Value of echocardiography for diagnosis and prognosis of chronic Chagas disease cardiomyopathy without heart failure

R J Viotti, C Vigliano, S Laucella, B Lococo, M Petti, G Bertocchi, B Ruiz Vera, H Armenti

Accepted 20 October 2003

Objective: To establish the usefulness of echocardiography for the clinical classification of patients with chronic Chagas disease and to determine the predictors of mortality and clinical events.

Methods: 849 patients with chronic Chagas disease were enrolled at Hospital Eva Perón, San Martín, Buenos Aires, Argentina. The presence of Trypanosoma cruzi antibodies was tested by an enzyme linked immunosorbent assay (ELISA), indirect haemagglutination, and indirect immunofluorescence assays. A patient was considered to be infected when at least two of the tests were positive. Patients with chronic Chagas disease were divided into three groups: group 0: reactive serology and abnormal ECG; group 1: reactive serology and abnormal ECG and chest radiography without cardiac enlargement; group 2: reactive serology and abnormal ECG and chest radiography with cardiac enlargement. Clinical events were defined as new ECG abnormalities, change in clinical status resulting in transfer to another group, and death. Morphologically characterised segmental lesions were seen in 12 patients with a clinical event, p < 0.0001, left ventricular diastolic dimension (mean (SD) 4.88 (0.54) cm vs 5.44 (0.83) cm, p < 0.0001), left ventricular systolic dimension (mean (SD) 2.98 (0.62) cm vs 3.64 (1.03) cm, p < 0.0001), and ejection fraction (mean (SD) 0.68 (0.10)% vs 0.60 (0.16)%), p < 0.0001) were predictors of clinical events. Segmental lesions were observed in 211 of 849 patients (25%). Segmental lesions were seen in 66 (13%) and systolic dysfunction was seen in four of 505 (0.8%) patients with normal ECG. Significant differences were found between the groups of patients (group 0: reactive serology and normal ECG and chest radiography without cardiac enlargement and no signs of heart failure; group 1: reactive serology and abnormal ECG and chest radiography without cardiac enlargement; group 2: reactive serology and abnormal ECG and chest radiography with cardiac enlargement and no signs of heart failure).

Conclusion: Echocardiography was useful to characterise and to determine the prognosis of patients with chronic Chagas disease without heart failure.

Echocardiography is a useful complementary method for the follow up of patients with chronic Chagas disease cardiomyopathy; however, echocardiograms are not part of the clinical classification of patients with chronic Chagas disease. Moreover, few studies involving large numbers of patients have been carried out to establish the prognostic value of the echocardiographic variables for the evolution of cardiomyopathy or death. Most of the studies examined patients with heart failure or advanced heart disease. Similarly, few reports analysed the type, location, and differences between segmental lesions. Our objectives were both to establish the diagnostic and prognostic value of echocardiography in chronic Chagas disease without heart failure and to characterise the location and morphology of segmental lesions.

METHODS

Patients

Eight hundred and forty-nine patients with chronic Chagas disease were enrolled at Hospital Eva Perón, San Martín, Buenos Aires, Argentina. The presence of T. cruzi antibodies was tested by an enzyme linked immunosorbent assay (ELISA), indirect haemagglutination, and indirect immunofluorescence assays. A patient was considered to be infected when at least two of the tests were positive. Patients with...
congenital, hypertensive, coronary, alcoholic, or other associated cardiomyopathies were excluded from this study. ECG, stress testing, chest radiography, and two dimensional echocardiography were performed on admission in all patients with confirmed positive serology for T. cruzi.

Patients were classified into clinical groups according to Kuschnir: group 0, reactive serology and normal ECG and chest radiography without cardiac enlargement; group 1, reactive serology and abnormal ECG and chest radiography without cardiac enlargement; and group 2, reactive serology and abnormal ECG and chest radiography with cardiac enlargement and neither radiological nor clinical signs of heart failure. Group 3 patients, presenting with heart failure, were excluded because they were considered to be at the end stage of the disease and therefore to have a high expected mortality.

During follow up, complete right bundle branch block, left anterior fascicular block, complete left bundle branch block, ventricular premature complexes of Lown degree II or greater, electrical inactivation areas, atrial flutter, atrial fibrillation, atrial tachycardia, sinus bradycardia < 50 beats/min, sustained and non-sustained ventricular tachycardia, second degree type II atrioventricular block, complete atrioventricular block, and placement of pacemakers were considered either previous ECG abnormalities related to the disease or new ECG abnormalities related to the disease. Once divided into clinical groups, patients were periodically evaluated as follows: group 0, ECG every six months to one year; group 1, ECG every four months; and group 2, ECG every three months. Holter monitoring or repeated stress testing and echocardiogram were optional for patients belonging to group 2.

