையோ வெளியிடமாட்டோம் என்பதையும் தெரிவித்துக் கொள்கிறோம்.

இந்த ஆராய்ச்சியில் பங்கேற்பது தங்களுடைய விருப்பத்தில் பேரில் தான் இருக்கிறது. மேலும் நீங்கள் எந்நேரமும் இந்த ஆராய்ச்சியில் இருந்து பின்வாங்கலாம் என்பதையும் தெரிவித்துக் கொள்கிறோம்.

இந்த சிறப்புப் பரிசோதனையின் முடிவுகளை ஆராய்ச்சியின் போது அல்லது ஆராய்ச்சியின் முடிவின் போதும் தங்களுக்கு அறிவிப்போம் என்பதையும் தெரிவித்துக் கொள்கிறோம்.

இந்த ஆய்வைப்பற்றிய சந்தேகங்களுக்கு தொடர்புகொள்ள வேண்டியவர்:
நா.பாலமுருகன்

பங்கேற்பாளர் கையொப்பம்..... இடம் : தேதி:

பங்கேற்பாளர் பெயர் மற்றும் விலாசம் :

ஆராய்ச்சியாளர் கையொப்பம் இடம் : தேதி:.....

PATIENT CONSENT FORM

Study Detail : ANALYSIS OF ABDOMINAL TUBERCULOSIS AND
INCIDENCE OF TUBERCULOSIS IN PERIANAL FISTULA

Study Centre : Rajiv Gandhi Government General Hospital, Chennai.

Patient's Name :

Patient's Age :

Identification Number :

Patient may check (☑) these boxes

The details of the study have been provided to me in writing and explained to me in my own language

I understand that my participation in the study is voluntary and that I am free to withdraw at any time without giving reason, without my legal rights being affected.

I understand that sponsor of the clinical study, others working on the sponsor's behalf, the ethical committee and the regulatory authorities will not need my permission to look at my health records, both in respect of current study and any further research that may be conducted in relation to it, even if I withdraw from the study I agree to this access. However, I understand that my identity will not be revealed in any information released to third parties or published, unless as required under the law. I agree not to restrict the use of any data or results that arise from this study.

I agree to take part in the above study and to comply with the instructions given during the study and faithfully cooperate with the study team and to immediately inform the study staff if I suffer from any deterioration in my health or well being or any unexpected or unusual symptoms.

I hereby consent to participate in this study.

I hereby give permission to undergo complete clinical examination , biochemical and radiological tests

Signature of Investigator

Signature/thumb impression

Study Investigator's Name:

Patient's Name and Address:

DR.BALAJI TIROUVINGADAME

ANNEXURE 6

ஆராய்ச்சிஒப்புதல் கடிதம்

ஆராய்ச்சிதலைப்பு: வயிற்று பகுதியில் காசநோயினால் ஏற்படும் பாதிப்பு, கண்டறிதல், வெவ்வேறு விதமான தீர்வுகள் பற்றிய ஓர் ஆய்வு.

சென்னைமருத்துவக் கல்லூரி நோய் குறியியல் துறையில் பயிலும் முதுகலை மருத்துவர் பா.பாலாஜி திருவேங்கடம் அவர்கள் மேற்கொள்ளும் இந்த ஆய்வில் பங்குகொள்ள ஆகியநான் முழுமனதுடன் சம்மதிக்கிறேன்.

எனக்கு விளக்கப்பட்ட விஷயங்களை நான் புரிந்துக் கொண்டு எனது சம்மதத்தை தெரிவிக்கின்றேன்.

இந்த ஆராய்ச்சியில் பிறரின் நிர்பந்தமின்றி எனது சொந்த விருப்பத்தின் பேரில் தான் பங்கு பெறுகிறேன் மற்றும் நான் இந்த ஆராய்ச்சியில் இருந்து எந்நேரமும் பின் வாங்கலாம் என்பதையும் இதனால் எந்த பாதிப்பும் ஏற்படாது என்பதையும் நான் புரிந்துக் கொண்டேன்.

நான் கல்லீரல்சீழ் தொற்று நோய் குறித்த இந்த ஆராய்ச்சி விவரங்களை கொண்ட தகவல் தாளை பெற்றுக் கொண்டேன்.

