Extent and severity of atherosclerotic involvement of the aortic valve and root in familial hypercholesterolaemia

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Abstract

Objective—To compare the frequency of valvar and supra-valvar aortic stenosis in homozygous and heterozygous familial hypercholesterolaemia (FH).

Design—Analysis of life time cholesterol exposure and prevalence of aortic atherosclerosis in 84 consecutive cases attending a lipid clinic.

Setting—A tertiary referral centre in London.

Patients—Outpatients with FH (six homozygous, 78 heterozygous).

Interventions—Maintenance of lipid lowering treatment.

Main outcome measures—Calculated cholesterol $\times$ years score (CYS) and echocardiographic measurement of aortic root diameter, aortic valve thickness, and transaortic gradient.

Results—Four homozygotes with a mean (SD) CYS of 387 (124) mmol/l $\times$ years had severe aortic stenosis (treatment started after seven years of age), whereas the other two had echocardiographic evidence of supra-valvar thickening but no aortic valve stenosis (treatment started before three years of age). On multivariate analysis, mean transaortic gradient correlated significantly with CYS (mean = 523 (175) mmol/l $\times$ years) in heterozygotes ($p = 0.0001$), but only two had severe aortic valve and root involvement.

Conclusions—In patients with familial hypercholesterolaemia, aortic stenosis is common in homozygotes, and aortic root involvement is always present despite the lower CYS than in heterozygotes. It appears to be determined by short term exposure to high cholesterol concentrations in early life. Conversely, aortic root and valve involvement are rare in heterozygotes and occur only with severe, prolonged hypercholesterolaemia, possibly accelerating age related degenerative effects.

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Familial hypercholesterolaemia (FH) is characterised by dominantly inherited hypercholesterolaemia, early appearance of cutaneous and tendon xanthomata, and a predisposition to premature cardiovascular disease. Inheritance of one mutant gene causes heterozygous FH

(1:500), while inheritance of two mutant genes gives rise to phenotypic homozygosity (1:1 000 000).

Premature coronary artery disease is common in FH, while atheromatous involvement of the aortic valve and root is almost always present in homozygotes.¹ Lipid infiltration with consequent thickening of the aortic cusps is considered to be a unique feature of homozygotes and can affect the valve's mobility.² However, atherosclerotic involvement of the aortic root is an even more common complication, rarely reported in heterozygotes, and leads to ostial stenosis and "supra-valvar" aortic stenosis. Previous reports of aortic valve and root involvement in homozygous FH were based on cardiac catheterisation and cross sectional echocardiography.³⁻⁵ Aortography shows characteristic aortic root funnelling when the proximal ascending aorta is infiltrated by atheroma. The gradient across the aortic valve indicates the severity of its involvement, but the invasive nature of catheterisation restricts the use of this diagnostic procedure to symptomatic patients or those with clinical evidence of severe aortic stenosis. However, the increasing use of Doppler echocardiography over the past decade has made the early detection and precise assessment of aortic stenosis much easier,^{6,7} and provides an excellent tool for the screening and follow up of these patients.

To test the hypothesis that lifelong exposure to high cholesterol concentrations largely determines the magnitude of aortic valve and root disease, we studied the frequency and haemodynamic significance of valvar and supra-valvar aortic stenosis in FH homozygotes and heterozygotes, using cross sectional and Doppler echocardiography.

Methods

PATIENTS

Eighty four consecutive patients with FH were studied over a one year period during 1995 to 1996. All were outpatients at the lipid clinic at Hammersmith Hospital, which most had been attending regularly for several years. All patients had normal thyroid, renal, and hepatic function and none had diabetes mellitus.

HOMOZYGOTE GROUP

This consisted of six patients (two male, four female) with a mean (SD) age of 22.8 (7.7) years (range 12 to 32 years). Pretreatment mean serum cholesterol concentration was 21.1 (4.8) mmol/l and the geometric mean

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concentration of lipoprotein(a) (Lp(a)) was 24.4 (2.6) mg/dl. All patients except case 6 were receiving statins, while cases 3 and 4 were also on biweekly low density lipoprotein (LDL) apheresis. Case 6 was on a combination of an anion exchange resin, fibrates, and nicotinic acid. The mean serum cholesterol concentration at the time of echocardiography was 10.5 (2.0) mmol/l.

