

Clinical, Microbiological and radiological study of community acquired Pneumonia

Dr.Kaliparambil Sugathan Roshni¹; Dr.P.C.Mishra²; Dr.S.C.Mohapatra³;
Dr.A.Swetha⁴; Dr.S.Prasanna Kumar⁵, Dr.Sushanth Vemuganti⁶.

¹Dr. Kaliparambil Sugathan Roshni, Junior Resident, General Medicine, Maharaja's Institute of Medical Sciences, Nellimarla, Vizianagaram Dist, AP, India.

²Dr. P.C.Mishra, M.D., General Medicine, Professor, Maharaja's Institute of Medical Sciences, Nellimarla, Vizianagaram Dist, AP, India.

³Dr. S.C.Mohapatra, M.D., General Medicine, Professor & HOD, Maharaja's Institute of Medical Sciences, Nellimarla, Vizianagaram Dist, AP, India.

⁴Dr. A.Swetha, General Medicine, Maharaja's Institute of Medical Sciences, Nellimarla, Vizianagaram Dist, AP, India.

⁵Dr. S. Prasanna Kumar, Junior Resident, General Medicine, Maharaja's Institute of Medical Sciences, Nellimarla, Vizianagaram Dist, AP, India.

⁶Dr. Sushanth Vemuganti, Junior Resident, General Medicine, Maharaja's Institute of Medical Sciences, Nellimarla, Vizianagaram Dist, AP, India.

Abstract:

Background: The true incidence of pneumonia acquired in the community is unknown, but this is a common clinical problem worldwide especially in developing countries and remains a leading cause of death in India.

Aims and Objectives: The aim and objective of the study were to check clinical, radiological, and bacteriological profile of patients of community-acquired pneumonia (CAP).

Materials and Methods: The present study was undertaken in Department of Medicine General Medicine Department, Maharaja's Institute of Medical Sciences, Nellimarla, Vizianagaram, Vizianagaram Dist, India from October 2014 to September 2016.. For the study, 60 indoor patients of >15 years of age group were selected from Medicine Department, diagnosed as CAP. In all studied patient's chest-X-rays, routine laboratory test, sputum, and blood culture were done.

Results: Despite the use of standard protocols, microbiological diagnosis of CAP was confirmed only in 28 (45.5%) of patients by sputum and blood culture. Sputum was the most common etiological source of organism isolation (22) followed by blood (6), *Streptococcus pneumoniae* was the commonest pathogen 10 (36.4%). Followed by *Klebsiella pneumoniae* 8 (29.%), *Staphylococcus aureus* 6 (20%) and other Gram-negatives bacilli* 8 (14.5%.) *(*Haemophilus influenzae* 5.5%, *Pseudomonas* 1.8%, *Acinetobacter* 1.8%, *Enterobacter* 1.8%, *Escherichia coli* 1.8%, *Citrobacter* 1.8%). CAP was found predominantly in males (67.5%) and elderly age group (68.3%). Maximum number of patients presented with cough (92.5%), fever (90%), dyspnea (59.2%), expectoration (55%), pleuritic chest pain (14.2%), most common predisposing factors associated with CAP in the following chronological order-smoking (40.8%) > chronic obstructive airway disease (35.8%) > cardiovascular disease (16.7%) > alcoholism (12.5%) > diabetes mellitus (6.7%) > neurological disorders (2.5%). Lobar pneumonia especially right lower lobe consolidation was the most common radiological finding observed in 48.3% patients, followed by left lower lobe infiltration ($P < 0.0001$).

Conclusion: *S. pneumoniae* was the most common pathogen, but the emergence of the higher incidence of Gram-negative organism specially *K. pneumoniae* has occurred in our geographical area (India). Age, smoking, and underlying co-morbid conditions especially chronic obstructive pulmonary disease were significantly associated with the development of CAP ($P < 0.01$). Radiographic changes usually cannot be used to distinguish bacterial from the nonbacterial pneumonia.

Keywords: Bacteriology, Blood culture, Chronic obstructive pulmonary disease, Community-acquired pneumonia, Radiology smoking, Sputum culture

Date of Submission: 26-01-2018

Date of acceptance: 09-02-2018

I. Introduction

Pneumonia is a disease known to mankind from antiquity. Pneumonia “the captain of men of death,” “The friend of the aged, allowing them a merciful relief from those cold gradations of decay, that make the last state of all so distressing” as described by William Osler, is one of the most common infectious disease encountered in the clinical practice ¹.

Community acquired pneumonia (CAP) is an acute illness acquired in the community with symptoms suggestive of Lower Respiratory Tract Infection (LRTI) together with presence of a chest radiograph of intra pulmonary shadowing which is likely to be new and has no clear alternative cause.^{1,2}

Community acquired pneumonia is a common disorder with an incidence of about 20% to 30% in developing countries compared to an incidence of 3% to 4 % in developed countries.^{6,43,44} The aetiology of CAP remains uncertain in many patients. Even with the use of extensive laboratory testing and invasive procedures, etiological confirmation being achieved in no more than 45% to 70% of patients.^{51,52} Streptococcus pneumonia is the most commonly isolated pathogen responsible for 35% to 60% of cases.^{53,54} The bacteriological profile of community-acquired pneumonia is different in different countries and changing with time within the same country, probably due to frequent use of antibiotics, changes in environmental pollution, increased awareness of the disease and changes in life expectancy.

Prognostic scoring systems for CAP have been developed to address these issues. The two prominent tools for this purpose are the pneumonia severity index (PSI) and the BTS rule, which has recently been modified to the CURB-65 rule.^{22, 23}

Treatment of CAP continues to be a challenge even in 21st century. Recommendations for CAP therapy are different, depending on whether patients require hospitalization or are treated as outpatients²⁴. Other issues for treatment recommendations include the emergence of antibiotic resistance among Streptococcus pneumonia and mono versus combination antibiotic therapy.

II. Aims And Objectives

- To study the clinical presentation of Community Acquired Pneumonia (CAP) in Maharaja Institute of Medical Sciences; Vizianagaram; Andhra Pradesh.
- To study the risk factors associated with CAP.
- To assess the spectrum of organisms responsible for CAP.
- To study the complications.
- To study the response to treatment and duration of hospital stay.

III. Materials And Methods

Total of 50 patients of age group >15 years, diagnosed as Community Acquired Pneumonia (CAP) and admit in to ICU and Medical wards, Maharajah’s Institute of Medical Sciences, Nellimarla, Vizianagaram from October 2014 to September 2016.

3.1 Inclusion criteria

Patients with new or progressive pulmonary infiltrates on chest radiograph together with at least two of the following:

- Fever, Cough, Production of sputum or
- Leucocytosis $\geq 11,000/ \text{mm}^3$.

3.2 Exclusion criteria

- Patients with radiographic or laboratory evidence suggestive of tuberculosis, acquired immunodeficiency syndrome (AIDS), leukaemia and those with chest infiltrates due to other causes such as congestive heart failure, pulmonary infarction or obstructive pneumonia due to lung cancer, and patients receiving immunosuppressive treatment were excluded from the study.

III.3 Method of the study:

In all the patients, detail history was taken and clinical examination was done. All of them were subjected to undergo chest radiograph, complete haemogram, renal and liver function tests, fasting blood sugar and serum electrolytes estimation were done. All efforts were made to obtain sputum at the time of initial clinical evaluation or within 24 hours of admission. In patients who could not expectorate sputum spontaneously, sputum was induced by nebulisation with 3% hypertonic saline. Sputum originating from lower respiratory tract containing > 25 polymorphonuclear leucocytes and < 10 epithelial cells per low power field (total magnification x 100) was subjected to Gram’s staining.

Sputum was also subjected to bacterial culture on blood agar and MacConkeys agar media. Two blood culture samples were also obtained from each patient at the time of initial visit from different vene-puncture sites and were cultured on blood agar and MacConkey's agar media. All patients were hospitalized and one full course of antibiotics treatment according to sensitivity given.

III.4 Investigations used:

- Sputum - Direct smear by Gram and Ziehl - Neelsen stains, Culture and antimicrobial sensitivity testing.
- Blood culture - Frequently positive in pneumococcal pneumonia
- Chest X-Ray - P/A view

A high neutrophil leukocytosis favors a diagnosis of bacterial (particularly pneumococcal) pneumonia; patients with pneumonia caused by atypical agents tend to have a marginally raised or normal white cell count. A marked leucopenia indicates either a viral aetiology or overwhelming bacterial infections.