![Figure 1](A) Two chamber apical view from a 67 year old female patient showing hollow punch lesion type I (arrow) without left ventricular (LV) dilatation or systolic dysfunction in the mid inferior wall during diastole. LA, left atrium. (B) The same patient during systole. (C) Four chamber apical view with intravenous contrast agent showing apical hernia with thrombus (Th), LV dilatation, and normal systolic function in a 44 year old female patient. The wall of the hernia has no muscular layer (arrows). RA, right atrium; RV, right ventricle. (D) Four chamber apical view with extensive thinning of apical segments (arrows); LV enlargement; and systolic dysfunction in a 60 year old male patient. (E) Four chamber apical view with all three kinds of segmental lesions, LV enlargement; and systolic dysfunction in a 71 year old female patient. Small arrow, type I lesion in the interventricular septum; double arrow, type II lesion on apical segments; large arrow, type III lesion on the lateral wall.
Table 1  Univariate analysis for mortality and clinical events in patients with chronic Chagas disease

| Variable                      | Survival | Death | No clinical event | Clinical event | Total |
|-------------------------------|----------|-------|-------------------|----------------|-------|
| Age at admission (years)      | 43.39 (11.11) | 49.12 (13.11)*** | 42.65 (11.21) | 47.50 (10.13)†† | 849 |
| EARD (ECG abnormalities related to the disease) | 304/833 (36%) | 14/16 (87%)*** | 220/699 (31%) | 98/150 (65%)†† | 849 |
| NEARD (new ECG abnormalities related to the disease) | 112/833 (13%) | 9/16 (56%)*** | – | – | 849 |
| Change in clinical group      | 68/833 (8%) | 15/16 (94%)*** | – | – | 849 |
| RVDD (cm)                     | 1.85 (0.55) | 2.12 (0.53) | 1.83 (0.52) | 1.97 (0.65)† | 753 |
| LVDD (cm)                     | 4.96 (0.62) | 5.99 (0.68)*** | 4.88 (0.54) | 5.44 (0.83)† | 849 |
| LVSD (cm)                     | 3.06 (0.72) | 4.71 (0.90)*** | 2.98 (0.62) | 3.64 (1.03)† | 840 |
| LASD (cm)                     | 3.44 (0.57) | 3.82 (0.82) | 3.39 (0.54) | 3.74 (0.69)† | 817 |
| Radius: thickness ratio       | 2.88 (0.70) | 3.97 (1.50) | 2.83 (0.65) | 3.24 (1.02)† | 823 |
| PWT (cm)                      | 0.98 (0.22) | 0.81 (0.18) | 0.89 (0.21) | 0.89 (0.27) | 825 |
| SWT cm                        | 0.98 (0.24) | 0.95 (0.20) | 0.98 (0.23) | 0.99 (0.27) | 820 |
| Systolic dysfunction          | 60/832 (7%) | 8/16 (50%)*** | 30/698 (4%) | 38/150 (25%)† | 848 |
| Segmental lesions             | 197/833 (24%) | 14/16 (87%)*** | 141/699 (20%) | 70/150(47%)†† | 849 |
| EF (%)                        | 0.67 (0.11) | 0.42 (0.17)*** | 0.68 (0.10) | 0.60 (0.16)†† | 840 |

Data are mean (SD) or proportion. Evaluation by χ² test: clinical events, change in clinical group, EARD (ECG abnormalities related to the disease), NEARD (new ECG abnormalities related to the disease), systolic dysfunction, and segmental lesions. Evaluated by unpaired Student’s t test: LASD (left atrial systolic dimension), LVDD (left ventricular diastolic dimension), LVSD (left ventricular systolic dimension), PWT (diastolic posterior wall thickness), RVDD (right ventricular diastolic dimension), and SWT (diastolic septal wall thickness).

*p<0.01, **p<0.05, ***p<0.0001 compared with alive group; †p<0.05; ††p<0.001 compared with patients without clinical events.

A Toshiba SSH 140 A ultrasound (Toshiba, Tokyo, Japan) and 2.5–3.7 MHz transducers were used to run the echocardiograms.