HETEROZYGOTE GROUP

This group consisted of 78 patients (49 male, 29 female) with a mean age of 52.2 (13.8) years (age range 22 to 84 years). The diagnosis was based on primary hypercholesterolaemia (> 7.5 mmol/l) with an autosomal dominant pattern of inheritance, presence of tendon xanthomata in the patient or a first degree relative, and a family history of premature coronary artery disease. Fourteen patients had proven mutations at the LDL receptor gene locus, determined as described previously.⁸ At the time of examination most patients were receiving either an HMG-CoA reductase inhibitor (statin) (43%), an anion exchange resin (8%), a fibrate (5%), or combinations of these classes of drug. Five heterozygotes were also on nicotinic acid.

ECHOCARDIOGRAPHY

In all patients cross sectional Doppler (colour, pulsed, continuous) echocardiography was performed with a Toshiba SSH-160A electronic scanner with a 3.75 or 2.5 MHz transducer (Toshiba Inc, Tokyo, Japan) and recorded on $\frac{3}{4}$ inch videotape for subsequent review.

Initially, we undertook a comprehensive echocardiographic study to obtain optimal imaging of the aortic root, ascending aorta, and aortic valve, without knowledge of the patients' clinical characteristics. For this purpose, we used a 3.75 MHz transducer from parasternal long and short axis projections, with the transducer focus at the level of the aortic valve. For better imaging of the proximal ascending aorta, we used a higher left parasternal window in most patients, with a parallel gradual decrease of the overall gains for better delineation of the inner walls of the aortic root. From these projections and with decreased gains (approximately by 10 db), we measured the internal diameter of the aortic ring (annulus), the diameter at the mid-level of the sinuses of Valsalva (mid-sinus level), and the diameter at the supra-avalvular ring (sinotubular junction). The minimum luminal diameter between aortic valve and supra-avalvular level was then noted in order to document the origin of the maximum gradient. We confirmed the minimum aortic narrowing with colour Doppler superimposed on grey scale imaging to identify the maximal aliasing or turbulence region. We also used short axis projections of the aortic valve, sinuses, and ascending aorta on cine-loop format and assessed the thickness of the aortic valve by measuring the most thickened cusp at the maximally affected area.

After optimal description of the aortic root and valve anatomy and identification of the

level of maximal narrowing, we employed a 2 MHz stand alone continuous wave Doppler transducer to measure the maximum transaortic velocity, assuming that it corresponds to the minimum aortic narrowing described. For this, all possible transducer positions were used including apical, right parasternal, high parasternal, and suprasternal positions, and the highest transaortic velocity was recorded.

For optimal assessment of the overall involvement of the aortic valve and root, we devised an aortic valve root score. Aortic valve thickening was scored as follows: 1, normal thickness (≤ 2 mm); 2, localised or nodular thickening involving one cusp; 3, diffuse thickening (2–4 mm) involving one or more cusps; 4, diffuse thickening > 4 mm involving one or more cusps. Valve mobility was scored as: 1, normal; 2, mildly restricted; 3, moderately restricted; 4, severely restricted (almost fixed). The aortic root was measured at mid-sinus level, evaluated subjectively for evidence of increased echogenicity/brightness, and gauged as follows: 1, normal appearance with no evidence of narrowing (diameter > 20 mm); 2, increased echogenicity but no evidence of narrowing (diameter > 20 mm); 3, supra-avalvular narrowing (diameter 16–20 mm); 4, marked supra-avalvular narrowing (diameter < 16 mm). The aortic valve root score was calculated by summing all three scores and thus ranged from 3 (normal) to 12 (worst).