IV. Observation And Results

Total 50 patients of age group >15 years, diagnosed as Community Acquired Pneumonia (CAP) and admit in to ICU and Medical wards, Maharajah's Institute of Medical Sciences, Nellimarla, Vizianagaram

Table 1 Age Distribution of Patients

AGE (YEARS)	MALES		FEMALES		TOTAL	
	NO.	%	NO.	%	NO.	%
<20	1	1.66	-	-	1	1.66
21-30	5	8.33	3	2	8	13.33
31-40	6	10	2	3.33	8	13.33
41-50	8	13.33	5	8.33	13	21.66
51-60	20	33.3	2	3.33	22	36.66
61-70	5	8.33	1	1.66	6	10
>70	2	3.33	-	-	2	3.33
TOTAL	47	78.33	13	21.66	60	100

Table -2 : sex distribution

	Mean (yrs)	Standard deviation (yrs)	Range (yrs)
Male	49.77	13.66	19 – 80
Female	42.85	12.88	24 – 66
Total	48.27	13.69	19 – 80

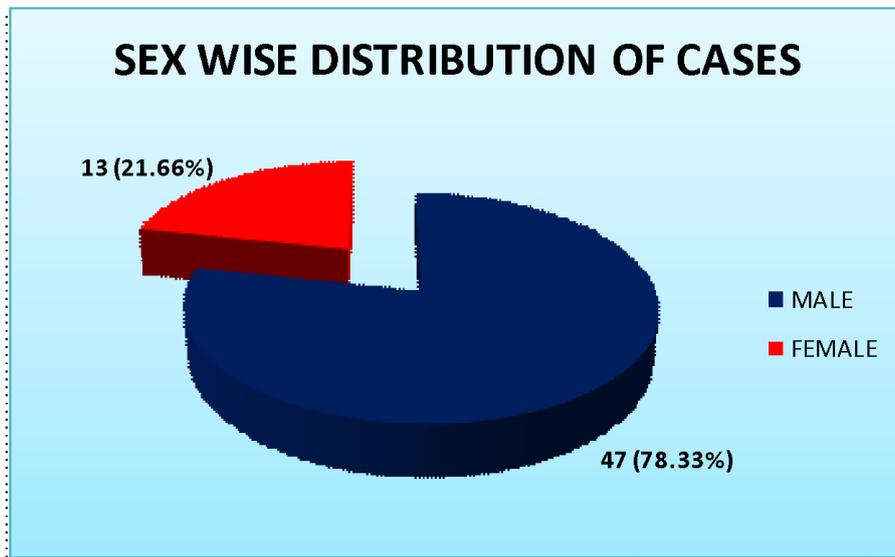
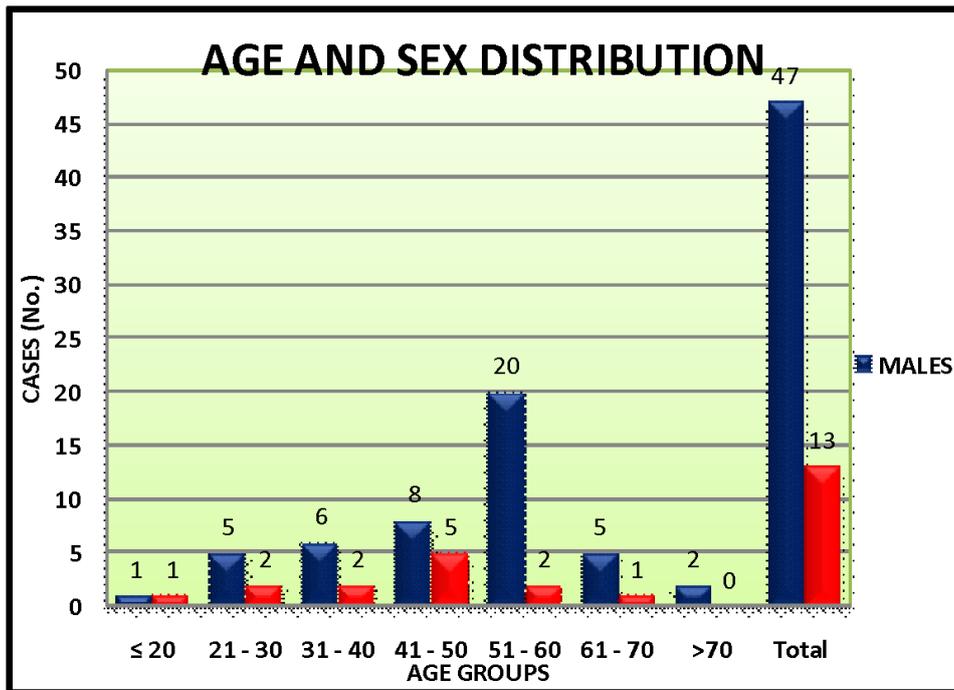