**Prognostic variables**

Age on admission, years of follow up, presence of either ECG abnormalities related to the disease or new ECG abnormalities related to the disease, change in clinical status resulting in transfer to another group, right ventricular diastolic dimension, left ventricular diastolic and systolic dimensions, left atrial systolic dimension, diastolic posterior wall thickness, diastolic septal wall thickness, radius to thickness ratio, left ventricular systolic dysfunction (as reported by the physician), ejection fraction (Teichholz), and the presence of segmental lesions were analysed to study mortality and clinical events. These variables were also used to compare the clinical groups on admission. New ECG abnormalities related to the disease, change in group, and death were defined as clinical events, taking into account only one per patient. New onset heart failure was considered to be a clinical event that was included in the variable change in clinical group. Only cardiac death was taken into account, without distinguishing the mechanisms of cardiac arrest (heart failure, sudden death, and thromboembolic events).

**Type and location of segmental lesions**

The types of regional contraction abnormality, as established by the American Society of Echocardiography, were defined as hypokinesis, akinesis, dyskinesis, and aneurysmal. Segmental lesions found in the apical, posterior, inferior, anterior, lateral, and septal cardiac walls were recorded. To improve the identification of morphological differences between segmental lesions, 12 patients were selected for further analysis by two dimensional echocardiography (ATL 3000 with 2.5 MHz transducer) with a second harmonic and contrast agent (4 g of intravenous contrast agent (Leovist NR) administered over a 10 minute period).

**Statistical analysis**

A univariate analysis was performed to determine differences in mortality and clinical events. Student’s t test was applied for continuous variables and χ² test for categorical variables. Only variables shown to be statistically different in the univariate analysis were used in the multivariate analysis with logistic regression. For the analysis of logistic regression of mortality, ejection fraction was considered an offset variable because of its well known prognostic value for mortality and its colinearity with left ventricular systolic dimension. To compare continuous variables (described above under “Prognostic variables”) between clinical groups,

**Table 2  Multivariate analysis for mortality and clinical events in patients with chronic Chagas disease**

| Variable                      | Mortality β Coefficient | p Value | Clinical events β Coefficient | p Value |
|-------------------------------|-------------------------|---------|-------------------------------|---------|
| Age at admission              | -0.00895                | 0.81    | 0.01                          | 0.18    |
| EARD                          | 1.4                     | 0.11    | 0.73                          | 0.003*  |
| NEARD                         | 0.51                    | 0.55    | –                             | –       |
| Change in clinical group      | 2.31                    | 0.0068* | –                             | –       |
| LVDD                          | -1.46                   | 0.08    | 3.05                          | 0.003*  |
| LVSD                          | 2.33                    | 0.01*   | -3.61                         | 0.02*   |
| RVDD                          | –                       | –       | 0.06                          | 0.78    |
| LASD                          | –                       | –       | 0.32                          | 0.12    |
| Systolic dysfunction          | -1.02                   | 0.29    | 0.46                          | 0.28    |
| Segmental lesions             | 1.02                    | 0.29    | 0.15                          | 0.39    |
| Radius: thickness             | 0.47                    | 0.22    | 0.33                          | 0.07    |
| EF†                           | -7.38††                 | 0.003†† | -14.78                        | 0.01*   |

*Significantly different; †offset variable in mortality; ‡EF regarded as an independent variable.

www.heartjnl.com
a one way analysis of variance followed by a Scheffe test was carried out, p < 0.05 being considered significant. Data were analysed by Statistix 7.0 analytical software.

RESULTS

Prognostic variables in chronic Chagas disease

We evaluated 849 patients with chronic Chagas disease (333 men and 516 women) with a mean (SD) age of 43.5 (11.1) years (range 18–74 years) and with a mean (SD) follow up of 9.9 (5.3) years (range 2–23 years). Table 1 shows the univariate analysis for mortality and clinical events. All variables except posterior wall and interventricular septum thickness (considered for the comparison between the alive and the dead groups, as well as between patients with and those without clinical events) were significantly different. Change in clinical group, left ventricular systolic dimension, and ejection fraction were the only independent predictors of mortality. ECG abnormalities related to the disease, left ventricular diastolic dimension, left ventricular systolic dimension, and ejection fraction were predictors of clinical events (table 2).

