



Low Dose Computed Tomography (LDCT) Versus Chest X-ray in Chronic Smokers- Can LDCT be Used as a Motivation to Quit Smoking?

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ABSTRACT

Background- Smoking is one of the biggest health threats to mankind. Early diagnosis of pulmonary changes induced by smoking can help in timely intervention and prediction of course of smoking-induced diseases.

Aim-The aim of this study was to determine the relative prevalence of various common radiological findings associated with smoking and compare their detection on digital chest X-ray and Low Dose Computed Tomography (LDCT)

Materials and Methods-Chest X-ray PA view and LDCT scan of 50 chronic symptomatic/asymptomatic male smokers were analysed over a period of 1 year from February 2016 to February 2017.

Summary- Chest X-ray was less sensitive in detecting early emphysematous changes, small pulmonary nodules and minimal fibro-bronchiectatic changes. In our study, spectrum of respiratory findings in smokers on LDCT included bronchial thickening (68%) emphysema (66%), interstitial thickening (61%), pulmonary micro- and macro- nodules, air space opacification, bronchiectasis, mediastinal lymphadenopathy and ground-glass opacities. The sensitivity of LDCT in diagnosing malignant lesions was 88.4% and specificity was 82.5%. The sensitivity of Chest Xray in diagnosing malignant lesions was 50% and specificity was 87.5%.

Conclusion- LDCT diagnoses malignant lesions much earlier and provides a better graphical picture of the pathology under study to young and middle aged smokers willing to quit smoking. Since the baseline risk of lung cancer development is small and the dose exposure in a single LDCT scan is well below the annual adult radiation dose limit, the risk-benefit ratio is very favourable.

Keywords-smoking, smoking-induced diseases, Chest X-ray, LDCT, risk-benefit ratio.

Introduction

The tobacco epidemic is one of the biggest public health threats the world has ever faced. Nearly 80% of more than 1 billion smokers worldwide live in low-and middle income countries, where the burden of tobacco-related illnesses and death is heaviest. Smoking is responsible for more than 80% of all lung cancers worldwide.

Smoking related diseases can be classified as- Interstitial lung diseases (Respiratory bronchiolitis,

Desquamative interstitial pneumonia, Pulmonary Langerhans histiocytosis, Acute eosinophilic pneumonia, Combined pulmonary fibrosis and emphysema), Neoplasms (Lung cancer, Tracheal tumours) and Chronic Obstructive Pulmonary Disease (Emphysema, Chronic bronchitis, Constrictive bronchiolitis).²

The routine protocol of most clinicians in a patient presenting with respiratory complaints is to order a preliminary Chest Xray PA view.³ However, since it

is a two-dimensional imaging modality, much information is lost due to overlap of organs, patient motion and inadequate breath-hold. Hence, early and subtle changes are often missed and the patient is falsely convinced of being disease-free.⁴

The role of LDCT in lung cancer screening is established by NLST (National Lung Screening Trial).⁵ Screening is the testing of an individual who is at risk for a disease, but who does not exhibit the signs of the disease.⁶ It involves scanning of an individual with average dose of 1.5 to 3.4 mSv (further improvements have led to further dose reduction upto a level of 0.2mSv) which is 4 to 12 times more than Chest X-ray and 5 times less than conventional CT chest.⁷

LDCT is comparable in sensitivity and specificity of lung nodule detection with conventional chest CT.⁶ In addition it offers all other advantages of a CT scan such as three dimensional reconstruction, single breath-hold and exploits the inherent high contrast of thoracic structures. However, it suffers from over-diagnosis and a high false positive rate.⁸

Since cigarette smoking results in a focal accumulation of macrophages within the walls of respiratory bronchioles and adjacent alveoli, the common finding on chest X-ray is of “dirty chest”, an overall increase in non-specific lung markings. Remy Jardin et al⁹ found that the morphological substrate of cigarette smoking- induced changes in chest radiography were parenchymal micro-nodules and intra-lobular opacities. Destructive process such as emphysema results in increased parenchymal lucency with preserved or increased lung volumes.

In a study by J Kirchner et al,⁹ there was a significant positive correlation between the increase of overall lung markings on chest radiography and the cigarette consumption quantified by pack-years. There is a positive linear correlation between bronchial wall thickening and intra-lobular opacities as seen in CT with the increase of overall lung markings on radiography.

Aims

- To study the effects of smoking on respiratory system.