Table 3 : : Distribution in Rural and Urban Areas

SL. NO.	URBAN/RURAL	NO. OF CASES	PERCENTAGE
1	RURAL	41	68.33%
2	URBAN	19	31.66%
TOTAL		60	100%

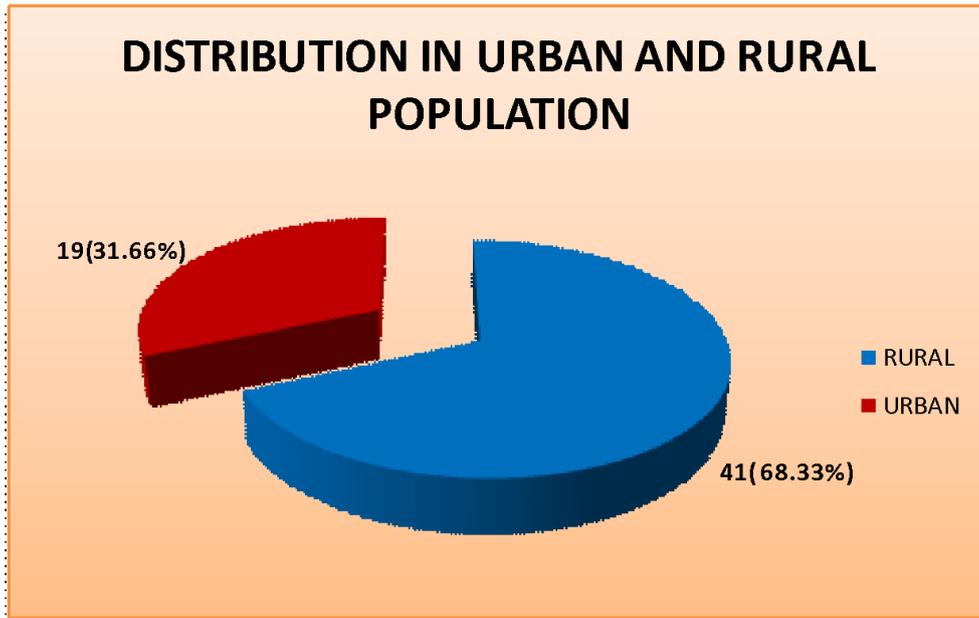


Table 4 Duration of Hospital Stay

Duration (Days)	Number	%
5-10	50	83.33%
11-15	7	11.66%
16-21	3	5%
Total	60	100%

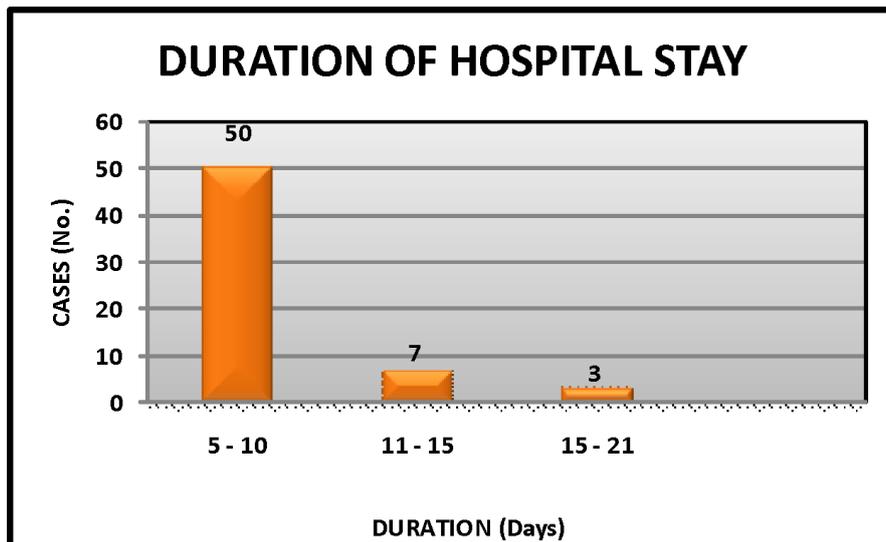


Table -5: Predisposing Factors

Signs Distribution of Patients

Signs	Number	%
HYPERTENSION	4	6.66%
DIABETES	6	10%
COPD	11	18.33%
SMOKING	22	36.66%
ALCOHOLISM	13	21.66%

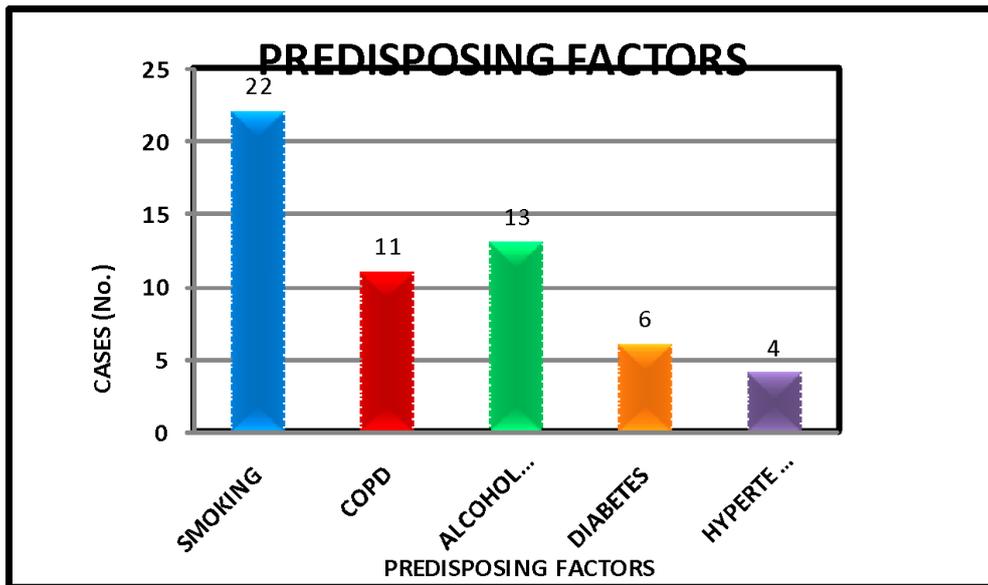


Table 6 : Presenting Symptoms

Signs	Number	%
FEVER	60	100%
COUGH	60	100%
EXPECTORATION	52	86.66%
DYSPNOEA	49	81.66%
CHEST PAIN	38	63.33%

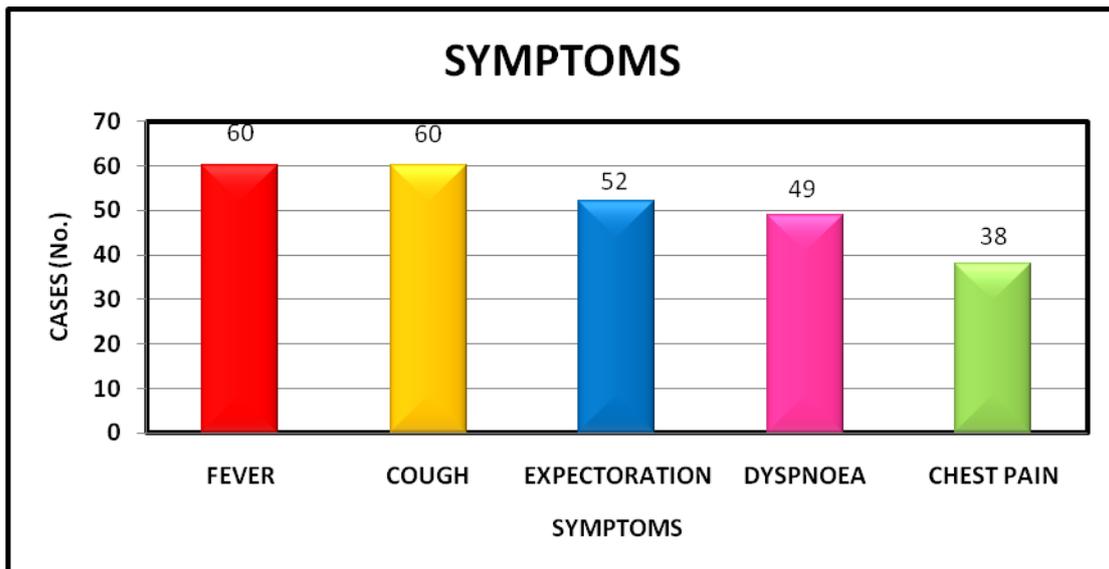


Table -7 : General Examination

Signs	Number	%
PALLOR	14	23.33%
ICTERUS	2	3.33%
CLUBBING	4	6.66%
CYANOSIS	5	8.33%
PEDAL EDEMA	1	1.66%

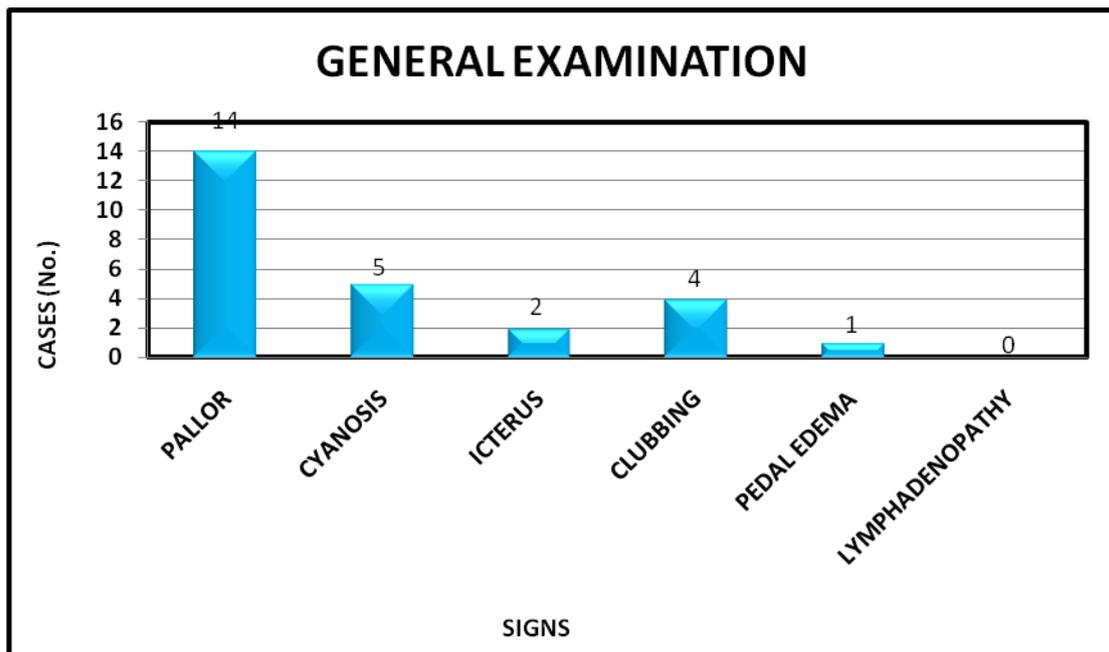


Table-8 : Systemic Examination

FEATURE	NO. OF CASES	PERCENTAGE
INCREASED VOCAL FREMITUS & VOCAL RESONANCE	50	83.33%
BRONCHIAL BREATHING	53	88.33%
WHISPERED PECTORILOQUY	34	56.66%
CREPITATIONS	41	68.33%

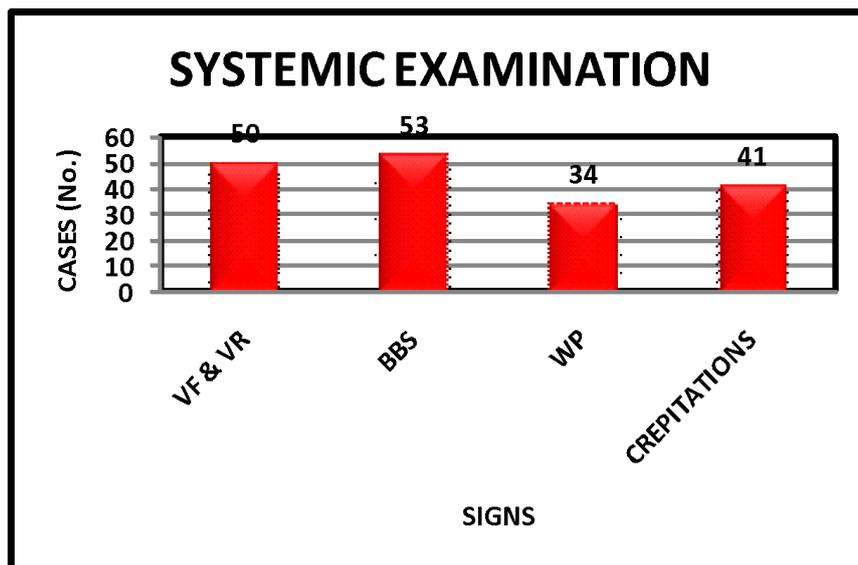


Table – 9 Sputum Staining

STAINING	NO. OF CASES	PERCENTAGE%
GRAM POSITIVE	30	50
GRAM NEGATIVE	14	23.33
MIXED	1	1.66
NO ORGANISM	15	25

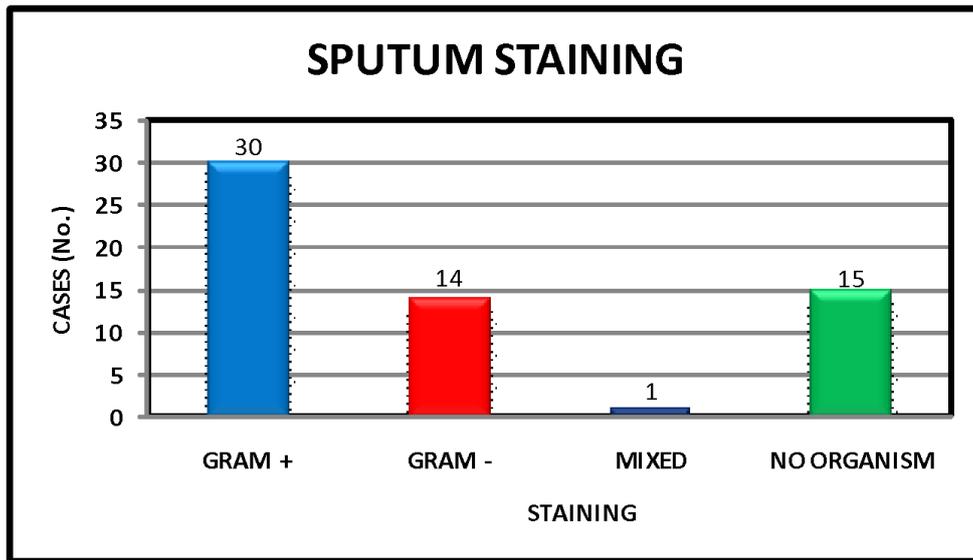


Table 10 :Sputum Culture

SL. NO.	ORGANISM	NO. OF CASES	PERCENTAGE
1	Streptococcus pneumoniae	21	35%
2	Staphylococcus aureus	9	15%
3	Klebsiella pneumoniae	7	11.66%
4	Pseudomonas aeruginosa	5	8.33%
5	E. Coli	2	3.33%
6	Mixed	1	1.66%
7	No organism	15	25%
TOTAL		60	100