Type and location of segmental lesions

As table 3 shows, segmental lesions were observed mainly in the apical and posterior walls. The most common lesions were hypokinetic lesions and aneurysms. Only aneurysms

| Table 3 | Type and location of segmental lesions |
|---------|--------------------------------------|
| Location                           | Hypokinesis (105 patients) | Akinesis (20 patients) | Dyskinesis (14 patients) | Aneurysms (73 patients) |
| Apical wall                          | 71/126 (56%)                | 12/32 (37%)             | 12/20 (60%)              | 52/105 (49%)           |
| Posterior wall                       | 23/126 (18%)                | 10/32 (31%)             | 2/20 (10%)               | 26/105 (25%)           |
| Inferior wall                        | 8/126 (6%)                  | 8/32 (25%)              | 2/20 (10%)               | 13/105 (12%)           |
| Anterior wall                        | 1/126 (0.8%)                | 0/32 (0%)               | 0/20 (0%)                | 2/105 (2%)             |
| Lateral wall                         | 4/126 (3%)                  | 0/32 (0%)               | 0/20 (0%)                | 4/105 (4%)             |
| Interventricular septum              | 19/126 (15%)                | 2/32 (6%)               | 4/20 (20%)               | 8/105 (8%)             |
| Total segmental lesions              | 126                         | 32                     | 20                       | 105                    |

Table 4 | Relation between the echocardiogram and the clinical status of patients with chronic Chagas disease grouped by Kuschnir classification

| Variable                          | Group 0 (505 patients) | Group 1 (257 patients) | Group 2 (87 patients) | Statistic |
|-----------------------------------|------------------------|------------------------|------------------------|-----------|
| Age at admission (years)          | 40.53 (10.9)*          | 47.18 (10.26)          | 49.93 (9.36)           | F = 51.75 |
| Follow up (years)                 | 9.64 (5.13)            | 10.67 (5.53)†          | 9.37 (5.49)            | p = 0.0001 |
| NEARD                             | 33/505 (6%)            | 55/257 (21%)           | 33/87 (38%)            | χ² = 75.27 |
| Change in clinical group          | 30/505 (6%)            | 25/257 (10%)           | 28/87 (32%)            | χ² = 57.95 |
| Mortality                         | 0/505 (0%)             | 4/257 (1%)             | 12/87 (14%)            | χ² = 76.57 |
| Clinical events‡                  | 39/505 (8%)            | 66/257 (26%)           | 45/87 (52%)            | χ² = 115.05 |
| RVDD (cm)                         | 1.80 (0.48)            | 1.86 (0.56)            | 2.16 (0.75)†           | p = 0.0001 |
| LVDD* (cm)                        | 4.81 (0.50)            | 4.98 (0.56)            | 5.94 (0.74)            | p = 0.0001 |
| LVSD* (cm)                        | 2.91 (0.54)            | 3.05 (0.66)            | 4.30 (0.97)            | p = 0.0001 |
| LASD (cm)                         | 3.35 (0.49)            | 3.45 (0.49)            | 3.99 (0.93)†           | p = 0.0001 |
| PWT (cm)                          | 0.89 (0.25)            | 0.90 (0.16)            | 0.85 (0.18)            | p = 0.16 |
| SWT (cm)                          | 0.96 (0.22)            | 1.00 (0.26)            | 0.98 (0.22)            | p = 0.11 |
| Radius: thickness ratio           | 2.83 (0.65)            | 2.80 (0.61)            | 3.69 (1.11)†           | p = 0.0001 |
| E point septal separation* (cm)   | 0.49 (0.33)            | 0.70 (0.43)            | 1.35 (0.64)            | p = 0.0001 |
| Fractional shortening (%)         | 39.36 (8.47)           | 37.57 (11.30)          | 26.65 (12.04)‡          | p = 106.77 |
| Systolic dysfunction              | 4/505 (0.8%)           | 22/257 (8%)            | 42/87 (48%)            | p = 0.0001 |
| Segmental lesions                 | 66/505 (13%)           | 84/257 (33%)           | 61/87 (70%)            | p = 0.0001 |
| EF (%)                            | 0.69 (0.09)            | 0.68 (0.11)            | 0.52 (0.16)†           | p = 0.0001 |

*Significantly different from groups 1 and 2; †significantly different from groups 0 and 2; ‡only one clinical event was considered per patient; ††significantly different from groups 0 and 1, *significant difference between groups 0, 1, and 2.
were found in the anterior wall; instead, most cases of akinesia were recorded in both the inferior and posterior walls. One hundred and forty seven of 283 segmental lesions were found in the apical wall (52%), 61 in the posterior wall (21%), 33 in the interventricular septum wall (12%), 31 in the inferior wall (11%), 8 in the lateral wall (3%), and 3 in the anterior wall (1%). Twenty eight of 211 (13%) patients with segmental lesions had lesions involving more than one wall (25 patients with two combined lesions and three with three combined lesions).