- To determine the relative prevalence of various common radiological findings associated with smoking and to correlate it with the amount of cigarette consumption measured in pack-years.
- To study and compare the findings on two modalities – Digital X-ray Chest and LDCT.
- To calculate the risk-benefit ratio of using either modality in terms of its sensitivity, specificity and dose of ionizing radiation.
- To target especially young chronic smokers and explain them the effects of smoking to their lungs, the importance of early diagnosis and motivate them to quit smoking.

Materials and Methods

The present study was conducted in a tertiary care center with 50 subjects from February 2016 to February 2017.

Description of tools:

- Digital X-rays (PA view of chest) procured through SIEMENS mm-10 160mAs Xray (HF) machine
- Non-contiguous Low dose computed tomography axial cuts with reformation procured through 16 slice MDCT machine (GE healthcare).

Inclusion Criteria

1. Age : 35 years or more
2. Sex : male/female
3. Active smoking history of 10 years or more (present or past).
4. Symptomatic/asymptomatic.

Exclusion criteria

1. Known case of primary/secondary lung cancer or any chronic lung disease.
2. No previous CT imaging for lung pathology.
3. Scanning protocol

For conventional radiography

- kvp 40-60
- mAs 20-30
- Exposure dose on an average 0.02 mSeivert

For low dose computed tomography:

- Number of detectors: 16
- kVp : 80

- mA : 30–50 CT-AEC (SD = 70)
- Seconds/rotation : 0.75
- mAs : 22.5–37.5
- Pitch factor : 1.438
- Collimation : 1 mm× 16
- Reconstruction slice thickness (mm) : 1.5 mm
- Slice interval (mm) : 3
- Lung field : 1600/-600
- Mediastinal : 400/35
- Exposure dose(range): 1.5 -3.4 mSv
- Area covered: From apex of lung to diaphragm.

The number of true positives and negatives were calculated for both Chest Xray and LDCT based on clinico-histopathological correlation. Based on it, the sensitivity and specificity of both diagnostic modalities were calculated. Chi-square test was applied for association of amount of cigarette consumption with pack-years with value of $X^2 = 4.23$, p value <0.05 and degrees of freedom: 1.

Observations

The distribution of various pulmonary findings, relative prevalence, detectability on Chest x-ray and LDCT is as described below-

Table 1 % Distribution of smokers according to pack-years

Smoking pack-years	No. of patients (n=50)	%Percentage
11 – 15	6	12
16 – 20	6	12
21 – 25	11	22
26 – 30	11	22
31 - 35	8	16
36 - 40	6	12
41 - 45	2	4

Majority of smokers had a smoking exposure of 21-30 pack years.

Table 2 % Distribution according to symptoms

Symptom	No. of patients (n=50)	% Percentage
Dry cough	34	68
Cough with expectoration	17	34
Dyspnoea	32	64
Hemoptysis	9	18
Weight loss	8	16
Fever	1	2
Chest pain	2	4

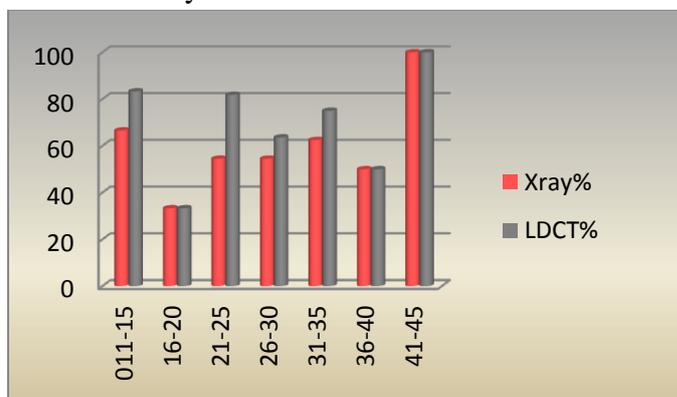
Most common presenting complaint was dry cough followed by dyspnoea.

Table 3 % Distribution of common radiological findings

Radiological finding	Chest X-ray(n=50)		LDCT(n=50)	
	No.	%	No.	%
Bronchial thickening	28	56	34	68
Interstitial thickening	20	40	30	60
Emphysema	21	42	33	66
Pulmonary nodules	16	32	25	50
Areas of air space opacification	7	14	9	18
Consolidation	6	12	5	10
Ground-glass opacities	0	0	4	8
Mediastinal lymphadenopathy	10	20	16	32
Pleural effusion	2	4	2	4

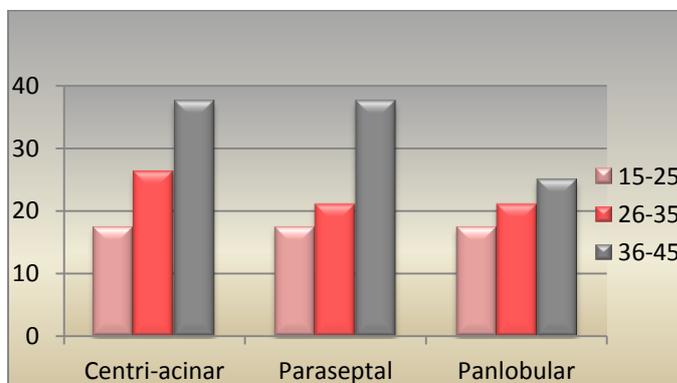
The three most common findings were the presence of bronchial thickening, emphysema and interstitial septal thickening.