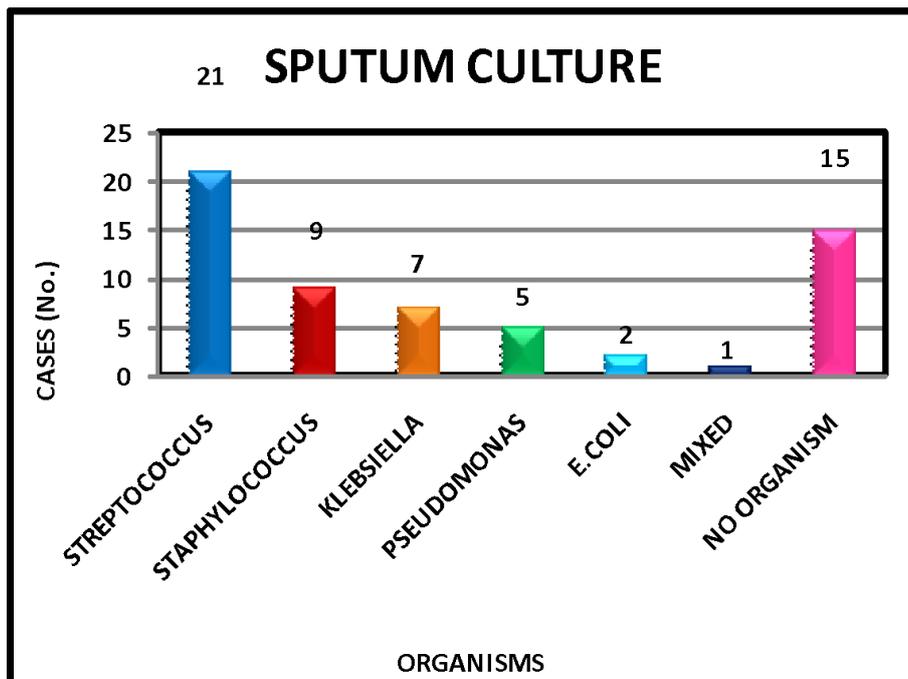


Table 11: Anatomical Localisation

S.No.	MODE OF LOCALIZATION	No. OF CASES	PERCENTAGE
1.	Unilateral involvement • Right lung • Left lung	55 38 17	91.66% 63.33% 28.33%
2.	Bilateral involvement	5	8.33%
3.	Right upper lobe	2	3.33%
4.	Right middle lobe	11	18.33%
5.	Right lower lobe	22	36.66%
6.	Left upper lobe	3	5%
7.	Left lower lobe	14	23.33%
8.	More than one lobe	8	13.33%

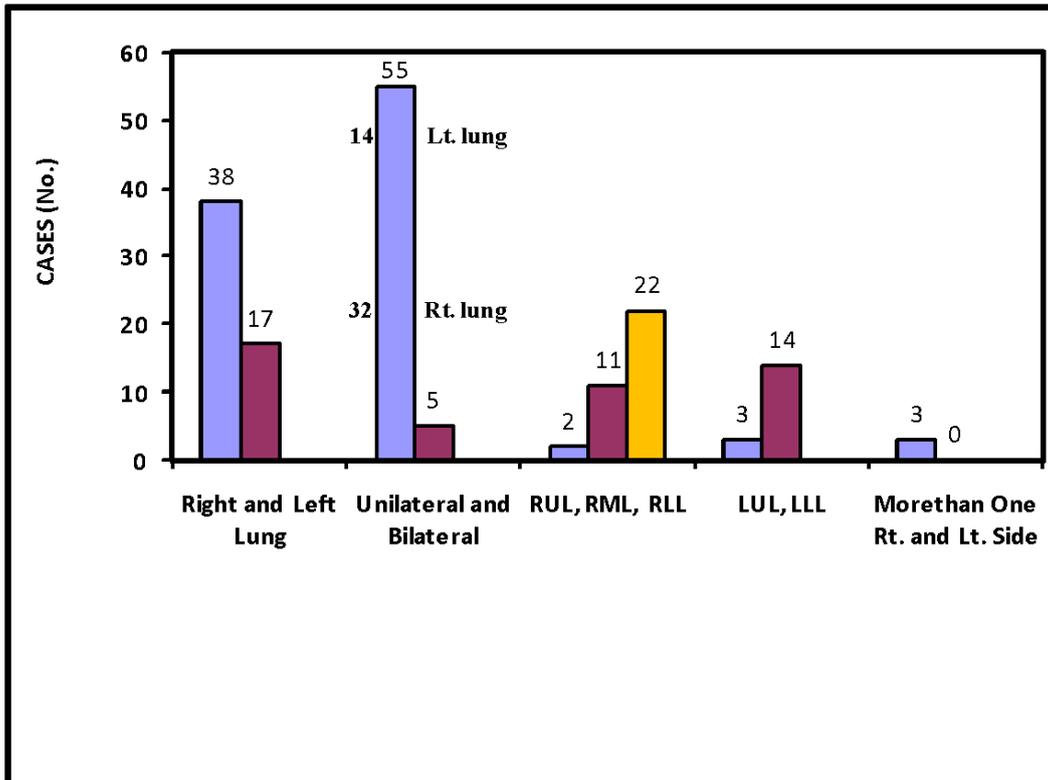


Table 12: Complications

SL. NO.	COMPLICATION	NO. OF CASES	PERCENTAGE
1	Pleural Effusion	2	3.33%
2	Empyema	-	-
3	Lung abscess	2	3.33%
4	Circulatory failure	6	10%
5	Jaundice	2	3.33%
6	Acute renal failure	2	3.33%

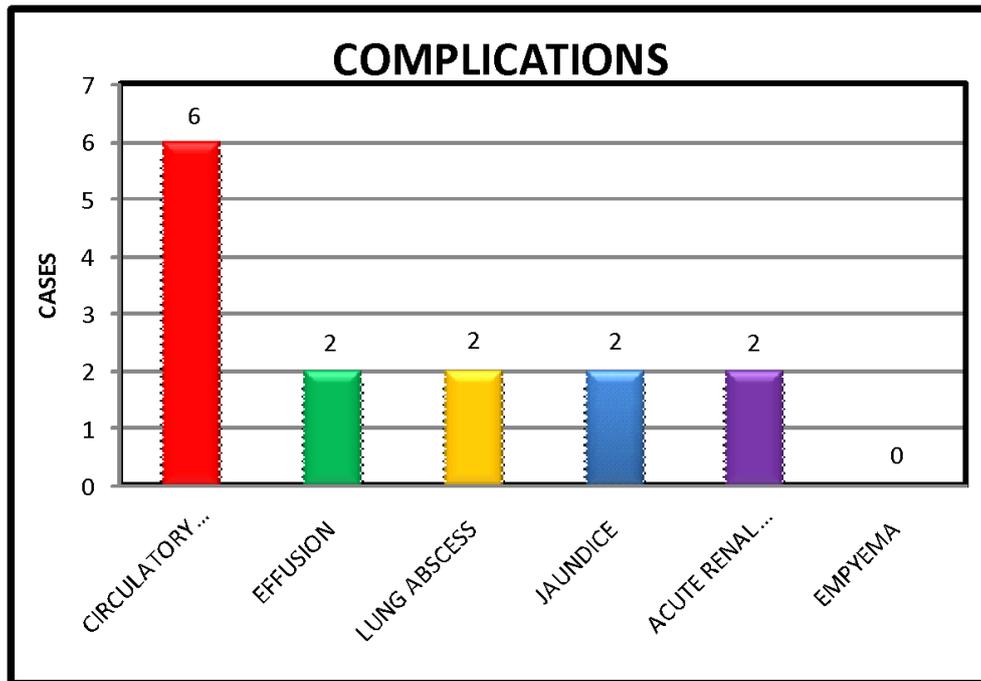
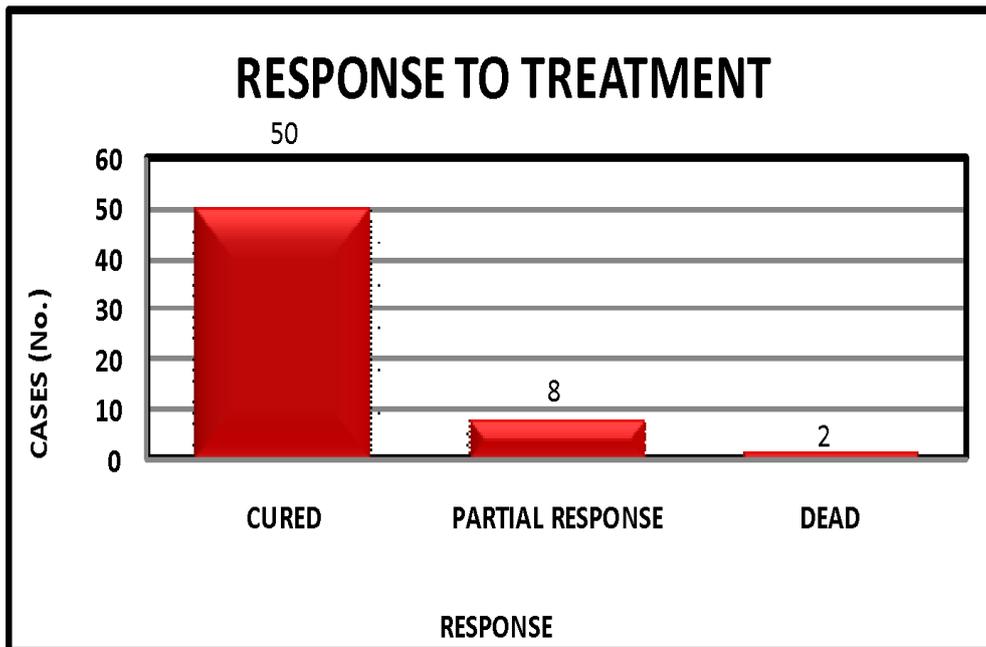
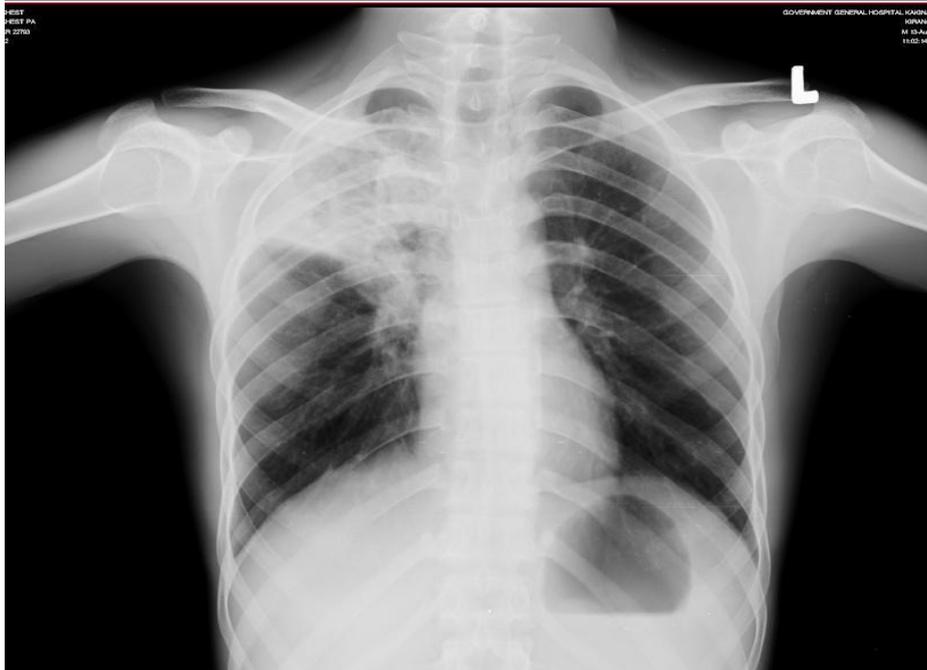


