Abstract

CAD is mainly due to atherosclerosis, an inflammatory process that is based on the interaction between immune mechanisms and metabolic risk factors. Abnormal lipid levels, particularly elevated low-density lipoprotein [LDL-C] and decreased high-density lipoprotein [HDL-C], as well as higher level of inflammatory markers especially high sensitive CRP [hsCRP] are well-established risk factors for cardiovascular disease [CVD]. The present work aims to study the relation between lipid profile, high sensitive C-reactive protein [hs-CRP] with coronary plaque characteristics as assessed by Multi slice Coronary CT angiography. Two hundreds and twenty patients presented with typical chest pain were evaluated by MSCT coronary angiography for assessment of calcium score, characters of atherosclerotic plaques as well as identification of vulnerable plaques. Lipid profile and hs-CRP were measured for those patients.

Keywords: Atherosclerosis, Lipid profile, hs-CRP, Vulnerable plaques, MSCT coronary angiography.

1. Introduction

Coronary artery disease [CAD], the leading cause of mortality worldwide, places a serious economic burden on health care systems. CAD is mainly due to atherosclerosis, which is the primary cause of mortality and morbidity in cardiovascular disease [CVD] [1]. Dyslipidemia is a well-established risk factor for the development of coronary artery disease [CAD], and this has been demonstrated in several clinical and epidemiological studies. High plasma low-density lipoprotein [LDL-C] concentrations and low high-density lipoprotein [HDL-C] concentrations have been pointed out as one of the strongest independent risk factors for coronary atherosclerotic disease[2]. Other data suggested that non-HDL cholesterol [non-HDL-C] is a better parameter for assessing CVD risk rather than TC and HDL-C [3]. Also, some studies suggest that lipid ratios, including TC/HDL-C and LDL-C/HDL-C ratios are risk factors with better predictive value for coronary atherosclerotic progression or regression than each lipid parameter used independently [4]. Concurrently, C-reactive protein [CRP] has also been proposed as a method of identifying individuals at higher cardiovascular risk. CRP is produced and secreted from the liver and smooth muscle cells surrounding atherosclerotic plaques.[5]. Studies have demonstrated that CRP correlates with various aspects of atherogenesis as well as significant coronary stenosis [6]. Major advances in CAD prevention require early detection of the vulnerable plaques. A noninvasive assay to directly detect coronary atherosclerosis would therefore be beneficial. Coronary CTA provides comprehensive information noninvasively regarding the location, severity, and characteristics of coronary atherosclerotic plaques [7].

2. Patients and methods

The study will include 220 [two hundreds and twenty] patients presented to MDCT coronary angiography for evaluation of chest pain. Based on the results of MSCT, we selected 30 patients with normal CTA for the control group. According to WHO diagnostic criteria, the standard CAD group included at least one coronary artery cavity narrowing ≥50% in the left main, left anterior descending [LAD], left circumflex [LCX], or right coronary artery [RCA] as determined by coronary imaging technique. From a group of patients with CAD, we evaluated their CT scales to select 40 patients with soft plaque, 40 patients with mixed plaque, and 40 patients with hard plaque according to the following inclusion and exclusion criteria.

2.1 Inclusion criteria

Sinus rhythm., Their heart rate less than 70 bpm spontaneously or Beta blocker induced., They can hold breath for more than 20 seconds., Weight less than 150 kg.

2.2 Exclusion criteria

Dye allergy..., Renal impairment [creatinine >1.5], Difficulties in performing CT like inadequate breath holding., Previous history of
coronary invasive maneuvers [PCI or CABG]. Respiratory failure or heart failure, Presence of arrhythmias., Treatment with statin

2.3 Methodology
All the patients will be subjected to the following:
* Personal data
* Clinical examination.
* 12-Lead ECG.
* Routine lab investigations:
  - Serum creatinine.
  - Lipid Profile
  - High sensitive C reactive Protein [hs-CRP]
  - Cardiac enzymes to exclude patients with acute coronary syndrome.

* Coronary CT angiography
The CT angiography will be performed to all patients utilizing a dual source scanner [Somatom Definition Flash, Siemens] with slice configuration of 128 x 0. 625 mm and gantry rotation time of 330 ms.

2.4 The following parameters will be assessed:
* Coronary calcium score: Based on total calcium score the coronary artery disease is graded
  - No evidence of CAD → 0 calcium score
  - Minimal → 1-10
  - Mild → 11-100
  - Moderate → 101-400
  - Severe → >400 [8].
* Coronary plaque type and composition.
Using Hounsfield unit three types of plaques will be identified:
* Non-calciﬁed lipid-rich plaques had a density of ≤ 60 HU.
* Mixed ﬁbrous plaques had a density of [61 ± 119]
* Calcified plaques had a density of ≥ 120 HU.