Morphological characterisation of segmental lesions

Three types of segmental lesions were differentiated by the morphological and extension of the segmental damage: type I, thinning of the wall or “hollow punch” (fig 1A, B); type II, apical hernia projecting from the external cardiac edge with normal systolic function (fig 1C); and type III, extensive wall thinning, without hernia, similar to myocardial infarction expansion (fig 1D). In one patient all three kinds of segmental lesions were observed (fig 1E).

Relation between the echocardiogram and the clinical status of patients with chronic Chagas disease

Table 4 shows the relation between the echocardiographic findings and patients’ clinical status according to the Kuschmir classification on admission. Among group 0 patients, 13% had segmental lesions and 0.8% had systolic dysfunction. Significant differences were found in the left ventricular diastolic and systolic dimensions and E point septal separation with reference to groups 0, 1, and 2, even though in groups 0 and 1 the measurements were within the normal range. All the echocardiographic parameters except diastolic posterior wall thickness and diastolic septal wall thickness were significantly increased in group 2 in comparison with groups 0 and 1.

DISCUSSION

The most significant prognostic index of mortality in our study was a change in clinical group. Both left ventricular systolic dimension and ejection fraction were independent predictive values, suggesting that systolic dysfunction may be an important factor in cardiac death. However, systolic dysfunction as reported by the physician did not have a prognostic value probably because this is a qualitative measure. Conversely, left ventricular diastolic dimension, left atrial systolic dimension, radius to thickness ratio, and presence of segmental lesions were not predictors of mortality, as suggested by other authors.\textsuperscript{17} \textsuperscript{24} \textsuperscript{25} This suggestion was based either on the well known association between left ventricular enlargement, increased diastolic stress, cardiac damage, and death or on the relation between segmental lesions, complex ventricular arrhythmias, and sudden death. ECG abnormalities related to the disease, left ventricular systolic and diastolic dimensions, and ejection fraction were predictors of clinical events. However, left ventricular diastolic dimension and ECG abnormalities related to the disease were not predictors of mortality. This difference between index of mortality and clinical events is unclear. On the one hand, change in clinical group may indicate the unavoidable evolution towards the end phase of the disease and death. This evolution probably is the clinical expression of a chronically active diffuse myocarditis.\textsuperscript{26} \textsuperscript{27} On the other hand, in the less advanced phases of cardiomyopathy, this active myocarditis would be clinically manifested as ECG abnormalities related to the disease or increased left ventricular diastolic dimension. However, as the cardiac damage worsens, the systolic function may be impaired, thus being the main cause of death.

Regarding the segmental lesions, no important differences were found with previous reports showing that segmental lesions are found mainly in the apical and posterior walls,\textsuperscript{18} with hypokinesia and aneurysms being most common.\textsuperscript{29} \textsuperscript{30} It is important to point out that segmental lesions were found in asymptomatic patients with a normal ECG (group 0), suggesting that any person with positive serology for \textit{T cruzi} should be to considered to have chronic Chagas disease, bearing in mind that pathological findings depend on both the accuracy and sensitivity of the clinical tests used.

In this study, three types of segmental lesions were morphologically defined. To our knowledge, this is the first time that type 1 segmental lesions have been characterised. Echocardiography proved to be a useful complementary test for the clinical classification of chronic Chagas disease and for characterising the type of segmental lesions. Moreover, the echocardiogram may become a good predictor of the outcome of the disease.

Limitations of the study

Although the aim of our study was evaluating the prognostic value of the most commonly used echocardiographic variables, right ventricular systolic function and diastolic dysfunction would also have been important to analyse. Unfortunately, it was not possible to estimate these variables because of limitations in economic and technological resources.

The ejection fraction was calculated by the Teichholz method. The usefulness of this method is limited in the presence of segmental lesions.

ACKNOWLEDGEMENTS

We thank María Gabriela Alvarez for her permanent collaboration, Horacio Tamagusuku for technical assistance, and José Luis Leone and Fernando Corbella for critical reading of the manuscript.