Table 13: Response to Treatment

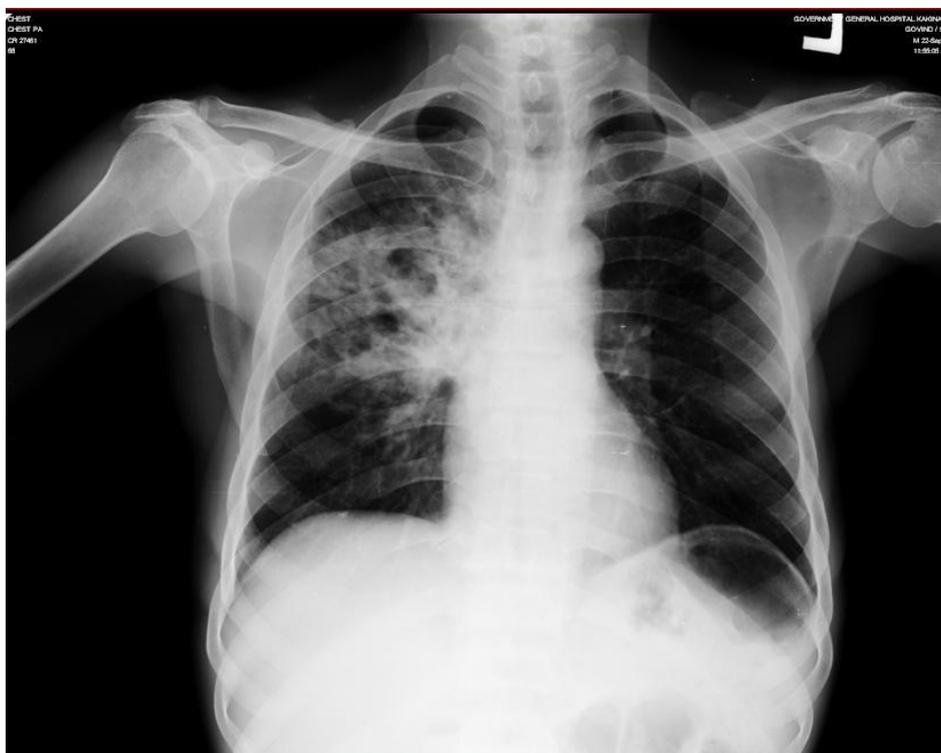
SL. NO.	RESPONSE	NO. OF CASES	PERCENTAGE
1	CURED	50	83.33
2	PARTIAL RESPONSE	8	13.33
3	DEAD	2	3.33
	TOTAL	60	100



CHEST RADIOGRAPHS



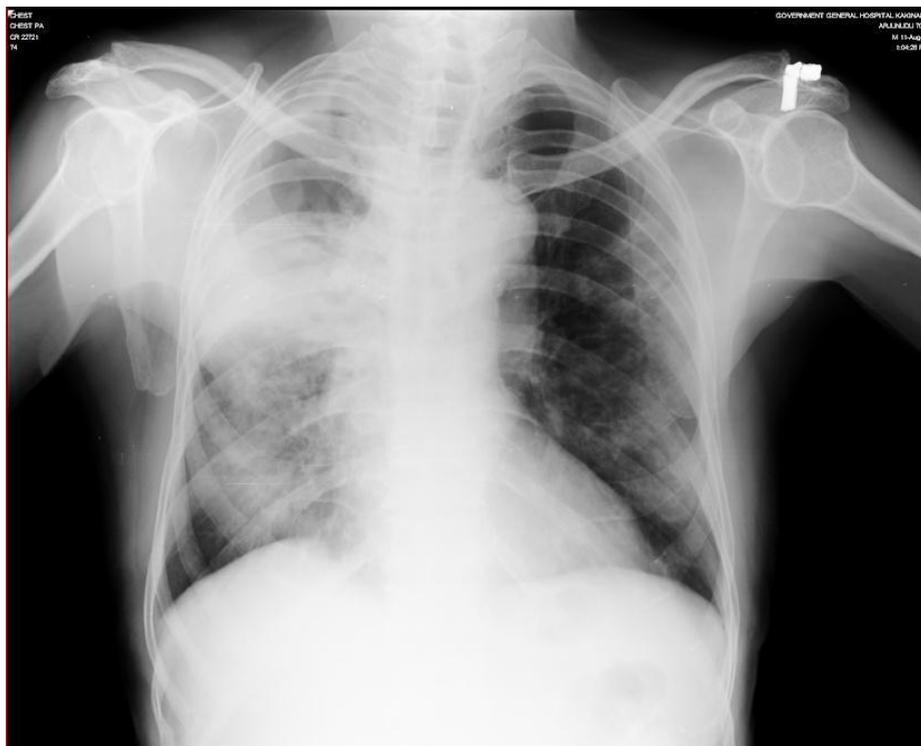
X ray chest PA view showing consolidation Rt. Upper zone



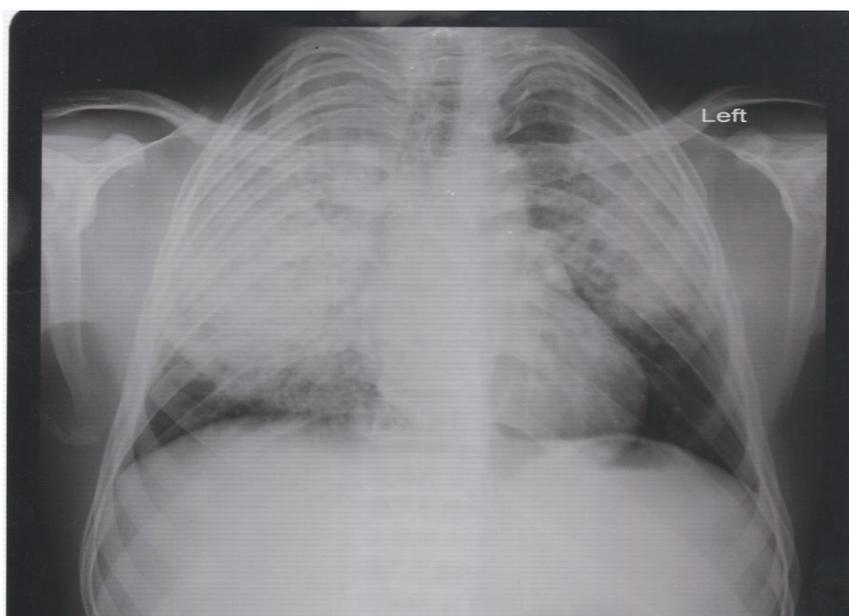
X ray chest PA view showing pneumonia Rt. Middle zone



X ray chest PA view showing Rt. Lower zone consolidation



X ray chest PA view showing consolidation Rt. middle and lower zones



Chest X ray PA view showing bilateral pneumonic consolidation

V. Discussion

Community Acquired Pneumonia (CAP) is a common medical problem in tropical countries like India. This prospective study of Community Acquired Pneumonia consisted of 60 patients admitted in Maharaja Institute of Medical Sciences, Nellimarla, Vizianagaram district; AP. All cases met inclusion and exclusion criteria.

There are many studies done in different parts of the world on community acquired pneumonia. Few of the important studies are quoted for comparison and discussion.

5.1. AGE:

It is well documented that pneumonia is commonly occurring disease in the community & its incidence rises sharply with extremes of age.^{17,25}

The present study included patients of CAP with age ranging from 19- 80 years with mean age of 48.27 ± 13.69 years which is similar to the studies done by Shrestha R et al⁶⁸ (51.3 yrs), Bansal S et al⁵³ (52.7 ± 18.1 yrs) and Shah et al⁶³ (53.68 ± 14.74 yrs). The mean age of presentation in males is 49.77 ± 13.66 yrs whereas in females it is 42.85 ± 12.88 yrs. This is comparable to the study done by Shah et al⁵⁵ (60.8 ± 13.6 yrs in males and 48.3 ± 17 yrs in females).

In this study, majority of patients with CAP were middle aged and elderly (71% are more than 50 years of age). Similar age distribution was also observed by Dey et al⁷ and Shah et al⁶³ in their study of CAP with 59% and 67% of patients respectively in the age group of > 50 years.

5.2. SEX INCIDENCE:

In this study of 60 patients of CAP, it was observed that majority of patients are males (78.33%) compared to the females (21.66%) with a male to female ratio of 3.6: 1. Similar male preponderance of CAP was also observed by various studies like Joshua et al²², Shah et al⁶³, Aroma oberoio et al⁶⁰ and Shrestha R et al⁶⁸.