2.4.1 Signs of vulnerable plaque
* Positive Remodeling
  Positive remodeling [outward expansion] is defined as 5% increase in the luminal cross-section at the site of plaque compared with the normal proximal segment of the vessel.[10].
* Napkin Ring sign
  The napkin ring sign is deﬁned by inhomogeneous plaque containing a core of lower attenuation material and an outer rim with higher attenuation material. [11].

2.5 Statistical analysis
Data will be collected, processed and analyzed using Statistical Package For Social Science [SPSS] version 23 [12].

3. Results and discussion
Low density lipoprotein [LDL] from the soft plaque group, mixed plaque group and hard plaque group showed values signiﬁcantly higher than control group [p=0.001].

Total cholesterol [TC] from the soft plaque group and mixed plaque group showed values signiﬁcantly higher than the hard plaque group and control group [p=0.00001]. However total cholesterol in soft plaque showed higher value than mixed plaques but the difference is statistically insigniﬁcant [p=0.0238].

High density lipoprotein [HDL] and Total triglycerides [TG] showed no signiﬁcant difference among the four groups. [p=0.467 , p=0.991 respectively].

Table (1) Lipid proﬁle in different plaques

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<tr>
<td>Total cholesterol [mg/dl]</td>
<td>230.5±23.09</td>
<td>225.5±12.58</td>
<td>173±14.62</td>
<td>150.5±5.77</td>
<td>0.00001</td>
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<td>LDL [mg/dl]</td>
<td>124.52± 40.21</td>
<td>123.36± 42.13</td>
<td>113.30± 33.71</td>
<td>99.77±28.53</td>
<td>0.001</td>
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<td>HDL [mg/dl]</td>
<td>38.27±12.8</td>
<td>40.50±12.668</td>
<td>41.37±10.27</td>
<td>41.74±9.29</td>
<td>0.467</td>
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<tr>
<td>Total Triglycerides [mg/dl]</td>
<td>197.25±68.23</td>
<td>193.98±63.21</td>
<td>194.86±41.31</td>
<td>187.75±26.87</td>
<td>0.991</td>
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Hs-CRP from the soft plaque group, mixed plaque group and hard plaque group showed values signiﬁcantly higher than the control group [p<0.001]. Also, the mean level of hs-CRP in the soft plaque and mixed plaque
group was significantly higher than the hard plaque group \( p<0.001 \). However, the difference between the soft plaque group and the mixed plaque group was statistically insignificant \( p=0.362 \). The results and the levels of CRP were statistically analyzed. The

### Table 2: Hs-CRP in different types of plaques

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<tr>
<td>Hs-CRP [mg/L]</td>
<td>5.21±1.17</td>
<td>4.91±1.78</td>
<td>1.71±1.32</td>
<td>1.44±1.662</td>
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There were no significant differences in age, sex, smoking, family history of CAD, hypertension, diabetes among the four groups.

### Table 3: Effect of other different factors on type of plaque

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<tr>
<td>Age [years]</td>
<td>62.9±10.1</td>
<td>62.3±10.3</td>
<td>63.6±6.9</td>
<td>58.4±6.9</td>
<td>0.108</td>
</tr>
<tr>
<td>Sex [male/female, n]</td>
<td>29/11</td>
<td>27/13</td>
<td>28/12</td>
<td>22/8</td>
<td>0.946</td>
</tr>
<tr>
<td>Hypertension [n]</td>
<td>24</td>
<td>23</td>
<td>25</td>
<td>17</td>
<td>0.957</td>
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<tr>
<td>Diabetes Mellitus [n]</td>
<td>25</td>
<td>22</td>
<td>23</td>
<td>17</td>
<td>0.917</td>
</tr>
<tr>
<td>Family Hx of CAD</td>
<td>8</td>
<td>7</td>
<td>6</td>
<td>6</td>
<td>0.932</td>
</tr>
<tr>
<td>Smoking/Nonsmoking [n]</td>
<td>26/14</td>
<td>25/15</td>
<td>21/19</td>
<td>18/12</td>
<td>0.692</td>
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Some studies have found the diagnostic accuracy of MSCT to be excellent compared with catheter angiography, but there has not been an abundance of clinical outcomes studies. For any new risk marker to be considered useful for risk prediction, it must, at the very least, have an independent statistical association with risk after accounting for established readily available and inexpensive risk markers [13]. CAD is one of the most common causes of death in humans. Recently, its incidence and mortality are increasing. ACS, accounting for 30% to 40% of CAD, has been the main cause of CAD patients' poor prognosis and sudden death. Coronary atherosclerosis, plaque formation, plaque instability and disruption, a thrombus in the artery, obliteration of the artery, heart muscle ischemia and even necrosis are the basic pathological mechanisms of CAD. The basic pathology of ACS and sudden coronary death involve vulnerable plaque, vulnerable cardiac muscle, and vulnerable blood [14].