LABORATORY PROCEDURES

Blood samples were obtained after an overnight fast of 12 hours for lipid and lipoprotein analysis within three months of echocardiography. Serum cholesterol, triglyceride, and high density lipoprotein (HDL) cholesterol were assayed enzymatically by automated procedures, HDL cholesterol being assayed after preliminary precipitation of very low density lipoprotein and LDL with heparin and manganese.⁹ LDL cholesterol was calculated¹⁰ and Lp(a) was assayed using a commercial enzyme linked immunosorbent assay (Tint Elize, Biopool, Umeå, Sweden).

The "cholesterol-years" score (CYS) for assessing the lifelong exposure to hypercholesterolaemia was calculated by modifying the formula of Hoeg *et al.*,¹¹ as follows: the total cholesterol concentration (mmol/l) of each patient at the time of original diagnosis was multiplied by the age of patient at diagnosis; the mean cholesterol concentration in the patient while on treatment before the introduction of statins was multiplied by the number of years of non-statin treatment; finally, the mean cholesterol concentration on statin treatment was multiplied by the number of years of statin treatment up to the time of echocardiographic examination. The sum of the pretreatment, non-statin treatment, and statin treatment cholesterol $\times$ years scores gave the total CYS. The average lifetime CYS was derived by dividing the total CYS by the current age of the patient.

Table 1 Age, sex, and serum lipids of patients with homozygous and severe heterozygous FH at the time of diagnosis. The mean (SD) values in the remaining 75 heterozygotes are also shown

Patient	Age (years)	Sex	Phenotype	At diagnosis (mmol/l)			
				TC	TG	HDL-C	LDL-C
1	28	F	Homozygous	17.5	0.72	1.04	16.1
2†	26	F	Homozygous	17	0.9	0.9	15.7
3	12	F	Homozygous	28.7	1.72	0.65	27.3
4	26	M	Homozygous	24.4	0.7	0.9	23.2
5	32	M	Homozygous	22.3	0.9	0.7	21.2
6	15	F	Homozygous	17	1.5	1.3	15
7	70	F	Severe heterozygous*	18.1	1.9	1.2	16
8	61	F	Severe heterozygous	17.4	1.1	1	15.9
9	62	M	Severe heterozygous	19	3.5	0.9	16.5
10–84	51.6 (14)	47M/ 28F	Heterozygous	10.4 (1.8)	1.8 (1.7)	1.16 (0.26)	8.5 (2.0)

HDL-C, high density lipoprotein cholesterol; LDL-C, low density lipoprotein cholesterol; TC, total cholesterol; TG, triglyceride.

*TC >17 mmol/l.

†Patient No 2 was reported previously.¹³

STATISTICAL ANALYSIS

Values are expressed as mean (SD). Logarithmic transformation of Lp(a) values was performed in order to normalise its highly skewed distribution and enable calculation of the geometric mean. Differences between means were compared with unpaired Student's *t* test. The significance of any differences in proportions was tested with the χ^2 statistic. Spearman's rank correlation coefficients were used to assess relations between variables. Multivariate stepwise regression analysis was used to assess independence of associations. A *p* value of < 0.05 was considered statistically significant.

Results

HOMOZYGOTES

Tables 1 and 2 show the lipid characteristics of the six homozygous subjects, and table 3 shows the echocardiographic findings. None of the homozygotes was hypertensive or a smoker. Case 6 had undergone cardiac surgery in 1984 and Doppler information was not available at that time. Therefore in table 3, we give her peak to peak transaortic gradient obtained by cardiac catheterisation before surgery and the corresponding lipid values.

Four homozygotes (cases 1, 2, 5, and 6) had severe aortic stenosis as indicated by mean gradients of > 50 mm Hg and a peak gradient of 130 mm Hg in case 6. Their mean (SD) CYS was 387 (124) mmol/l × years. The aortic cusps were diffusely thickened (3.2 to 4.6 mm), more so at the tips but without commissural fusion; their aortic valve root scores ranged from 10 to 11. Figure 1 shows the histological findings of

the excised aortic valve of case 2. There was focal deposition of calcium, fibrosis, and chronic inflammatory cells. Lipid stains showed central cholesterol clefts and oxidised lipid both within macrophages and extracellularly. None of the patients had a bicuspid aortic valve. The anterior mitral leaflet was also thickened to a lesser degree in cases 2, 5, and 6, but the overall valve mobility was normal.