This could be attributed to the well-established fact that cigarette smoking and alcoholism, as well as underlying lung disease e.g. COPD predispose to pneumonia and are more common in males. In this study group majority of male patients are exposed to one or more of the above mentioned predisposing factors.

5.3. URBAN AND RURAL POPULATION:

In this study, 68.33% of the patients of CAP are from rural area and majority was daily wage workers and manual laborers belonging to low socio-economic status. Remaining 31.66% of patients belong to urban population.

The increased frequency of lung infection in rural population may be attributed to malnutrition, poverty, overcrowding and low socio-economic status.

5.4. DURATION OF HOSPITAL STAY:

In our study, most of the patients (83.33 %) required hospitalization for less than 10 days, 11.66 % required stay between 11-15 days and only 5% for more than 15 days with the mean duration of 7.74days. This is in par with the study conducted by Bansal S et al⁵³ (9.11 days) and Shrestha et al⁶⁸ (8 days). The contribution of pneumonia by virtue of man hours among workers is significant. Most of the patients with longer duration of hospital stay have developed either pulmonary complications or systemic complications.

5.5. PREDISPOSING FACTORS:

In the analysis of various predisposing factors in the cases taken up for this study, smoking proved to be a significant predisposing factor for CAP accounting for 36.66% of cases. This correlates very well with the Bochud study⁴², which mentioned the incidence of smoking as 35.3% among its cohorts. Incidence of other predisposing factors of CAP in present study like alcoholism (21.66%), COPD (18.33%) and Diabetes (10%) is comparable with Bochud study⁴² which observed alcoholism in 12.9%, COPD in 6.5% and diabetes in 2.9% of CAP patients. Other studies by Aroma oberoi et al⁶⁰ and Shah et al⁶³ also observed smoking as the most common risk factor for CAP (26.6% and 65% respectively).

Table14 showing comparison of predisposing factors in various studies

Factor	Bochud et al ⁴² 2001	Aroma oberoi et al ⁶⁰ 2006	Shah et al ⁶³ 2010	Present study
Smoking	35.3%	26.6%	65%	36.66%
COPD	6.5%	14%	57%	18.33%
Alcoholism	12.9%	23%	1%	21.66%
Diabetes	2.9%	13.7%	13%	10%

Smoking is a well-known and important risk factor for CAP through alterations in mechanisms of the host defence system⁶⁴. Almirallet al⁶⁵ reported that even in persons without COPD, the proportion of CAP cases attributable in the population to ever having consumed any type of tobacco was 23% (95% confidence intervals 3.3 to 42.7%). Alterations in the immune system and inflammatory mechanisms in smokers are well known. Tobacco smoking is the most important risk factor for the development of COPD⁶⁶ and these patients were known to be at a higher risk of pneumonia and other respiratory infections. Both smoking and COPD are predisposing risk factors for CAP⁶⁷. These patients have altered mucocilliary mechanisms in the lower respiratory tract and stasis of secretions in lung parenchyma, which predispose to pneumonia. COPD patients who are on regular inhaled corticosteroids are also susceptible to respiratory infections.

Diabetes is also one of the risk factors for community acquired pneumonia. Diabetes is a predisposing factor for various infections in the body by altering immune and inflammatory mechanisms. In the present study 10% of the patients have diabetes mellitus as predisposing factor for CAP.

5.6. PRESENTING COMPLAINTS:

Fever and cough (100% each) are the most common presenting symptoms of CAP in the present study. Other presenting symptoms include expectoration (86.66%), breathlessness (81.66%) and chest pain (63.33%). Similar incidence of presenting symptoms of CAP was also observed in other studies done by Mac Fariene et al³⁰, Joshua. P. et al²², Bansal et al⁵³ and Shah et al⁶³.

Table 15 showing Comparison of Presenting Symptoms in various studies

Symptoms	Mac Fariene Study ³⁰ 1982	Joshua P et al ²² 2003	Bansal et al ⁵³ 2004	Shah et al ⁶³ 2010	Present study
Fever	86%	88%	90%	95%	100%
Cough	92%	92%	97%	99%	100%
Expectoration	54%	65%	87%	65%	86.66%
Dyspnoea	67%	71%	48%	-	81.66%
Chest pain	62%	-	34%	75%	63.33%

5.7. GENERAL PHYSICAL EXAMINATION:

In this study, pallor is the most common general examination finding observed (23.33%) followed by cyanosis (8.33%), icterus (3.33%), clubbing (6.66%) and pedal edema (1.66%). Pedal edema observed in one

patient who developed acute kidney injury. These findings are comparable to Bansalet al⁵³ study with pallor (11%), icterus(3%) and cyanosis(27%) among its cohorts.

Our results are also comparable to the study done by Abdullah et al⁶⁹ in elderly population which demonstrated pallor (26%), icterus(12%), cyanosis(8%), clubbing(4%) and pedal edema(8%) in its cohorts.

Anaemia seen in 23.33% of cases of CAP in our study may be an indicator of extent of malnutrition and the resultant impaired immunity in the community.

5.8. VITAL SIGNS:

In this study, examination of vital data revealed that 86.66% had tachypnea (respiratory rate > 24/min), 78.33% had tachycardia (pulse rate > 100/min), 10% had hypotension (SBP < 90 mmHg) and all the cases had elevated body temperature (> 38⁰ C).

These findings are comparable to those obtained by Abdullah et al⁶⁹ in their study of elderly population which demonstrated elevated temperature in 68%, tachypnea in 84%, tachycardia in 70% and hypotension in 16%. Several other comparable studies include those done by Bansal et al⁵³ which showed tachypnea in 24%, hypotension in 13% and Shah et al⁶³ which showed tachycardia in 92% among their patients.

The above mentioned vital signs, tachycardia, tachypnoea and high-grade fever associated with chills and rigors are well known to occur in patients with acute lung infections. Thus, meticulous recording of the respiratory rate and blood pressure at the time of initial evaluation and careful monitoring thereafter will be helpful in reducing the mortality in patients with CAP.

5.9. SYSTEMIC EXAMINATION:

In this study of CAP, the examination of respiratory system revealed various features of pneumonia like bronchial breath sounds in 88.33%, increased VF and VR in 83.33% and inspiratory crackles in 68.33% of patients. These physical signs in case of pneumonia are well documented to occur frequently as evident from other studies.

Similarly study done by Bansal et al⁵³ observed an incidence of bronchial breath sounds and crepitations in 47% and 98% of their subjects respectively. However, study done by Abdullah et al⁶⁹ observed crepitations as the common respiratory finding in 94% of their elderly subjects of CAP. The clinical findings in our study are comparable to the study done by Speteri et al as shown below.

Table 16 showing comparison of respiratory signs in various studies

Signs	Speteri et al ⁷⁰ 1988	Bansal et al ⁵³ 2003	Present study
BRONCHIAL BREATH SOUNDS	72%	47%	88.33%
INCREASED VOCAL FREMITUS	85%	-	83.33%
VOCAL RESONANCE	85%	-	83.33%
WHISPERED PECTORILOQUY	85%	-	56.66%
CREPITATIONS	72%	98%	68.33%

5.10. INVESTIGATIONS:

A. Hematological Tests

In our study of CAP, hematological investigations revealed polymorphonuclea rleukocytosis (leukocyte count >11,000 cells/mm³) in 70%, raised ESR(>20mm at 1 hour) in 73.33% and anemia(Hb< 11gm/dL)in 41.66% of the patients in the study group.

Joshua et al²² also observed similar incidence of leukocytosis in 58% of the cases while Shah et al reported an incidence of 43% in their study of CAP which are comparable to the present study.

Abdullah et al⁶⁹ reported leukocytosis in 84%, anemia in 32%, raised ESR in 76% of his elderly subjects. A high leukocyte count (>15000 cells/mm³) strongly indicates a bacterial cause.

B. Sputum Staining

In the present study of CAP, staining of sputum revealed Gram positive organisms in majority of cases (50%) compared to Gram-negative organisms (23.33%) and mixed etiology (1.66%). Similar observations were made by Larry G. Reimer et al¹⁶ in their study which found Gram positivity in 76%, Gram negativity in 14% while mixed etiology in 10% of the cases.

In our study, in about 25% of cases no organism could be seen on Gram staining and culture was negative in these cases. Probably the pathogens were atypical bacteria, fungi etc. in these cases.

In a similar study done on adult population by Abdullah et al⁶⁹, it was found that sputum shows Gram positivity in 32%, Gram negativity in 12%, mixed staining in 8% and no organism in 48% of the cases. This study is comparable to our present study.

So Gram's staining is a useful tool in predicting the probable organism before culture reports are available and these can be made use of, in initiating appropriate empirical therapy.

C. Sputum Culture

In the present study, sputum culture revealed *Streptococcus pneumoniae* as the most common pathogen in CAP accounting for 35%. Next common are *Staphylococcus aureus* (15%) and *Klebsiella* (11.66%) followed by *Pseudomonas* (8.33%) and *E.coli* (3.33%). No organisms were grown in 25% of the cultures.

These observations are similar to that of study done by Larry G. Reimer et al¹⁶ and Sanraj K. Basi et al²⁰ as depicted below.

Sputum Culture	Larry G. Reimer ¹⁶ 1998	Sanraj K. Basi ²⁰ 2004	Present study
Streptococcus	15-76%	73%	35%
Staphylococcus	3-14%	32%	15%
Klebsiella	3-14%	-	11.66%
Pseudomonas	-	8%	8.33%
E-Coli	6-20%	4%	3.33%
Mixed	-	3%	1.66%

Streptococcus pneumonia is the most common organism identified in our study. Studies from USA and UK have reported isolation rates for streptococcus ranging from 39% to 75%^{31,74,75,76}.

