Gastric wall fat halo sign as a risk factor for cardiovascular diseases

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Abstract

A B S T R A C T

Objective: To investigate the relationship between gastric wall fat halo sign and potentially associated cardiovascular disease (CVD) in thoracic computed tomography (CT). Material and Methods: Between October 2018 and June 2019, 62 patients with gastric wall fat halo sign and 97 controls were prospectively evaluated. Patient height, weight, body mass index (BMI), sex, age, ascending aorta, descending aorta, main pulmonary artery, right and left pulmonary artery, long and short cardiac axis and maximum transverse thorax diameters; and ascending, arcus, descending aorta and coronary artery calcium scores were recorded for the two groups. Results: No significant differences were found in sex, age, height, body weight or BMI between the two groups (p > 0.125). Patients with gastric wall fat halo sign had significantly larger diameters of the ascending aorta, the descending aorta, the main pulmonary artery, the right and left pulmonary arteries, and the short and long cardiac axes and a higher cardiothoracic ratio (CTR) than the control group (p < 0.001). Additionally, the calcium scores of the ascending, arcus, and descending aortas and the coronary arteries were significantly higher detected in patients group (p < 0.001). Conclusion: The gastric wall fat halo is the result of excessive fat accumulation and can be observed in overweight people, especially those with increased visceral fat tissue. Additionally, patients with a gastric wall fat halo have a higher cardiovascular risk because of increased vascular diameters, CTR, heart sizes and calcium scores.
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Keywords: Cardiovascular disease, CT, gastric wall fat halo sign, obesity

What is ready known about this topic?

A fat halo sign refers to fat accumulation in the submucosal layer of the gastric, colonic and small bowel wall. The fat halo can also be commonly detected in overweight people, especially those with increased visceral fat tissue.

What does this article add?

Obesity is a remarkably heterogeneous condition, and visceral fat tissue has been revealed to be more deeply related to cardiovascular diseases than subcutaneous or total fat tissue. To the best of our knowledge, no study has investigated the gastric wall fat halo sign as a risk factor for cardiovascular diseases.

INTRODUCTION

Despite preventative and therapeutic efforts, cardiovascular diseases (CVDs), especially heart disease and stroke, are prominent causes of mortality and morbidity around the world. The main risk factors and pathological mechanisms resulting in CVD can be traced back to childhood. There are several controllable and uncontrollable risk factors for CVD, such as male sex, family history, old age, smoking, hypertension, physical inactivity, hypercholesterolemia, diabetes mellitus and obesity.

The unfavorable consequences of overweight and obesity have increased worldwide and can be seen beginning in childhood. Obesity is thought to be an independent risk factor for CVD, and increased body mass index (BMI) facilitates the development of comorbidities that contribute to CVD. The generally preferred anthropometric technique to evaluate relative weight and classify obesity is BMI, which is the ratio of total body weight to height squared. Additionally, obesity is a remarkably heterogeneous condition, and visceral fat tissue has been revealed to be more deeply related to CVD than subcutaneous or total fat tissue.

A fat halo sign refers to fat accumulation in the submucosal layer of the gastric, colonic and small bowel wall. The fat halo in the intestinal wall was first associated with chronic inflammatory bowel diseases, but it can also be commonly detected in overweight people, especially those with increased visceral fat tissue. Additionally, fat halo sign can be observed in approximately 21% of patients who underwent computed tomography (CT) examinations.

There are only a few studies on gastric wall fat halo sign. To the best of our knowledge, no study has investigated the gastric wall fat halo sign as a risk factor for CVD. In this study, we aimed to investigate the relationship between gastric wall fat halo sign and potentially associated CVD based on thorax CT.

MATERIAL AND METHODS

Study Population

Between October 2018 and June 2019, data from 71 patients who underwent unenhanced thorax CT and who had gastric wall fat halo sign were prospectively recorded. Three patients with chronic inflammatory bowel disease history and six patients with implanted coronary artery stents were not included in the study. Therefore, 62 patients were included in the final study.