Authors’ affiliations

R J Viotti*, C Vigliano, L Laucella, B Lococo, M Petti, G Bertocchi, B Ruiz Vera, H Armenti, Cardiology Department, Hospital Eva Perón, San Martin, Buenos Aires, Argentina

*Also Clinica Bessone, San Miguel, Buenos Aires, Argentina

REFERENCES

1. Feldman AM, Mac Namara D. Myocarditis. N Engl J Med 2000;343:1388–98.
2. Anon. Tropical disease research progress 1975–1994 highlights 1993–1994. Twelfth programme report of the UNDP/World Bank/WHO Special Programme for Research and Training in Tropical Diseases (IDR) Geneva: World Health Organization, 1995:125–34.
3. Elsner M. La miocardiopatia chagásica: perspectiva histórica. Medicina 1999;59(suppl II):25–40.
4. Yoshida N, Schenkman S. American trypanosomiasis. Curr Opin Infect Dis 1997;10:351–6.
5. Jones EM, Colley DG, Tostes S, et al. Amplification of a \textit{T}rypanosoma cruzi DNA sequence of inflammatory lesions in human chagasic cardiomyopathy. Am J Trop Med Hyg 1993;48:348–57.
6. Brandariz S, Schijman A, Vigliano C, et al. Detection of parasite DNA in \textit{T} chagas’ heart disease. Lancet 1995;346:1370.
7. Cossio G, Delgado AM, Zague A, et al. Trypanosoma cruzi-induced host immune system dysfunction: a rationale for parasite immunosuppressive factor(s) encoding gene targeting. J Biomed Biotechnol 2001;1:11–7.
8. Tarleton RL, Zhang L. Chagas disease etiology: autoimmune or parasite persistence? Parasitol Today 1999;15:94–9.
9. Andrade ZA. Immunopathology of Chagas disease. Mem Inst Oswaldo Cruz 1999;94(suppl II):71–80.
10. Laranje PA, Dias E, Nóbrega G, et al. Chagas disease: a clinical, epidemiologic and pathologic study. Circulation 1956;14:1035–60.
11. Puigbó J, Giardino H, Sécurez C, et al. Aspectos clínicos en la enfermedad de \textit{Chagas}. In: Madoery R, Madoery C, Cámara M, eds. Actualizaciones en la enfermedad de \textit{Chagas}. Simposio satelitc. Organismo oficial del Congreso Nacional de Medicina. 1993:27–38.
12. Manczulo EC. Chuit R. Risk of death due to chronic chagasic cardiopathy. Mem Inst Oswaldo Cruz 1999;94(suppl I):317–20.
13. Storino R. Chagas crónico. In: Storino R, Mili J, eds. Enfermedad de Chagas. Buenos Aires: Mosby, 1994:249–54.
Aneurysm of the left main coronary artery in Takayasu arteritis

A 30 year old woman was admitted to our hospital for evaluation of syncope. Her blood pressure was 100/70 mm Hg in the left arm, while it was not recordable in the right arm. Systolic bruit was present over the right supraclavicular region. There was a blowing diastolic murmur over the left sternal border. ECG was normal. Laboratory studies gave either negative or normal results.

She underwent angiographic evaluation with provisional diagnosis of Takayasu arteritis (TA). Aortogram revealed mild aortic regurgitation, total occlusion of the right subclavian artery, narrowing of the proximal left subclavian artery (below, left panel), and stenosis of the supra renal aorta (middle panel) consistent with a diagnosis of TA. Coronary angiogram showed a large aneurysm of the left main coronary artery (right panel).

Most of the discrete coronary artery aneurysms are atherosclerotic in origin; other causes of coronary aneurysms include Kawasaki disease, Takayasu disease, polyarteritis nodosa, and systemic lupus erythematosus. TA is a chronic inflammatory vasculitis characterised by stenosis or obliteration of large and medium sized arteries. Morbidity results from arterial stenosis and organ ischaemia as well as from aneurysm formation. The cardiac sequelae of TA are far more common due to aortic regurgitation and inadequately treated hypertension than arteritis affecting the coronary vessels. Coronary arteries are affected in approximately 10% of the cases. Most of the lesions cause luminal narrowing; however, coronary aneurysm formation is extremely rare. The left main coronary artery is the least frequently involved artery. Aneurysm formation results from extensive destruction of elastic fibres in the media of the involved arterial wall as well as from arterial hypertension. A coronary aneurysm often causes stasis of blood flow and results in mural thrombus and myocardial infarction; saccular aneurysm may get thrombosed, ruptured or enlarged, resulting in myocardial infarction, cardiac tamponade or sudden death.

C G Sajeev
M N Krishnan
K Venugopal
cgsajeev@hotmail.com