Our study had reported that *Stapylococcus aureus* to be the second most common isolate in 15% of the cultures. This is in par with the results obtained by Sanraj K Basi et al²⁰ (32%), Larry G Reimer¹⁶ (3-14%), Bansal et al⁵³(17%). But studies during the last three decades from India have reported a higher prevalence of Gram negative organisms among culture positive pneumonias^{54, 56, 61}.

Most of the patients from whom Gram negative bacteria was isolated were over 50 years of age, smokers or alcoholics or had COPD. It has been reported that old age, smoking, alcoholism and COPD impair pulmonary defenses predispose to CAP caused by Gram negative bacteria.

The overall identification of bacterial pathogens was 75%, which is comparable to that of Shimla⁵³ (75.6%), Chandigarh⁶⁰ (47.7), or other parts of the world like UK⁷¹(62%), Singapore⁷²(68%), and Philippines⁷³(56%). Our study is in par with many other studies in different regions of India and the world.

5.11. LOCALIZATION BASED ON CHEST X-RAY:

Our study based on chest X-ray PA view showed pneumonia to be more common on the right side (63.33%) with predominant involvement of right lower lobe in 36.66% of cases. Left side was involved in 28.33% while bilateral pneumonia was noticed in 8.33% of cases. Most commonly involved lobe is right lower lobe (36.66%) followed by left lower lobe (23.33%), right middle lobe (18.33%), left upper lobe (5%) and right upper lobe (3.33%) in decreasing order. Multilobar involvement is observed in 13.33% of cases.

Our study is comparable to the study done by Bansal et al⁵³ which demonstrated right lower lobe infiltration to be the most common (48.6%) followed by left lower lobe (21%), multilobar involvement (15.7%), right upper lobe (8.5%).

Major involvement of right lower lobe is attributed to the anatomical position of the right main bronchus, which is short, more or less vertical facilitating aspiration in to the basal bronchial segments.

5.12. ANALYSIS OF MAJOR COMPLICATIONS:

Our study showed lung complications in 6.66% of patients with CAP. Among them, 3.33% had lung abscess and 3.33% had pleural effusion. None of the patients developed empyema in our study group.

Systemic complication in the form of circulatory failure was noticed in 10% of cases, probably a result of delayed presentation to the hospital and extensive lung involvement with delayed antibiotic therapy. These patients developed sepsis and septic shock. They were treated with vasopressors and higher antibiotics.

Acute renal failure occurred in 3.33% of patients in the study group. Both these patients developed ARF because of late presentation and as they belong to elderly age group and have developed bilateral pneumonia. Both of them recovered from acute kidney injury with effective antibiotic therapy without any requirement for dialysis.

These figures are comparable to those obtained by Bansal et al⁵³ who observed circulatory failure in about 5% of their patients. Our results are also comparable to the study done by Abdullah et al⁶⁹ which reported shock(16%), effusion(12%) and lung abscess(4%) among elderly subjects.

5.13. RESPONSE TO TREATMENT:

In the present study 83.33% of cases showed good clinical response while partial response was observed in 13.33% of cases. Only 2 patients with circulatory failure expired while on treatment with a mortality rate of 3.33%. Both of them belong to elderly age group and they have predisposing factors like COPD and smoking. The organisms isolated in both of them are staphylococcus aureus and pseudomonas aureginosa. Good clinical response to treatment emphasizes the curable nature of CAP.

In the study by Bochudet al⁴², 45% were cured, while 38% had minimal residual lesion and no improvement or worsening was reported in 5% of cases.

Mortality due to CAP in various hospital based studies has been variable. While the British Thoracic Society multi-centric study recorded a low mortality of 5.7%⁷⁷ which is on par with our study (3.33%), higher mortality (ranging from 21% to 25%) has been reported in other studies^{78, 79}. However, in another Indian study a significantly higher mortality was noticed in patients aged 50 years or above and in those with underlying co-morbid conditions⁵⁶. According to the study conducted by British Thoracic Society and the Public Health Laboratory Services⁸⁰, patients had a 21 fold increased risk of mortality if they had respiratory rate 30 breaths per minute or more and diastolic blood pressure less than or equal to 60 mm of Hg.

This emphasizes the need for further investigations in patients who had bad prognostic factors at the time of admission so as to establish the etiology, start early treatment and thereby reducing mortality.

Our observations will also be useful to monitor the trends of CAP in the population of the region and will help the physicians to start rational empirical treatment for patients with CAP.

VI. Conclusion

In essence, CAP is a common malady affecting elderly, rural, males with history of smoking, COPD and alcoholism. It commonly presents with all the classical symptoms and signs of pneumonic consolidation. Cases presenting with peripheral circulatory failure have a higher mortality.

Majority of the pneumonias are caused by *Streptococcus and Staphylococcus* in our study. Clinical and radiological response to a combination of Betalactam and macrolides or fluroquinolones was excellent, requiring less than a 10 days stay as an inpatient. Finally, cessation of smoking and alcoholism, early detection of cases with appropriate empirical treatment pending laboratory reports for as little as a week will go a long way in curing this condition.

References

- [1]. Seaton A, Seaton D. and AG Leich, Crofton & Douglas's. Respiratory Diseases. 5th edition. Vol-1 Chapter 13: 356-429.
- [2]. Lionel A.Mandell, Richard Wunderink, Harrison's Principles of Internal Medicine, 18th edition Pneumonia, Chapter 257, page 2130-2141.
- [3]. PaiDhungat AJ. API Text Book of Medicine Pneumonias, 7th edition, Chapter 7 page 301- 305.
- [4]. John G. Berlett. Oxford Text Book of Medicine- Pneumonia- Normal host, 4th edition chapter 17, 1357-1367.
- [5]. Medical Clinics of North America (Sep. 1993): Pneumonia.
- [6]. Garibaldi RA. Epidemiology of community acquired respiratory tract infection in adults: incidence, etiology and impact Am J Med 1985; 78: Suppl 6b, 32-37.
- [7]. Dey et al. clinical presentation and predictors of outcome inn adult patients with community-acquired pneumonia. Natl Med-India. 1997 July-Aug; 104: 169-172.
- [8]. Metlay JP et al. Does this patient have CAP? Diagnosing pneumonia by history and physical examination. JAMA 1997; 278: 1440-1445.
- [9]. David Jonson Gray's anatomy. The anatomical basis of clinical practice, 39th edition, Chapter 56, 62 and 63. Anatomy of thorax, lungs & diaphragm.
- [10]. BD Chaurasia. Human anatomy, regional and applied, chapter-16, The lungs: 199-207.
- [11]. William F. Ganong, Medical physiology Respiration, Chapter 34, Pulmonary Function 21st edition, 649-698.
- [12]. Metlay J. P, Schulz R. et al. Influence of age on symptoms at presentation in patients with community acquired pneumonia. JAMA & ARCHIVES - Arch Intern Medicine Vol. 157W13, July 14, 1997
- [13]. John Bartlet Treatment of Community Acquired Pneumonia, Chemotherapy 2000; 46 (suppl 1): 24-31.
- [14]. Niederman et al. Guidelines for the management of adults with Community acquired pneumonia. Assessment of severity, antimicrobial therapy, and prevention. Am J Respir Crit Care Med 2001; 163: 1730-1754.
- [15]. Ananthanrayana R and Jayaram Paniker CK. Part III, Textbook of Microbiology. Page 178-389.
- [16]. Larry G. Reimer and Karen C. Carrol, Role of the microbiology laboratory in the diagnosis of lower respiratory tract infection, Clinical Infectious Diseases 1998; 26: 742-8.
- [17]. John G. Barlett, Scott F. et al. Practice guidelines for the management of CAP in adults. Clin Infect Dis 2000; 31: 347-82.
- [18]. John G. Bartlett and Linda M. Mundy. Community Acquired Pneumonia NEJM. Dec 1995; 333; 1618-1624.
- [19]. Roger G. et al. Practical considerations and guidelines for the management of community acquired pneumonia. Drugs, Jan 1998; 31-43.
- [20]. Sanraj K. Basi et al. Patients admitted to hospital with suspected pneumonia and normal chest radiographs; epidemiology, microbiology, and outcomes, American Journal of Medicine, 1st Sept 2004; 117: 305-311.
- [21]. Thomas P et al. seminar on community-acquired pneumonia Lancet 2003; 362.
- [22]. Joshua P. Metlay and Micheal J. Fine testing strategies in the initial management of patients with CAP. Annals of Internal medicine Jan, 2003; 138, 2: CINAHL.
- [23]. Jose Vilar et al. Radiology of bacterial pneumonia. Eur J Radiol 51 (2004)102-113.
- [24]. Mandell LA, Marrie TJ, et al Canadian guidelines for the initial management of CAP. Clin Infect Dis. 2000; 383-421.