நான் என்னுடைய சுயநினைவுடன் மற்றும் முழு சுதந்திரத்துடன் இந்த மருத்துவ ஆராய்ச்சியில் என்னை சேர்த்துக் கொள்ள சம்மதிக்கிறேன்.

பங்கேற்பாளர் கையொப்பம்..... இடம் : தேதி:

பங்கேற்பாளர் பெயர் மற்றும் விலாசம் :

ஆராய்ச்சியாளர் கையொப்பம் இடம் : தேதி:.....

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CERTIFICATE – II

This is to certify that this dissertation work titled “**DESCRIPTIVE STUDY ON ANALYSIS OF ABDOMINAL TUBERCULOSIS AND INCIDENCE OF TUBERCULOSIS IN PERIANAL FISTULAE**” of the candidate **Dr.B.BALAJI TIROUVINGADAME** with Registration Number 221511015 for the award of M.S. in the branch of GENERAL SURGERY. I personally verified the urkund.com website for plagiarism Check. I found that the uploaded thesis file contains from introduction to conclusion pages and result shows 10 percentage of plagiarism in the dissertation.

Guide & Supervisor sign with Seal.

**INSTITUTIONAL ETHICS COMMITTEE
MADRAS MEDICAL COLLEGE, CHENNAI 600 003**

EC Reg.No.ECR/270/Inst./TN/2013
Telephone No.044 25305301
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CERTIFICATE OF APPROVAL

To
Dr.B.Balaji Tirouvingadame
Post Graduate in M.S.(General Surgery)
Institute of General Surgery
Madras Medical College/RGGGH
Chennai 600 003

Dear Dr.B.Balaji Tirouvingadame,

The Institutional Ethics Committee has considered your request and approved your study titled **"ANALYSIS OF ABDOMINAL TUBERCULOSIS AND INCIDENCE OF TUBERCULOSIS IN PERIANAL FISTULA "** - NO.20022017

The following members of Ethics Committee were present in the meeting hold on **07.02.2017** conducted at Madras Medical College, Chennai 3

- | | |
|---|---------------------|
| 1.Dr.C.Rajendran, MD., | :Chairperson |
| 2.Dr.M.K.Muralidharan,MS.,M.Ch.,Dean, MMC,Ch-3 | :Deputy Chairperson |
| 3.Prof.Sudha Seshayyan,MD., Vice Principal,MMC,Ch-3 | : Member Secretary |
| 4.Prof.S.Suresh, MS., Prof.of Surgery, MMC, Ch-3 | : Member |
| 5.Prof.Baby Vasumathi,MD.,Director, Inst. of O & G | : Member |
| 6.Prof.K.Ramadevi,MD.,Director,Inst.of Bio-Che,MMC,Ch-3 | : Member |
| 7.Prof.R.Padmavathy, MD, Director,Inst.of Pathology,MMC,Ch-3 | : Member |
| 8.Prof.S.Mayilvahanan,MD,Director, Inst. of Int.Med,MMC, Ch-3 | : Member |
| 9.Tmt.J.Rajalakshmi, JAO,MMC, Ch-3 | : Lay Person |
| 10.Thiru S.Govindasamy, BA.,BL,High Court,Chennai | : Lawyer |
| 11.Tmt.Arnold Saulina, MA.,MSW., | :Social Scientist |

We approve the proposal to be conducted in its presented form.

The Institutional Ethics Committee expects to be informed about the progress of the study and SAE occurring in the course of the study, any changes in the protocol and patients information/informed consent and asks to be provided a copy of the final report.

Member Secretary – Ethics Committee

**MEMBER SECRETARY
INSTITUTIONAL ETHICS COMMITTEE
MADRAS MEDICAL COLLEGE
CHENNAI-600 003**

KEY TO MASTER CHART

ESR	-	Erythrocyte sedimentation rate
BM	-	Barium Meal
BE	-	Barium Enema
CT ABD	-	CT Abdomen
D/L	-	Diagnostic Laparoscopy