For the control group, 97 patients with no gastric wall fat halo sign were selected from the clinical database of patients who underwent unenhanced thorax CT. Age, sex, height (m) and body weight (kg) were noted for the patient and control groups. Additionally, BMI (kg/m²) was calculated as the ratio of total body weight to height squared for both groups. This study was approved by the Institutional Ethics Committee.

CT Protocol
The thorax CT examinations were performed with the participants in the supine position, and the patients were scanned from the lung apices to the adrenal gland level during a breath hold at the end of inspiration with using 16 slices (Siemens Somatom Sensation, Forchheim, Germany). All patients were examined using the standard scanning protocol without intravenous contrast agent. The CT protocol was as follows: 120 kVp, tube current of 150–165 mAs, maximum 2.5 mm collimation, slice thickness of 1.5 mm and rotation time of 0.5 s. Then, the images were reconstructed into multiplanar reformations.

Analysis of CT Images

The thorax CT images were evaluated by two radiologists (A.K. 8 years of thoracic radiology experience and M.K. 7 years of thoracic radiology experience) to reach a joint consensus. The images were transferred to the Syngo workstation (Siemens Medical Solutions). Linear fat accumulation in the submucosal layer of the gastric wall with attenuation of <-10 Hounsfield units (HUs) was described as a gastric wall fat halo sign (Figure 1). The measurements of the outer wall of the ascending aorta, the descending aorta, the main pulmonary artery, and the right and left pulmonary arteries were recorded in the axial plane at the level of the pulmonary artery bifurcation (Figure 2). After detecting the axial slice level that represented the maximum size of the heart, the long and short cardiac axes and the maximum transverse thoracic diameter were measured at the same level (Figure 3). Then, the cardiothoracic ratio (CTR) was calculated as the ratio of the long diameter of the cardiac axis to the maximum transverse thoracic diameter. The calcification of the ascending, arcus, and descending aortas and total coronary artery calcification were quantified by the method defined by Agatston et al.\textsuperscript{13}. A region of interest (ROI) was positioned manually on the calcification of the ascending, arcus, and descending aortas and the coronary arteries. After selecting the total number of pixels over 130 HUs, the Agatston calcium score was computed with the software.

Statistical Analysis

All of the data were analyzed using the Statistical Package for the Social Sciences (SPSS 13.0 Statistical Software, SPSS Inc., Chicago, IL, USA) and the MedCalc package of Statistical Software version 16.8 (MedCalc Software bvba, Ostend, Belgium). Descriptive statistics, including the means and ranges, were calculated for age, height, BMI, CTR, the calcium scores of the ascending, arcus, and descending aortas and the coronary arteries, and the diameters of the ascending aorta, the descending aorta, the main pulmonary artery, the right and left pulmonary arteries, the short and long cardiac axes and the transverse thorax in the patient and control groups. The Kolmogorov–Smirnov test was used to identify deviations from anormal distribution. Additionally, the parametric Student’s t-test was used to compare the age, height, BMI, and the diameters of the ascending aorta, the descending aorta, the main pulmonary artery, the right and left pulmonary arteries, the short and long cardiac axes and the transverse thorax. The nonparametric Mann-Whitney U test and the parametric Student’s t-test were used to compare the calcium scores of the ascending, arcus, and descending aortas and the coronary arteries. A p value of less than 0.05 was considered to indicate a significant difference.

RESULTS

In this study, we analyzed 62 patients (45 male, 17 female) with gastric wall fat halo sign and 97 control patients (64 male, 33 female). The mean ages of the patient and control groups were 63.5 ± 8.1 and 62.1 ± 8.8 years, respectively. The mean height (m), body weight (kg) and BMI (kg/m\textsuperscript{2}) of the patient and control groups were 1.62 ± 0.1 m and 1.63 ± 0.1 m; 80.8 ± 13.4 kg and 77.4 ± 13.7 kg; and 30.6 ± 4.9 and 29.4 ± 5.6 kg/m\textsuperscript{2}, respectively. There were no statistically significant differences in sex or the mean values of age, height, body weight or BMI between the two groups (p > 0.125).