- [25]. Fine MJ et al. Prognosis and outcomes of patients with CAP. A meta-analysis-JAMA;257 (2): Jan 1996.
- [26]. Michiel J. Fine et al. A prediction rule to identify low risk patients with community acquired pneumonia. NEJM, Jan 1997; 336; 243-250.
- [27]. Hanry F. Chambers Goodman & Gilman's Pharmacological Basis of Therapeutics Anti-microbial agents-General considerations, 10th Edition, 1143-1163.
- [28]. Robert E. Siegel et al. A prospective randomized study of inpatients, IV antibiotics for CAP.CHEST/110/4/OCT, 1996.
- [29]. T. Franquet. Imaging of pneumonia: Trends and algorithms. Euro Respire J 2001; 18; 196-208.
- [30]. Macfartane JT, et al. Hospital study adult community acquired pneumonia. The community, Lancet. 1982; 2: 255-8.
- [31]. Humphry JH et al. Pneumonia in North London, Thorax 1948; 1: 314.
- [32]. Carpenter JL and Huang DY. Community-acquired pneumonia in a public municipal hospital in 1980s. South Med J 1991; 84: 299-306.
- [33]. Ewig S, Torres A. Severe CAP.Clin Chest Med 1999; 20: 575-97.
- [34]. Fick RB Jr. Reynolds HY. Changing spectrum of pneumonia - News media creation or clinical reality? Am J Med 1983; 74: 1-8.
- [35]. Larsen RA, Jacobson JA. Diagnosis of community-acquired pneumonia experience at a community hospital. Compr. Ther. 1984; 10: 20-25.
- [36]. Lim WS, Baudouin SV, George RC, Hill AT, Jamieson C, Le Jeune I, et al. BTS guidelines for the management of community acquired pneumonia in adults: update 2009. Thorax 2009;64(Suppl 3):iii1-55.
- [37]. Marcos I. Restrepo, MD, MS, Antonio Anzueto, MD Infect Dis Clin N Am 23 (2009) 503–520 doi:10.1016/j.idc.2009.04.003.
- [38]. World Health Organization. Disease and injury country estimates. http://www.who.int/healthinfo/global_burden_disease/estimates_country/en/index.html.
- [39]. Guidelines for diagnosis and management of community-and hospital-acquired pneumonia in adults: Joint ICS/NCCP(I) recommendations Dheeraj Guptaetal., for the Pneumonia Guidelines Working Group, Department of Pulmonary Medicine, Postgraduate Institute of Medical Education and Research (PGIMER), Chandigarh, Indian Chest Society, National College of Chest Physicians, India.
- [40]. Infectious Diseases Society of America/American Thoracic Society Consensus Guidelines on the Management of Community-Acquired Pneumonia in Adults .JDSA/ATS Guidelines for CAP in Adults • CID 2007;44 (Suppl 2).
- [41]. SUPPLEMENT TO JAPI • JANUARY 2012 • VOL. 60, Management of Community Acquired Pneumonia, Randeep Guleria, Jaya Kumar.
- [42]. Bochud P.Y, Moser F., Erard P, et al. Community-acquired pneumonia. Prospective study. Medicine 2001; 80(2):75-87.
- [43]. Karetzky M. Community-acquired pneumonia. In: Brandstetter RD, Karetzky M, Cunha BA, editors. The Pneumonias. New York: Springer-Verlag; 1993:pp 25-48.
- [44]. Regional situation on health statistics reporting. Health Situation in the South-East Asia Region 1994-1997. New Delhi: EHI/WHO-SEARO. September 2007.
- [45]. Tejerina E, Frutos V, Restrepo MI, et al. Prognosis factors and outcome of community acquired pneumonia needing mechanical ventilation. J Crit Care2005; 20(3):56–65.
- [46]. Marrie TJ, Carriere KC, Jin Y, et al. Factors associated with death among adults<55 years of age hospitalized for community-acquired pneumonia. Clin Infect Dis 2003; 36(4):413–21.
- [47]. Pascual FE, Matthay MA, Bacchetti P, et al. Assessment of prognosis in patients with community-acquired pneumonia who require mechanical ventilation. Chest 2000; 117(2):503–12.
- [48]. Mehta R, Groth M. Clinical application of a prognostic model for severe community-acquired pneumonia. Chest 2001; 119(1):312–3.
- [49]. Meehan TP, Fine MJ, Krumholz HM, et al. Quality of care, process, and outcomes in elderly patients with pneumonia. JAMA 1997; 278 (23): 2080–4.
- [50]. Mortensen EM, Coley CM, Singer DE, et al. Causes of death for patients with community-acquired pneumonia: results from the Pneumonia Patient Outcomes Research Team cohort study. Arch Intern Med 2002; 162(9):1059–64.
- [51]. Ishida T, Hashimoto T, Arita M, Ito I, Osawa M. Etiology of community-acquired pneumonia in hospitalized patients: a 3-year prospective study in Japan. Chest 1998; 114:588-93.
- [52]. Lieberman D, Schalaeffer F, Boldur I, Liebermam D, Horowitz, Friedman MG, et al. Multiple pathogens in adult patients admitted with community acquired pneumonia: a one year prospective study of 346 consecutive patients. Thorax 1996;51:179-84.
- [53]. Bansal S, Kashyap S, Pal LS, GoelA. Clinical and bacteriological profile of community acquired pneumonia in Shimla, Himachal Pradesh. Indian J Chest Dis Allied Sci 2004;46:17-22.
- [54]. Kulpati DDS, Khastgir T. Reappraisal of pneumonias. JAPI 1988;36:660-4.
- [55]. Shah BA, Ahmed W, Dhobi GN, Shah NN, Khursheed SQ, Haq I. Validity of pneumonia severity index and CURB-65 severity scoring systems in community acquired pneumonia in an Indian setting. Indian J Chest Dis Allied Sci 2010;52:9-17.
- [56]. Dey AB, Chaudhry R, Kumar P, Nisar N, Nagarkar KM. Mycoplasma pneumoniae and community-acquired pneumonia. Natl Med J India 2000;13:66-70.
- [57]. Capoor MR, Nair D, Aggarwal P, Gupta B. Rapid diagnosis of community-acquired pneumonia using the BacT/Alert 3D system. Braz J Infect Dis 2006; 10:352-6.
- [58]. Samuel KM. Notes on Clinical Laboratory Techniques. 4thedn. Madras: MGK Iyyer& Sons; 1986 : 168.
- [59]. Donalísio MR, Arca CH, Madureira PR. Clinical, epidemiological, and etiological profile of inpatients with community-acquired pneumonia at a general hospital in the Sumaremicroregion of Brazil. J Bras Pneumol 2011;37:200-8.
- [60]. Aroma Oberoi et al., Bacteriological Profile, Serology and Antibiotic Sensitivity Pattern of Micro-organisms from Community Acquired Pneumonia JK SCIENCE Vol. 8 No. 2, April-June 2006.
- [61]. Madhu SB, Gupta U, Guleria JS, Talwar V. Clinical and bacteriological profile of hospitalized community – acquired pneumonias a preliminary study. *Ind J Chest Dis & All Sci*1990 ; 32 (2) : 96-100.
- [62]. Woodhead MA, MacFarlane JT, McCracken JS, Rose DH, Finch RG. Prospective study of the etiology and outcome of pneumonia in the community. *Lancet* 1987 ; 1 : 671-74.
- [63]. Shah et al., Bacteriological and clinical profile of CAP in hospitalized patients. Lung india.vol 27, issue 2, Apr – Jun 2010.
- [64]. Marcy TW, Merrill WW. Cigarette smoking and respiratory tract infection. *Clin Chest Med* 1987; 8 : 381-91.
- [65]. Almirall J, Gonzalez CA, Balanzo X, Bolibar I. Proportion of community acquired pneumonia attributable to tobacco smoking. *Chest* 1999; 116 : 375-79.
- [66]. Sherman CB. The health consequences of cigarette smoking: Pulmonary diseases. *Med Clin North Am* 1992; 76 : 355-75.

- [67]. Ginesu F, Pirina P. Etiology and risk factors of adult pneumonia. *J Chemother* 1995; 7 : 277-85.
- [68]. Shrestha R, Paudel N, Barakoti B, Dhungana D, Sharma P. Etiology and clinical profile of inpatients with community acquired pneumonia in Manipal teaching hospital, Pokhara, Nepal. *Nepal Journal of Medical sciences* 2012;1(2):84-8.
- [69]. A Study of Community-Acquired Pneumonias in Elderly Individuals in Bijapur, India. *International Scholarly Research Network, ISRN Pulmonology*, Volume 2012, Article ID 936790
- [70]. Spiteri MA, Cook DG, Clarke SW. Reliability of eliciting physical signs in examination of the chest. *Lancet* 1988;1:873-5.
- [71]. Howard LS, Sillis M, Pasteur MC, Kamath AV, Harrison BD. Microbiological profile of community-acquired pneumonia in adults over the last 20 years. *J Infect* 2005; 50:107-113.
- [72]. Lee KH, Hui KP, Tan WC, Lim TK. Severe Community-acquired Pneumonia in Singapore. *Singapore Med J* 1996;37:374-7.
- [73]. Ong G, Antonio-Velmonte M, Mendoza MT. Etiologic agents of community acquired pneumonia in adults: The PGH experience. *Phil J Microbiol Infect Dis* 1995; 24:29-32.
- [74]. Bath JC, Boissard GP, Caldre MA Wood SC, Rowansky MJ, Chanock RM. Pneumonia in hospital practice in Edinburgh. *Br J Dis Chest* 1964; 58 : 1-16.
- [75]. Mufson MA, Chang V, Gill V, et al. The role of viruses, mycoplasma and bacteria in acute pneumonia in civilian adults. *Am J Epidemiol* 1976;86: 526-44.
- [76]. Dorff GJ, Rytel MW, Farmer SG, Scanlon G. Etiologies and characteristic features of pneumonia in a munipicle hospital. *Am J Med Sci* 1973; 266: 349-58.
- [77]. Mac Farlance J. Community acquired pneumonia. *Br J Dis Chest* 1987; 81 : 116-27.
- [78]. Ortqvist A, Hedlund J, Grillner L, et al. Aetiology outcome and prognostic factors in community acquired pneumonia requiring hospitalization. *Eur Respir J* 1990; 3: 1105-13.
- [79]. Pachon J, Prados MD, Capote F, Cuello JA, Garnacho J, Verano A. Severe Community acquired pneumonia : Etiology, prognosis and treatment. *Am Rev Respir Dis* 1990; 142: 369-73.
- [80]. Research Committee of the British Thoracic Society and the Public Health Laboratory Service. Community acquired pneumonia in adults in British hospitals in 1982-83: A survey of aetiology, mortality, prognosis factors and outcome. *Q J Med* 1987; 62 : 195-220.

Dr.Kaliparambil Sugathan Roshni "Clinical, Microbiological and radiological study of community acquired Pneumonia". IOSR Journal of Dental and Medical Sciences (IOSR-JDMS), vol. 17, no. 2, 2018, pp. 45-63.