Methods Sixty-five patients with ACS and 75 controls with SAP with similar atherosclerotic risk profiles were studied. CT angiography was performed using a dual-source CT scanner before invasive catheterisation. The lesion characteristics that were assessed included luminal cross-sectional area (L-CSA), vascular cross-sectional area (V-CSA), plaque area, and degree of stenosis with DSCT and quantitative coronary angiography (QCA), and plaque types, mean and minimal CT density (HU), remodelling index, and the presence of 'spotty' calcifications with DSCT.

Results All parameters, including L-CSA, V-CSA, plaque area, and degree of stenosis showed a good correlation between DSCT and QCA ($p<0.05$). In comparison with stable lesions in SAP, culprit lesions in ACS had a much larger mean V-CSA (20.5 ± 6.0 vs 14.8 ± 4.8 mm²), plaque area (15.3 ± 5.0 vs 11.1 ± 3.3 mm²), and remodelling index (1.3 ± 0.2 vs 1.0 ± 0.4) ($p<0.05$). The prevalence of non-calcified/calcified/mixed plaque was 30/0/35 for culprit lesions in ACS compared with 25/15/35 for stable lesions in SAP ($p<0.01$). The proportion of 'spotty' calcified plaque was 21.5% in ACS culprit lesions (14 of 65) compared with 1.3% for stable lesions in SAP (1/75). The mean HU and minimum HU of culprit lesions in ACS versus those in stable lesions of SAP were 88.6 ± 43.2 vs 154.2 ± 98.7 ($p<0.01$) and 45.9 ± 34.7 vs 98.2 ± 76.8 ($p<0.01$), respectively.

Conclusions DSCT is feasible to quantify plaque and distinguish culprit lesions in ACS, which display a greater proportion of non-calcified material and 'spotty' calcifications, lower CT attenuation, and a higher remodelling index compared with stable lesions in SAP.

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CHARACTERISATION BY DUAL-SOURCE CT OF CULPRIT LESIONS IN ACUTE CORONARY SYNDROMES COMPARED WITH STABLE ANGINA PECTORIS

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Objectives The advent of dual-source CT (DSCT) has enabled easy detection of atherosclerotic plaques and assessment of their composition and mechanical properties. This study assessed the value of DSCT in identifying the characteristics of culprit lesions in acute coronary syndrome (ACS) as compared with lesions in stable angina pectoris (SAP).