Echocardiographic evidence of supravalar narrowing was obtained in four homozygotes (cases 1, 2, 5, and 6) in whom there was highly echogenic material encircling the aortic root and mainly arising from the sinuses of Valsalva, with a considerable reduction of the diameter at mid-sinus level (fig 2A). Cases 2 and 6 underwent cardiac catheterisation, which confirmed the presence of supravalar atheroma by the funnelled appearance of the ascending aorta (fig 2B). The gradient in both cases was transvalvar

In cases 3 and 4 the aortic root was highly echogenic but less thickened than in the other homozygotes and the aortic cusps were normal. The diameter at mid-sinus level was the same as that of the aortic annulus; this is considered abnormal since a mean increase of 12.5% is expected at mid-sinus level compared with the annulus diameter.¹² These findings, in association with the accelerated velocity along the ascending aorta, were suggestive of supravalar aortic atheromatous involvement, although this was relatively mild, with an aortic valve root score of 5 in both cases.

None of the homozygotes had resting left ventricular wall motion abnormalities, which

Table 2 Age treatment started, serum lipids on treatment, and cholesterol–years score in patients with homozygous (Nos 1–6) and severe heterozygous (Nos 7–9) familial hypercholesterolaemia. The mean (SD) values in the remaining 75 heterozygotes are also shown

Patient	Age treatment started (years)	At echocardiography			
		TC (mmol/l)	LDL-C (mmol/l)	Lp(a) (mg/dl)	CYS (mmol/l × years)
1	9	12.8	11.2	60	465
2	16	11.5	9.9	23	400
3	3	8	7.1	63	194
4	2	8	6.8	35	480
5	12	11.3	9.6	5	475
6	7	10.3	8.3	14	207
7	63	6.7	4.8	49	1217
8	43	8.8	7	32	972
9	40	10.5	8.2	5	990
10–84	41.6 (13.3)	6.8 (1.7)	4.9 (1.6)	21.5 (4.1)*	501.4 (136)

CYS, cholesterol–years score; LDL-C, low density lipoprotein cholesterol; Lp(a), lipoprotein(a); TC, total cholesterol.

*Geometric mean.

Table 3 Echocardiographic findings in patients with homozygous (Nos 1–6) and severe heterozygous (Nos 7–9) familial hypercholesterolaemia, and mean (SD) values in the remaining 75 heterozygotes

Patient	Supravalvar atheroma present	Annulus diameter (mm)	Mid-sinus diameter (mm)	Diameter of supravalvar ring (mm)	Maximum thickness of aortic cusps (mm)	Mobility of aortic cusps	Vmax (m/s)	Peak gradient (mm Hg)	Mean gradient (mm Hg)	AVRS
1	Yes	17	14.4	17.2	3.4	Mildly restricted	4.4	77	52	10
2	Yes	20	14.5	16.5	4.6	Mildly restricted	6	147	92	11
3	Possibly	17	17	16.1	<2	Normal	2.5	25	13	5
4	Possibly	19	19	*	<2	Normal	2.9	33	19	5
5	Yes	18	15	15.5	3.2	Mildly restricted	4.7	88	51	10
6	Yes	17	15	16	3.2	Mildly restricted	–	130†	–	10
7	Uncertain	18.1	19.8	18.6	3.6	Severely restricted	4.9	98	63	9
8	Yes	17.4	16.1	18.2	3.8	Mildly restricted	5.4	118	74	9
9	No	25	30	23	3.3	Normal	1.5	9	5	5
10–84	6 with bright aortic root	22.4 (2.5)	31.5 (4.2)	26.5 (3.8)	2.7 (0.5)‡	Normal§	1.3 (0.3)	6.7 (3.4)	3.8 (2.3)	3.5 (0.87)

Vmax, peak transaortic velocity.