The mean diameters of the ascending aorta, the descending aorta, the main pulmonary artery, the right and left pulmonary arteries, the short and long cardiac axes and the transverse thorax, the CTR, and the calcium scores of the ascending, arcus, and descending aortas and the coronary arteries are presented in Table 1. Patients with gastric wall fat halo sign had significantly larger diameters of the ascending aorta, the descending aorta, the main pulmonary artery, and the right and left pulmonary arteries than the control group (p < 0.001). Although the short and long diameters of the cardiac axis and CTR were observed to
be significantly higher in patients with gastric wall fat halo sign (p < 0.001), no statistically significant
difference was detected in the transverse thoracic diameter between the two groups (p=0.861). Additionally,
the calcium scores of the ascending, arcus, and descending aortas and the total and coronary arteries were
significantly higher in patients with gastric wall fat halo sign than in patients without a gastric wall fat halo
sign (p < 0.001) (Figure 4).

DISCUSSION
In the present study, we investigated the relationship between gastric wall fat halo sign detected by thorax
CT and CVD. We found that gastric wall fat halo sign might be detected in patients with increased aorta
and pulmonary artery diameters and calcium scores.

Although there are several studies of the fat halo sign in the gastrointestinal tract, there are few studies
involving the fat halo sign in the gastric wall\textsuperscript{10-16}. To the best of our knowledge, this is the first study
investigating the relationship between gastric wall fat halo sign and associated CVD based on thorax CT.
The fat halo sign was first thought to indicate the presence of inflammatory bowel disease. However, fat
halo sign can be seen especially in overweight people. Additionally, gastric wall fat halo sign can be observed
in patients without gastrointestinal disease and with increased visceral fat tissue\textsuperscript{12,13}. After fat is stored in
normal adipose tissues, free fatty acids are discharged into the blood\textsuperscript{17}. The fat halo sign could be related
to an increase in the blood concentration of free fatty acids and could be an indicator of excessive fat
accumulation\textsuperscript{17}.

The ascending and descending aorta diameters are easily evaluated through chest CT. In our study, we
found that patients with gastric wall fat halo sign had significantly higher ascending and descending aorta
diameters than those without fat halo sign. One of the causes of aortic diameter increase is visceral obesity.
The increase in aortic diameter associated with visceral obesity could be related to the process of adaptation
to hypertension, increased blood volume and underlying structural or functional aortic abnormalities\textsuperscript{18}.
Additionally, visceral obesity can stimulate vascular aging, resulting in future cardiovascular events\textsuperscript{8}. Visceral
obesity and gastric wall fat halo sign are associated with each other, so increased aortic diameters may be
related to this relationship.

Although pulmonary hypertension may appear to be a sporadic disease with undetectable risk factors, it can
occur as a result of preexisting medical conditions\textsuperscript{19}. Pulmonary hypertension and obesity are generally two
coexisting conditions in the clinic, and increased BMI induces the development of pulmonary hypertension\textsuperscript{20}.
Chan et al. reported that there were more obese participants in the pulmonary hypertension group than in the
control group, whereas most patients were not obese in both groups\textsuperscript{21}. In this study, we revealed that patients
with gastric wall fat halo sign had significantly higher main, right and left pulmonary artery diameters than
those without fat halo sign. This relationship may be the reason for pulmonary artery enlargement in our
study.

Obesity is connected with most CVDs, such as coronary heart disease, hypertension and heart failure\textsuperscript{22}. In
particular, visceral adipose tissue is hormonally active tissue that contributes to vascular inflammation by
producing adipokines and is a risk factor for mortality and CVD\textsuperscript{23}. Additionally, obesity is related to some
hemodynamic changes, such as increased cardiac output, diastolic filling pressure, left ventricle hypertrophy
and ventricular dilatation\textsuperscript{24}. The CTR is usually used to evaluate cardiomegaly in chest radiography,
cardiomegaly indicating underlying CVD, is evidenced by hypertrophy or dilation of the heart\textsuperscript{25}. CT imaging
is a simple and effective method for assessing cardiomegaly. Despite a lack of significant difference in the
transverse thoracic diameter, we found that patients with gastric wall fat halo sign had significantly higher
short and long diameters of the cardiac axis and CTRs than the control group. These results may be related to
cardiovascular changes caused by visceral adipose tissue.