Figure 1 % of prevalence of bronchial thickening on Chest X-ray and LDCT



There was no significant difference between moderate and severe consumption of smoking as regards to prevalence of bronchial thickening on chest X-ray and LDCT.

Figure 2 % Distribution of different types of emphysema according to pack years



All types of emphysema were strongly related to smoking with a positive correlation to amount of cigarette consumption measured in pack-years.

Figure 3 % Distribution according to number, size and consistency of nodules

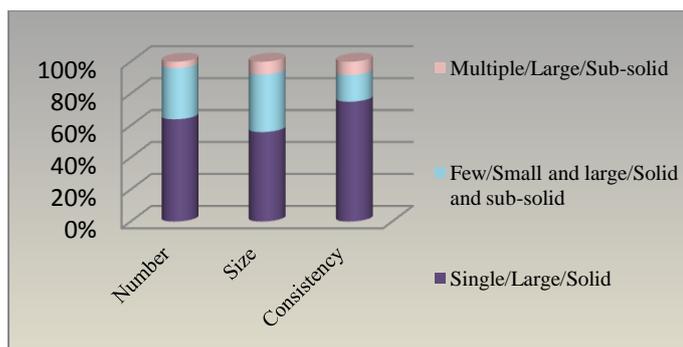
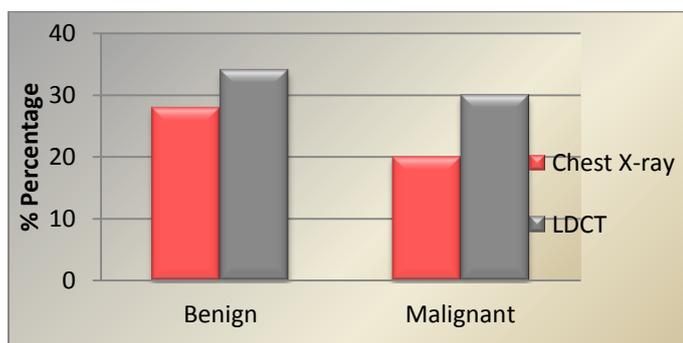
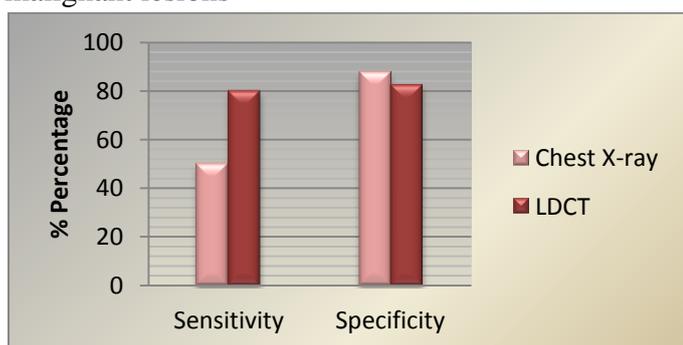


Figure 4 % Distribution of benign and malignant findings on Chest X-ray and LDCT



The proportion of benign lesions in smokers was higher than malignant lesions by diagnosed by both Chest X-ray and LDCT. LDCT could diagnose more benign and malignant lesions than chest X-ray.

Figure 5 Comparison between sensitivity and specificity of chest X-ray and LDCT in diagnosing malignant lesions

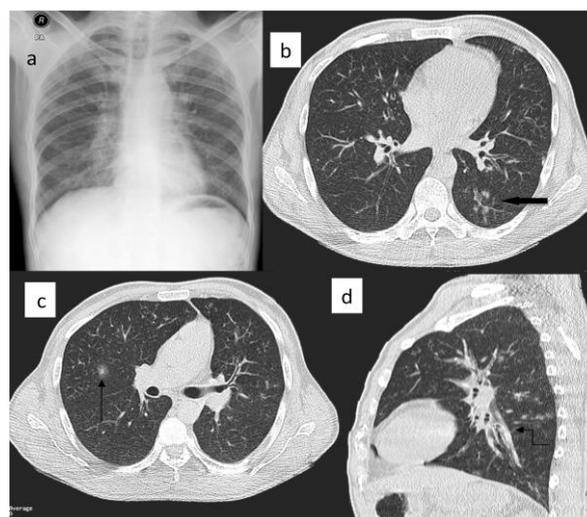


LDCT was sensitive in diagnosis of malignant lesions as compared to Chest X-ray, however, its specificity was slightly lower as compared to chest X-ray. With modern multi-detector CT, pulmonary nodules are detectable at a size of less than 2 mm, however, most of them are benign giving rise to false positive results