*No clear visualisation of supravalvar ring; †peak to peak gradient on cardiac catheterisation; ‡in the 14 heterozygotes with thickened aortic valve; §only in one case mildly restricted.

excluded previous myocardial infarction. Table 4 shows the coronary anatomy, the condition of the coronary sinuses, and the type of cardiac surgery undertaken in three homozygotes (cases 2, 5, and 6). In all cases the right coronary ostium was narrowed and in case 5 both ostia were stenosed. The distal coronary arteries were free from significant disease with the exception of case 2, where there was a 60% stenosis at the middle third of the left anterior

descending coronary artery. Cases 2 and 6 underwent successful aortic root reconstruction, aortic valve replacement, and coronary artery bypass grafting. Case 5 underwent coronary artery bypass grafting alone.

HETEROZYGOSES

Serum lipids and echocardiographic findings in three heterozygotes (cases 7, 8, and 9) with severe hypercholesterolaemia (cholesterol at original diagnosis > 17 mmol/l, total CYS > 900 mmol/l × years), and the mean values of the remaining 75 heterozygotes are presented in tables 1, 2, and 3. Nine (11.5%) were hypertensive and seven (9%) were smokers.

The mean CYS in all heterozygotes was 523 (175) mmol/l × years. Cases 7 and 8 had severe aortic stenosis, while case 9 had diffuse thickening of the non-coronary cusp (3.3 mm) without evidence of stenosis. Both the heterozygotes with severe aortic stenosis underwent cardiac catheterisation. In case 7, the aortic valve and root were heavily calcified and the left descending coronary artery was blocked proximally. In case 8, the aortic valve was diffusely thickened (3.8 mm) and the root had a funnelled appearance, typical of supravalvar narrowing. There was an 80% stenosis of the left descending coronary artery and the right ostium was blocked. The aortic valve root score ranged from 5 to 9 in these three patients.

Of the remaining 75 heterozygotes, only one had mild aortic stenosis (mean gradient 18 mm Hg). In this patient the right and non-coronary cusps had diffuse thickening (3.7 mm) and their mobility was slightly restricted. In addition, the aortic root was highly echogenic but without supravalvar narrowing. Among the others, there was increased echogeneity of the aortic root in six patients, without supravalvar narrowing (mid-sinus diameter > 20 mm). Sixteen patients had resting left ventricular akinesis on the echocardiographic study, implying previous transmural myocardial infarction. The mean aortic valve root score of these 75 patients was 3.5 (table 3).

Table 5 shows the correlations between maximum transaortic velocity, peak gradient, and mean gradient across the aortic valve with age and serum lipids. Multiple regression analysis was applied to assess the independence of association between mean aortic gradient and the variables in table 5. In a stepwise

Figure 1 (A) Section of the resected valve showing fibrosis (left), cholesterol clefts (centre), and nodular calcification (right) (×42). (B) Higher power view of the valve stained with oil red O showing insoluble lipid (ceroid) and cholesterol clefts (×210). (By kind permission of Dr D Parums.)

Figure 2 (A) Parasternal long axis view showing a highly echogenic aortic root (arrows) and thickening of the aortic valve. The focus zone of the ultrasound beam is at the anterior aortic wall. (B) Aortography of the same patient showing the characteristic funnelling of the proximal ascending aorta.

regression model with mean gradient as dependent variable, only CYS ($\beta = 0.042$, $p = 0.0001$) and age ($\beta = -0.47$, $p = 0.002$) were significant determinants of the mean gradient, accounting for 40% and 9% of its variation respectively. Figure 3 shows the correlation between mean gradient (after logarithmic transformation) and CYS.

Figure 3 Correlation between mean gradient across the aortic valve (after logarithmic transformation) and cholesterol-years score (CYS) in patients with heterozygous familial hypercholesterolaemia ($n = 78$).

Seventeen heterozygotes (22%) had thickening of aortic cusps, but only in two (2.6%) was this associated with severe aortic stenosis and in one with mild aortic stenosis. Five of the heterozygotes studied were over the age of 70 years, and of these three had localised aortic valve thickening but none had aortic stenosis. In an effort to minimise the contribution from senile degenerative aortic valve disease, we compared the serum lipids in the 73 heterozygotes aged ≤ 70 years according to the presence of aortic valve thickening. The heterozygotes with thickened aortic valves were older and had significantly higher total and LDL cholesterol at diagnosis and had higher total and average CYS/year, but there was no difference in serum lipids or Lp(a) concentrations at the time of echocardiography (table 6). There was also a significant positive correlation between aortic valve root score and total CYS in the heterozygotes aged ≤ 70 years (fig 4).