Research shows that almost all the cases of plaque disruption are based on the instability of plaque with little relation to the degree of coronary artery stenosis. A correlation has been shown between elevated systemic inflammation markers, accumulation of inflammatory cells within atherosclerotic soft plaque, and lower fibrous cap thickness [15]. Early identification and early intervention with soft vulnerable plaques is definitely important to improving the prognosis in ACS. We need a non-invasive means to replace early invasive inspection [Coronary angiography and IVUS] to identify and predict unstable plaque [16].

Selected biomarkers may be used to predict future cardiovascular events, but the gains over considering conventional risk factors are minimal. Many risk factors have been proposed as predictors of CHD. New risk factors or markers are frequently identified and evaluated as potential additions to standard risk assessment strategies [17].

Histopathology shows that vulnerable plaques are characterized by their connective tissue, a fairly high content of fat and inflammatory masses in soft plaques with the frequent occurrence of hemorrhage, calcification, necrosis and occasional thrombus on the surface. Studies have discovered that CRP is a typical phase synthetic protein in human. It is not only a symbol of inflammation but also a direct factor of arterial thrombosis. Serum Hs-CRP levels have a close correlation with inflammation and with the degree of tissue injury. CRP has been shown to participate in the whole atherosclerotic process, including damage of blood vessel endothelium and the formation, maturation, instability and final disruption of atheromatous plaque [18].

We diagnosed all the patients based on their MSCT results. The results and the levels of serum Hs-CRP were statistically analyzed. The study showed that mean levels of serum Hs-CRP in the soft plaque group and the mixed plaque group were statistically higher than...
those in the hard plaque group \([P < 0.01]\). Mean levels of serum Hs-CRP in the three plaque groups were statistically higher than in the control group \([P < 0.01]\). This agrees with previous research results by Sun et al which suggested that inflammation is involved in the occurrence and development of plaque and is the cause for ACS [19].

Concerning lipid profile , as expected, the majority of the normolipidemia group had almost no affected vessels. In comparison, all dyslipidemia groups except for the hypertriglyceridemia and low HDL patients , were associated with higher rates of coronary artery disease.

An earlier study by Paramsothy et al. showed that, of 4,795 MESA participants, without known clinical cardiovascular disease, those with combined hyperlipidemia, hypercholesterolemia, and MetS had increased relative risk for prevalent CAC compared with normolipidemia participants when adjusting for demographic and CVD risk factors.[20]

Similar to Paramsothy et al., we found that isolated hypertriglyceridemia was not associated with CAC extent [20]. These findings suggest that an isolated elevation in triglyceride levels may not be a pathologic risk factor for subclinical atherosclerosis, although hypertriglyceridemia may still be an important factor in cardiovascular disease.

In contrast to Noda et al. who studied two hundred and eighty-nine patients who underwent 64-slice multidetector CT for suspected coronary artery disease. They found that HDL-c cholesterol levels were more accurate for diagnosing the presence of high-risk coronary plaque[21]. However we found no significant association between low HDL-c and prevalent CAC which is similar to the results obtained by Paramsothy et al.

Further more, elevated LDL-c seems to be the principal determinant of CAC prevalence and its extent, with high TG levels having less influence . This is in line with Paramsothy et al.’s findings where the same dyslipidemia groups were associated with prevalent CAC [22].

4. Conclusion
The partial and systemic inflammation has a major effect on arteriosclerosis and its complications. The MSCT coronary angiography can effectually detect coronary atherosclerosis plaque and provide evidence of vulnerable patients. Hs-CRP, indirectly reflect the stabilization degree of plaque and reveal their involvement in the occurrence and development of vulnerable plaques. Also, Dyslipidemia except for hypertriglyceridemia has an important association with vulnerable atherosclerotic plaques and the development of Coronary artery disease. However the effect of HDL level and its relation with the atherosclerotic process and vulnerability of coronary plaques is still under debate and still need further studies.

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