Table 4 Dose exposure in chest X-ray and LDCT

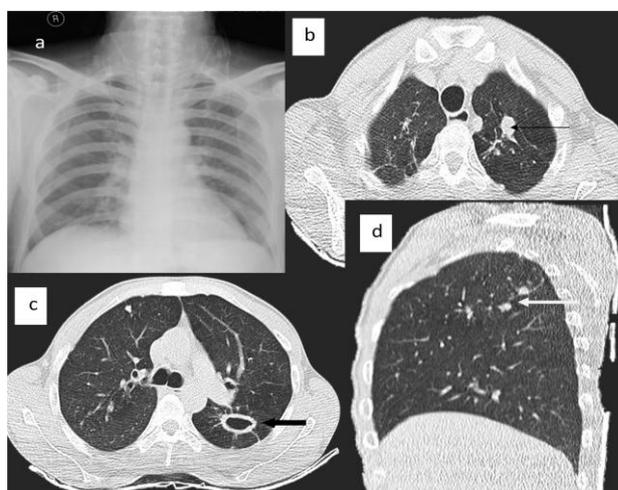
Dose range (X-ray)(mSv)	No.of patients	Dose range (LDCT) (mSv)	No.of patients
0.18	7	1.4	1
0.19	7	1.5	10
0.2	11	1.6	9
0.21	7	1.7	8
0.22	11	1.8	11
0.23	4	1.9	9
0.24	3	2	2
Total	50	-	50

Figure 6 Multiple nodules with interstitial septal thickening



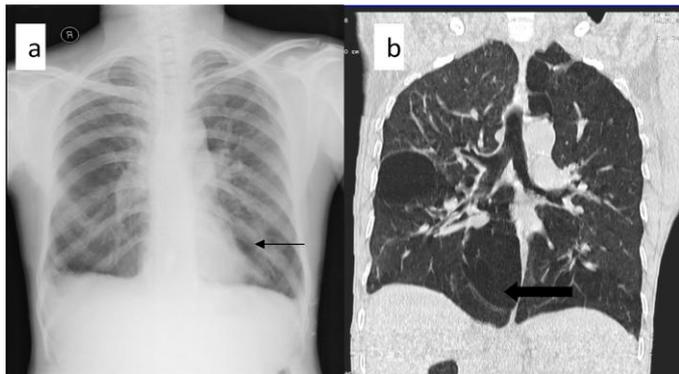
45 year old smoker with dry cough, chest X-ray (figure a) appears unremarkable. LDCT axial scans (figures b and c) show few solid micro-nodules in superior segment of left lower lobe in perilymphatic distribution (thick arrow) and a sub-solid nodule in anterior segment of right upper lobe (thin arrow). Coronal reformatted image (figure d) shows peri-bronchovascular cuffing (elbowed arrow).

Figure 7 Multiple parenchymal nodules with cavity formation



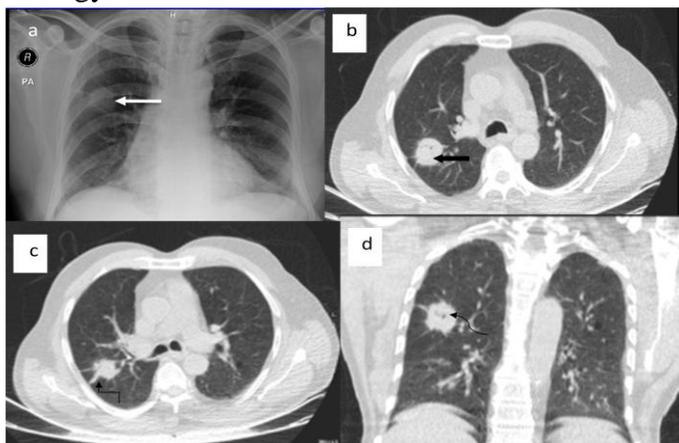
40 year old male smoker with complaints of dry cough. Chest X-ray (figure a) appears unremarkable. LDCT images (figures b and d) show few irregularly marginated solid nodules (thin arrows) in random distribution in apicoposterior segment of left upper lobe, (figure c) shows a cavitary lesion with outer irregular and inner smooth margin with perilesional interstitial thickening (thick arrow) in apico-posterior segment of left upper lobe.