Discussion

In this study we found paradoxically that homozygote patients with FH get severe aortic root and valve involvement at lower CYS scores than heterozygotes. This suggests that aggregates of LDL at the high concentrations found in homozygotes, but seldom in heterozygotes, are responsible, rather than the lifetime elevation of cholesterol levels found in the heterozygous group.

Histologically the aortic valve and root lesions consist of foam cells, cholesterol clefts, and fibrocalcific deposits. The presence of intracellular lipid and cholesterol clefts within the cuspal tissues is unique to homozygous FH and differentiates these lesions from other forms of aortic stenosis. Despite the significant thickening of the aortic cusps, there was no commissural fusion and good overall cusp mobility, though there was incomplete opening caused by atheromatous accumulation in the sinuses of Valsalva. This mechanism of aortic stenosis explains the clinical paradox of a well preserved aortic closure sound. Our findings are consistent with previous reports which ascribe the aortic stenosis in homozygotes to a

Table 4 Coronary anatomy, condition of sinuses of Valsalva, and type of cardiac surgery in three homozygotes

Patient	Coronary anatomy	Condition of sinuses of Valsalva	Type of operation	Age at operation (years)
2	Right ostial occlusion, LAD stenosis	Right coronary sinus obliterated	ARR (pericardial patch), AVR, 2 grafts	26
5	Right and left ostial stenosis	All sinuses infiltrated	4 grafts*	21
6	Right ostial occlusion	Right and non-coronary sinuses obliterated	ARR (pericardial patch), AVR, 1 graft	15

ARR, aortic root reconstruction; AVR, aortic valve replacement; LAD, left anterior descending coronary artery.

*The coronary artery bypass graft was performed in 1985 when the peak to peak gradient across the aortic valve was 20 mm Hg.

Table 5 Spearman's correlation coefficients between maximum transaortic velocity, peak gradient, mean gradient with age, lipids, and cholesterol-years score in all heterozygotes ($n = 78$)

Variable	Vmax	Peak gradient	Mean gradient
Age	$r = 0.29$; $p = 0.01$	$r = 0.22$; NS	$r = 0.21$; NS
TC (d)	$r = 0.52$; $p = 0.0001$	$r = 0.51$; $p = 0.0001$	$r = 0.51$; $p = 0.0001$
TG (d)	$r = 0.03$; NS	$r = 0.05$; NS	$r = 0.05$; NS
HDL-C (d)	$r = 0.01$; NS	$r = 0.08$; NS	$r = 0.09$; NS
LDL-C (d)	$r = 0.56$; $p = 0.0001$	$r = 0.56$; $p = 0.0001$	$r = 0.55$; $p = 0.0001$
TC (e)	$r = 0.07$; NS	$r = 0.08$; NS	$r = 0.09$; NS
LDL-C (e)	$r = 0.09$; NS	$r = 0.11$; NS	$r = 0.11$; NS
Lp(a)* (e)	$r = 0.07$; NS	$r = 0.07$; NS	$r = 0.08$; NS
CYS	$r = 0.53$; $p = 0.0001$	$r = 0.59$; $p = 0.0001$	$r = 0.58$; $p = 0.0001$

CYS, cholesterol-years score; HDL-C, high density lipoprotein cholesterol; LDL-C, low density lipoprotein cholesterol; Lp(a), lipoprotein(a); TC, total cholesterol; TG, triglyceride; Vmax, maximum transaortic velocity.

*After logarithmic transformation.