MASTER CHART															
Name	Age	Sex	Clinical Presentation	Lymphocytes	ESR	Ascitic fluid	Chest x-ray	BM	BE	CT ABD	D/L	Diagnosis	TREATMENT	TIMING	PERIANAL FISTULA
Rajesh	17	M	Non specific symptoms	Present	Raised		Normal			d(+)	d(+)	MESENTRIC LYMPADENITIS	MEDICAL MANAGEMENT		NO
Panneerselvam	32	M	ascites	Present	Raised	Positive	Normal					TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Malliga	31	F	Abdominal mass	Present	Raised		Normal		(+)	d(+)	d(+)	ILEOCAECAECAL T.B	RIGHT HEMICOLECTOMY	Elective	NO
Anbarasan	25	M	intestinal obstruction	Absent	Raised		Normal			d(+)	d(+)	INTERLOOP ADHESIONS	LAPAROSCOPY-ADHESIOLYSIS	Elective	NO
Priya	21	F	Abdominal pain	Absent	Normal		Positive			d(+)	d(+)	INTERLOOP ADHESIONS	LAPAROTOMY- ADHESIOLYSIS	Elective	NO
Kumar	19	M	Non specific symptoms	present	Raised		Normal			d(-)	d(+)	MESENTRIC LYMPADENITIS	MEDICAL MANAGEMENT		NO
Amina bee	33	F	intestinal obstruction	Absent	Raised		Normal	(+)		d(+)	d(+)	INTERLOOP ADHESIONS	LAPAROSCOPY-ADHESIOLYSIS	Elective	NO
Manivasagam	35	M	ascites	Absent	Raised	Positive	Normal					TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Raj	31	M	Abdominal mass	present	Raised		Normal		(+)	d(+)	d(+)	ILEOCAECAECAL T.B	RIGHT HEMICOLECTOMY	Elective	NO
Punniyakodi	41	M	intestinal obstruction	present	Raised		Normal			d(+)		ILEAL PERFORATION	LAPAROTOMY- LIMITED RESECTION	Emergency	NO
Natarajan	37	M	intestinal obstruction	Absent	Raised		Normal			d(+)		ILEAL STRICTURE	LAPAROTOMY-STRICTUROPLASTY	Emergency	NO
Meena	19	F	Abdominal pain	present	Raised		Positive			d(+)		INTERLOOP ADHESIONS	LAPAROSCOPY-ADHESIOLYSIS	Elective	NO
Murugan	26	M	ascites	present	Raised	Negative	Normal	(+)				T.B ABDOMEN	MEDICAL MANAGEMENT		NO
Renuka	31	F	Abdominal mass	present	Raised		Normal			d(+)	d(+)	ILEOCAECAECAL T.B	RIGHT HEMICOLECTOMY	Elective	NO
Yusuf	20	M	Non specific symptoms	Absent	Raised		Normal			d(+)		MESENTRIC LYMPADENITIS	MEDICAL MANAGEMENT		NO
Anitha	24	F	ascites	Absent	Raised	Positive	Normal					TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Suresh	39	M	Abdominal mass	present	Raised		Normal	(+)		d(+)	d(+)	ILEAL STRICTURE	LAPAROTOMY- STRICTUROPLASTY	Elective	NO
Rajan	44	M	ascites	Absent	Raised	Positive	Normal					TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Sivaraja	34	M	Abdominal mass	present	Raised		Positive			d(+)	d(-)	ILEOCAECAECAL T.B	RIGHT HEMICOLECTOMY	Elective	NO
Muthuraj	40	M	intestinal obstruction	Absent	Normal		Normal		(-)	d(+)	d(+)	INTERLOOP ADHESIONS	LAPAROSCOPY-ADHESIOLYSIS	Elective	NO
Mumtaj	36	F	Abdominal mass	present	Raised		Normal			d(-)	d(+)	T.B ABDOMEN	MEDICAL MANAGEMENT		NO
Durai	29	M	intestinal obstruction	Absent	Normal		Positive			d(+)		ILEAL PERFORATION	LAPAROTOMY-LIMITED RESECTION	Emergency	NO
Amala	19	F	Non specific symptoms	present	Raised		Normal	(+)		d(-)		T.B.ABDOMEN	MEDICAL MANAGEMENT		NO
Palpandi	30	M	ascites	Absent	Raised	Negative	Normal	(+)				T.B.ABDOMEN	MEDICAL MANAGEMENT		NO
Mynaa	25	F	ascites	present	Raised	Negative	Normal			d(+)		TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Masilamani	40	M	Abdominal mass	present	Raised		Positive	(+)	d(-)	d(+)		T.B.ABDOMEN	MEDICAL MANAGEMENT		NO
Akbar	23	M	ascites	present	Raised	Positive	Normal					TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO

Palani	35	M	intestinal obstruction	Absent	Raised		Normal		d(+)		ILEAL STRICTURE	LAPAROTOMY-STRICTUROPLASTY	Emergency	NO
Balakumar	18	M	Abdominal pain	present	Raised		Normal	(-)	d(-)	d(+)	T.B.ABDOMEN	MEDICAL MANAGEMENT		NO
Revathi	36	F	intestinal obstruction	Absent	Raised		Normal		d(+)		INTERLOOP ADHESIONS	LAPAROTOMY-ADHESIOLYSIS	Emergency	NO
Munirathinam	50	M	Abdominal mass	present	Raised		Normal	(+)	d(+)	d(-)	T.B.ABDOMEN	MEDICAL MANAGEMENT		YES
Alamelu	31	F	ascites	present	Raised	Positive	Normal				TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Boominathan	32	M	ascites	present	Raised	Negative	Normal		d(+)	d(+)	T.B.ABDOMEN	MEDICAL MANAGEMENT		NO
Vedammal	41	F	intestinal obstruction	Absent	Normal		Normal		d(+)		INTERLOOP ADHESIONS	LAPAROTOMY-ADHESIOLYSIS	Emergency	NO
Alagar	42	M	Abdominal mass	present	Raised		Positive		d(+)	d(+)	ILEOCAECAL T.B	RIGHT HEMICOLECTOMY	Elective	NO
Arul murugan	21	M	ascites	present	Raised	Positive	Normal				TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Sivanesan	30	M	Abdominal mass	present	Raised		Normal		d(-)	d(+)	ILEAL STRICTURE	LAPAROTOMY-STRICTUROPLASTY	Elective	NO
Thangamani	25	M	Abdominal pain	present	Raised		Positive	(-)	d(+)	d(-)	MESENTRIC LYMPADENITIS	MEDICAL MANAGEMENT		NO
Kamala	33	F	intestinal obstruction	Absent	Raised		Normal		d(+)		ILEAL STRICTURE	LAPAROTOMY-DIVERSION ILEOSTOMY	Emergency	NO
Ayyanar	36	M	ascites	present	Raised	Positive	Normal				TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Devi	18	F	Abdominal pain	Absent	Raised		Normal		d(-)	d(+)	INTERLOOP ADHESIONS	LAPAROTOMY -ADHESIOLYSIS	Elective	NO
Chinnathambi	35	M	intestinal obstruction	present	Normal		Normal		d(+)		ILEAL STRICTURE	LAPAROTOMY-DIVERSION ILEOSTOMY	Emergency	NO
Pushpa	40	F	Abdominal mass	Absent	Raised		Positive	(-)	d(-)	d(+)	T.B.ABDOMEN	MEDICAL MANAGEMENT		NO
Mohammed	40	M	Abdominal pain	Absent	Raised	Negative	Normal		d(+)		MESENTRIC LYMPADENITIS	MEDICAL MANAGEMENT		NO
Mannar	32	M	intestinal obstruction	present	Raised		Normal		d(+)	d(+)	ILEAL STRICTURE	LAPAROTOMY-DIVERSION ILEOSTOMY	Elective	NO
Samy	19	M	Abdominal pain	Absent	Raised		Normal	(-)	d(-)	d(+)	MESENTRIC LYMPADENITIS	MEDICAL MANAGEMENT		NO
Ayyavu	26	M	ascites	present	Raised	Positive	Normal				TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO
Mangai	50	F	Abdominal mass	Absent	Raised		Normal		d(+)	d(+)	ILEOCAECAL T.B	RIGHT HEMICOLECTOMY	Elective	NO
Ayyappan	22	M	Non specific symptoms	present	Raised		Positive	(+)	d(+)		T.B.ABDOMEN	MEDICAL MANAGEMENT		NO
Poongavanam	29	F	ascites	present	Raised	Positive	Normal				TUBERCULOUS ASCITES	MEDICAL MANAGEMENT		NO