Figure 8 Bullous emphysematous changes



38 year old male smoker with recurrent breathlessness. Chest X-ray (figure a) shows patchy areas of radiolucency in bilateral lungs (thin arrow). LDCT coronal image (figure b) shows moderate centri-acinar and panacinar emphysema with bulla formation (thick arrow).

Figure 9 Solitary pulmonary nodule, malignant etiology



50 year old with hemoptysis. Chest X-ray (Figure a) shows a rounded soft tissue radio-opacity in right upper lung zone. LDCT images (figures b, c and d) show a spiculated marginated nodule (thick arrow) with perilesional nodular thickening (elbowed arrow)

suggestive of lymphangitic spread and central necrosis (curved arrow).

Discussion

In our study 50 chronic smokers were subjected to both chest X-ray and LDCT according to standard protocols to evaluate and compare the spectrum of respiratory findings associated with smoking. Various recent studies on epidemiology of smoking related diseases, radiological patterns, correlation between tobacco burden and radiographic appearances, dose exposure were kept as reference.¹⁰

Majority of smokers (44%) had a smoking exposure equivalent to 21-30 pack-years and all smokers were males reflecting that smoking habit is widely prevalent among Indian men than women. No female smoker attended our institution during the study period fulfilling the inclusion criteria; hence the findings were completely obtained on male sub-population. Majority of smokers were active smokers (90%).

The most common presenting symptom was dry cough (68%) followed by breathlessness (64%). The spectrum of respiratory findings in smokers included bronchial thickening, emphysema, interstitial thickening, pulmonary micro and macro-nodules, areas of air space opacification, bronchieactasis, mediastinal lymphadenopathy and ground-glass opacities. No significant correlation was found between lung cysts, consolidation, parenchymal fibrosis, honeycombing, pleural effusion and smoking. ($p < 0.001$)

Only a small proportion of LDCT scans in smokers were completely normal. Chest X-rays were normal in more number of patients (36%) as compared to LDCT (12%). LDCT was able to detect subtle findings in 24% of patients whose X-rays were considered normal.

The most common finding in smokers was the presence of bronchial thickening (68%) on both Chest X-rays and LDCT. Bronchial thickening was not associated with the amount of cigarette consumption measured in pack-years. It was present in varying proportions in mild, moderate and heavy smokers.

The second most common finding on both Chest X-ray and LDCT was the presence of Emphysema (62%). All types of emphysema were associated with smoking. The prevalence of emphysema and its severity increased with the increase in cigarette consumption measured in pack-years. LDCT (36%) was superior in diagnosing early emphysematous changes as compared to X-rays (21%).

The third most common finding was increased interstitial opacities on LDCT or increase in non-specific parenchymal markings on Chest X-ray. The prevalence of interstitial opacities showed a positive correlation with cigarette consumption measured in pack-years. ($X^2=5.6$, $p<0.01$).

Pulmonary micro-nodules showed a positive correlation with smoking. Most of the nodules detected on Chest X-ray and LDCT were of benign etiology. LDCT could diagnose more benign and malignant nodules/areas of air space opacification as compared to Chest X-rays. However, LDCT had a slightly higher false positive rate, i.e., few malignant nodules diagnosed on LDCT turned out to be of benign etiology.

LDCT was sensitive in diagnosis of malignant lesions as compared to Chest X-ray, however, its specificity was slightly lower as compared to chest X-ray.

The effective radiation dose was approximately 7 times higher in LDCT as compared to Chest X-rays. However, LDCT diagnosed potential malignant lesions much earlier and provided a better graphical picture of the pathology under study. Since the base-line risk of lung cancer development is small¹¹ and the dose exposure in a single LDCT scan is well below the annual adult radiation dose limit, the risk-benefit ratio in employing LDCT over chest X-rays is very favourable.

Early diagnosis and evidence based-counselling ultimately leads to improved patient outcome. Young and middle-aged smokers were explained the damage smoking had done to their lungs, further health implications if they continued to smoke and hence effectively dissuaded to quit smoking.

CONCLUSION

For a young to middle aged chronic smoker LDCT can serve as a convenient diagnostic modality to evaluate early pulmonary changes induced by smoking, when X-ray findings are non-specific. Further studies are required to establish cost-benefit analysis and impact on quality of life when LDCT is used as a patient counselling tool.

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