(d), at initial diagnosis; (e), at the time of echocardiography.

valvar gradient secondary to supra-avalvular narrowing,^{3 13} although there are some cases where the gradient was predominantly supra-avalvular, apparently owing to gross narrowing at the sinotubular junction.¹⁴ Interestingly, although the aortic valve appeared normal in two homozygotes, the ascending aorta was highly echogenic with mild narrowing at mid-sinus level suggestive of supra-avalvular atherosclerosis. It is possible that the absence of aortic valve involvement and the relatively mild aortic root disease in those two patients is a result of the early introduction of lipid lowering treatment (≤ 3 years of age), consisting initially of statins and later the combination of statins with LDL apheresis.

Regression of coronary atherosclerosis has been reported with LDL apheresis in homozygotes^{15 16} and consequently one can speculate that a similar beneficial effect could be expected for the aortic valve and root involvement.¹⁷ In homozygotes the arrest or regression of aortic root involvement should be the main therapeutic aim, since early fatal coronary events are usually related to ostial occlusion secondary to lipid deposition in the sinuses of Valsalva. However, delayed introduc-

Figure 4 Correlation between cholesterol-years score (CYS) and aortic valve root score (AVRS) in heterozygotes for familial hypercholesterolaemia aged ≤ 70 years.

tion of aggressive lipid lowering treatment carries the theoretical risk of worsening the valvar aortic stenosis by converting the lipid rich aortic cusps into fibrotic and more rigid cusps, since the mobilisation and removal of cholesterol may paradoxically accelerate fibrosis, as has been shown in the coronary arteries of an FH patient.¹⁸ This underlines the need to start treatment at an early stage.

The anterior mitral leaflet showed mild generalised thickening in three homozygotes (cases 2, 5, and 6), probably owing to lipid infiltration, but no evidence of valvar dysfunction. However, a 9 year old homozygote has been reported with mitral regurgitation caused by lipid infiltration of the mitral leaflets along with marked fibrosis and atrophy of the papillary muscles.¹⁹

This is the first study to report a positive correlation between the mean transaortic gradient, severity of aortic valve root involvement, and CYS in heterozygotes with familial hypercholesterolaemia. These associations suggest that lifelong exposure to high cholesterol concentrations largely determines the severity of aortic stenosis in this group and can result in combined involvement of the aortic valve and root, which is manifested by some degree of aortic valve thickening and increased echogenicity of aortic root. However, severe root involvement is rare in heterozygous FH, in contrast with homozygous FH.

We analysed separately the lipid and echocardiographic findings in the three heterozygotes with the most marked hypercholesterolaemia (initial cholesterol concentrations > 17 mmol/l, CYS > 900 mmol/l $\times$ years), two of whom had severe aortic stenosis. Case 8 had all

Table 6 Serum lipids and lipoproteins, cholesterol-years score, and average cholesterol-years score in 73 heterozygotes for familial hypercholesterolaemia aged ≤ 70 years according to the presence of aortic valve thickening

Thickened aortic valve	Age (years)	Lipids at diagnosis				Lipids at echocardiography				Average CYS/year
		TC (mmol/l)	TG (mmol/l)	HDL-C (mmol/l)	LDL-C (mmol/l)	TC (mmol/l)	LDL-C (mmol/l)	Lp(a) (mg/dl)	CYS	
Yes (n = 14)	58 (9.5)	12.9 (3.3)	1.9 (0.76)	1.1 (0.29)	11.6 (3.6)	7.4 (2.4)	5.5 (2.2)	37.4 (2.9)*	692 (218)	12.2 (12.7)
No (n = 59)	48.5 (12.2)	10.3 (1.8)	1.9 (1.8)	1.2 (0.26)	8.3 (1.9)	6.7 (1.6)	4.8 (1.5)	22.4 (3.9)*	467 (129)	9.8 (1.7)
p Value	0.006	0.0002	NS	NS	0.0004	NS	NS	NS	0.0001	0.0001

Values are mean (SD); CYS, cholesterol-years score; HDL-C, high density lipoprotein cholesterol; LDL-C, low density lipoprotein cholesterol; Lp(a), lipoprotein(a); TC, total cholesterol; TG, triglyceride. *Geometric mean.