Atherosclerosis is a major cause of death and morbidity in industrialized societies due to its association with
cardiovascular events and peripheral vascular disease. Atherosclerosis has different stages, and calcification
indicates advanced atherosclerotic lesions\textsuperscript{26}. Atherosclerosis starts with fatty streaks and advances further to
calcified plaques in adulthood. Although the cause of vascular calcification is not fully understood, inflammation, dysregulated metabolism, and osteogenesis are some processes associated with vascular calcification. Thoracic aortic calcification represents a systemic atherosclerotic burden and results in an increased risk of cardiovascular events. Visceral fat tissue is an endocrine organ that can produce atherogenic agents. Additionally, this condition can induce vascular aging resulting in an increased risk of CVDs, hypertension, and calcified atherosclerotic plaques. It has been reported that the risk of aortic atherosclerosis increases as the amount of visceral fat tissue increases. Moreover, aortic wall calcification can be assessed with CT and lateral X-ray. In our study, we used non contrast thoracic CT to evaluate thoracic aortic calcification and found that patients with gastric wall fat halo sign had significantly higher calcium scores in the ascending, arcus, and descending aortas than patients without fat halo sign.

Coronary artery disease, caused by atherosclerosis, is a main cause of morbidity and mortality worldwide, so the early diagnosis of subclinical atherosclerosis is crucial for preventing coronary artery disease. Similar to thoracic aortic calcification, obesity is an independent risk factor for atherosclerosis and coronary artery calcification. The calcium score of coronary arteries indicates the burden of atherosclerotic plaque in arteries and is a helpful, noninvasive method for the assessment of subclinical atherosclerosis in asymptomatic patients. Additionally, the calcium score of coronary arteries is connected with cardiovascular events and obesity. In this study, we showed that higher calcium scores of coronary arteries were detected in patients with gastric wall fat halo sign related to visceral obesity.

This study has a number of limitations. First, the CT images were evaluated based on a consensus, and we did not evaluate the inter- or intraobserver variability in our study. The second limitation was that the number of patients was also relatively small. The third limitation was that participants in the patient and control groups did not undergo abdominal CT imaging, so visceral fat tissue measurement could not be performed in our study. Fourth, we did not have access to laboratory parameters such as lipid panels.

CONCLUSION

The gastric wall fat halo results from excessive fat accumulation and can be observed in overweight people, especially those with increased visceral fat tissue. Additionally, patients with a gastric wall fat halo have higher cardiovascular risk because of increased vascular diameters, CTR, heart sizes and calcium scores. However, extensive studies with larger populations are needed to clearly confirm the relationships of gastric wall fat halo sign with cardiovascular diseases.

References

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Figure Legends

Fig. 1. 45 year-old man with pneumonia. Unenhanced thorax CT image represents thin linear fatty infiltration in submucosal layer of gastric wall with attenuation value<-10 HU.

Fig. 2. 43 year-old man with cough complaint. The measurements of the ascending aorta, the descending aorta, the main pulmonary artery, and the right and left pulmonary arteries were recorded in the axial plane at the level of the pulmonary artery bifurcation.

Fig. 3. 51 year-old man with solitary pulmonary nodule. The measurements of the long and short cardiac axes and the maximum transverse thoracic diameter were obtained at the axial slice level that represented the maximum size of the heart.

Fig. 4. 59-year-old man with gastric wall fat halo sign.

A, Unenhanced sagittal CT image shows extensive calcific atherosclerotic plaques in aorta

B, Unenhanced axial CT image shows extensive calcific atherosclerotic plaques in coronary arteries

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