the characteristic features of valvar aortic stenosis secondary mainly to supra-avalvar narrowing which prevented the normal cusp opening. Case 7 was older (70 years), the aortic valve and root were markedly calcified, the aortic cusps had little mobility, and there was no significant narrowing at mid-sinus level, nor was there the characteristic funnelling of the aortic root on aortogram. On the basis of these findings it is unclear whether the involvement of aortic valve and root in this patient was the result of lipid infiltration followed by severe calcification or whether it was caused by degenerative aortic valve disease, or a combination of both. The third heterozygote (case 9), despite the early development of coronary artery disease (coronary artery bypass graft at 42 years, redone at 57 years), showed only mild diffuse thickening of the non-coronary aortic cusp without aortic stenosis or supra-avalvar narrowing. Of the remaining 75 heterozygotes, one had mild aortic stenosis without supra-avalvar narrowing, but the remainder were normal.

To shed further light on the mechanism of aortic valve thickening in heterozygotes we subdivided them according to the presence or absence of pathological valve thickening. Heterozygotes aged > 70 years were excluded since it is known that age contributes to the degenerative thickening of aortic valve in the general population.²⁰ Age, cholesterol, and LDL cholesterol at diagnosis and the CYS were the main predictors of aortic valve thickening. Lp(a) did not differ significantly between the subgroups, although there was a trend for higher levels in the heterozygotes with thickened aortic cusps. A recent study showed that Lp(a) was closely related to aortic valve sclerosis in a rural Japanese population,²¹ and the authors postulated that the high affinity of Lp(a) for glycosaminoglycans enhances the accumulation of Lp(a) in the connective tissue of aortic cusps. The absence of this association in our population presumably reflects differences in patient selection or treatment. To minimise the confounding effect of age further, we calculated the average CYS by dividing the CYS by age. This was significantly higher in the heterozygotes with aortic valve thickening, supporting our hypothesis that lifelong exposure to high cholesterol levels represents the main risk factor for the thickening of aortic cusps in this situation, possibly by accelerating age related degenerative aortic valve lesions. A recent study²² reinforces our hypothesis, suggesting that hypercholesterolaemia may be an aetiological factor in the development of aortic stenosis.

We conclude that aortic stenosis is very common in homozygotes, and aortic root involvement is always present. It appears that in homozygotes it is the exposure to a very high LDL cholesterol concentration rather than the lifelong cholesterol burden that determines aortic valve and root involvement. Early initiation of treatment slows the rate of progression, not only of the coronary atherosclerosis but of the aortic valve and root involvement. In

contrast, aortic valve and root involvement are rare in heterozygotes and occur only with severe and prolonged hypercholesterolaemia—that is, when the total CYS is > 900 mmol/l × years. The occurrence of similar lesions in homozygotes with a CYS only half as high suggests that at very high concentrations, seen only in homozygotes, LDL may undergo spontaneous aggregation when subjected to turbulent flow during systole. Aggregation can be induced in vitro by vortexing concentrated LDL, and such aggregates are avidly taken up by scavenger receptors on monocyte macrophages²³ and are thus potentially atherogenic. Although speculative, this proposal provides a possible explanation for the different distribution and much greater severity of aortic atherosclerosis in homozygotes compared with heterozygotes.

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IMAGES IN CARDIOLOGY

“Steam iron” appearance of chronic thoracic aortic dissection on angiography

A 42 year old man was referred for investigation of an abnormal chest x ray that showed profound widening of the mediastinum suggestive of a large thoracic aneurysm. At the age of 35 years he had collapsed while at work. At the time no specific cause had been found. He was otherwise fit and well. Thoracic computed tomography had showed dilatation of the aorta extending from the arch to the level of the renal arteries but did not confirm dissection. Trans-thoracic echocardiography had suggested an intimal flap within the descending aorta. Transoesophageal echocardiography was not routinely available at that time.

The image shows a contrast angiogram of the descending aorta revealing extensive dissection with compression of the true lumen by a large false lumen. The true lumen communicates with the false lumen through a number of stripped off intercostal arteries giving a series of jets akin to those seen when using a steam iron.

Following discussion with cardiothoracic surgeons it was decided to treat the patient conservatively.

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