

Apolipoprotein(a) Phenotypes, Lp(a) Concentration and Plasma Lipid Levels in Relation to Coronary Heart Disease in a Chinese Population: Evidence for the Role of the apo(a) Gene in Coronary Heart Disease

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Abstract

Elevated lipoprotein(a) (Lp[a]) concentrations are associated with premature coronary heart disease (CHD). In the general population, Lp(a) levels are largely determined by alleles at the hypervariable apolipoprotein(a) (apo[a]) gene locus, but other genetic and environmental factors also affect plasma Lp(a) levels. In addition, Lp(a) has been hypothesized to be an acute phase protein. It is therefore unclear whether the association of Lp(a) concentrations with CHD is primary in nature. We have analyzed apo(a) phenotypes, Lp(a) levels, total cholesterol, and HDL-cholesterol in patients with CHD, and in controls from the general population. Both samples were Chinese individuals residing in Singapore. Lp(a) concentrations were significantly higher in the patients than in the population (mean 20.7 ± 23.9 mg/dl vs 8.9 ± 12.9 mg/dl). Apo(a) isoforms associated with high Lp(a) levels (B, S1, S2) were significantly more frequent in the CHD patients than in the population sample (15.9% vs 8.5%, $P < 0.01$). Higher Lp(a) concentrations in the patients were in part explained by this difference in apo(a) allele frequencies. Results from stepwise logistic regression analysis indicate that apo(a) type was a significant predictor of CHD, independent of total cholesterol and HDL cholesterol, but not independent of Lp(a) levels. The data demonstrate that alleles at the apo(a) locus determine the risk for CHD through their effects on Lp(a) levels, and firmly establish the role of Lp(a) as a primary genetic risk factor for CHD. (*J. Clin. Invest.* 1992. 89:1040-1046.) Key words: atherosclerosis • risk factor • quantitative genetic trait • population genetics • genetic epidemiology

Introduction

Lipoprotein(a) (Lp[a])¹ is a macromolecular complex in human plasma that is assembled from a low-density lipoprotein (LDL) and apolipoprotein (a) (apo[a]) (1). Shortly after the discovery of the genetic Lp(a) system by Berg (2), it was recognized that high concentrations of Lp(a) in plasma are asso-

ciated with coronary heart disease and early myocardial infarction (3-6). This observation was confirmed by numerous studies in different ethnic groups, using different endpoints for definition of coronary artery disease (CAD), and different methods to measure Lp(a) in plasma (7). The vast majority of these studies are cross-sectional case-control studies. Because of the strong genetic determination of Lp(a) concentrations, Lp(a) is widely considered a primary genetic risk factor for coronary heart disease (CHD), despite the lack of any large prospective epidemiological studies.

Originally, Lp(a) was described as a qualitative autosomal dominant trait (2). Later studies showed that Lp(a) is a continuous quantitative trait under the control of a single major gene (8-12). The nature of this gene remained elusive until the discovery of a genetic size polymorphism of apo(a) and its association with Lp(a) levels in plasma (13-15). We have described six different apo(a) isoforms, designated F, B, S1, S2, S3, and S4, that vary in size from ~ 400 to over 800 kD. In the population, the sizes of apo(a) isoforms are inversely associated with Lp(a) levels, and in families, apo(a) isoforms and levels cosegregate (13-17). Large isoforms are associated with low Lp(a), and small isoforms with high Lp(a), in plasma. The human apo(a) gene codes for a large protein with a high degree of homology to the plasma zymogen plasminogen (18, 19). It contains a protease domain, one so-called kringle 5 domain, and multiple complete, or nearly identical, tandem repeats of a plasminogen like kringle 4 domain. The introns in apo(a) are also highly conserved (1, 20) (H. J. Menzel, J. Pfitscher, and G. Utermann, unpublished). The gene locus for apo(a) on chromosome 6 q2.6-2.7 is highly polymorphic. The alleles at this locus determine a genetic size polymorphism of apo(a) (1, 13). It has been shown by quantitative Southern blotting, and more recently by pulsed field gel electrophoresis, that this polymorphism results from differences in the number of tandem kringle 4 repeats in the apo(a) gene (20-22) (H. G. Kraft, S. Köchl, and G. Utermann, unpublished). Together, the family, extended population, and molecular genetic studies have shown that the apo(a) gene locus determines both the size of the apo(a) isoform, and the concentration of Lp(a) in plasma.

The mechanism by which apo(a) size determines Lp(a) concentrations is presently not understood. However, in healthy Caucasians only about 40% of the variability in Lp(a) levels is explained by the protein size polymorphism, whereas the rest is presently unexplained. Other genetic, endogenous, and exogenous factors that influence Lp(a) levels have been identified, including defective alleles at the LDL-receptor gene locus which result in a two- to threefold elevation (23). Lp(a) levels may also be elevated secondary to disease. In end-stage renal disease, Lp(a) levels are elevated two- to threefold over controls (24, 25) (H. Dieplinger and G. Utermann, unpublished). Fur-

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1. Abbreviations used in this paper: CHD, coronary heart disease; Lp(a), lipoprotein(a).

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ther, it has been shown that Lp(a) may behave like an acute phase reactant (26). The development of atherosclerosis is a longstanding process that takes decades before clinical symptoms are manifest. Therefore, the possibility that elevated Lp(a) levels are secondary to the disease process cannot be excluded. The same argument does not apply to the apo(a) isoform phenotypes that represent invariable genetic markers with a significant effect on Lp(a) levels. The present study was designed to test whether Lp(a) is a primary risk factor for CHD. To this end, we determined apo(a) types, Lp(a) levels, total cholesterol, and HDL-cholesterol in Chinese patients with CHD, and a population sample from Singapore. This population was selected because a much higher fraction of the variability of Lp(a) levels is explained by the apo(a) size polymorphism than in Caucasians (27). This demonstrated the expected association of CHD with elevated Lp(a) levels, but, more importantly, it also demonstrated an association of CHD with the apo(a) phenotype.

Methods

CHD-patients and population sample. Chinese patients with CHD were selected from a series of consecutive patients attending the Singapore Chest and Heart Clinics in the second half of 1990 for suspected CHD. Patients with a positive stress test (Bruce protocol) were evaluated for presence of CHD by coronary angiography. Inclusion criteria for this study were 50% or more stenosis of at least one of the major coronary arteries. Patients with less than 50% stenosis, valve disease, or cardiomyopathy, were excluded. 170 patients (136 men, 34 women) fulfilled the inclusion criteria. For 162 patients, the complete data set was available. Only those have been analyzed.

The healthy population sample (controls) was recruited from subjects who underwent routine medical examination in connection with their employment. Inclusion criteria were normal routine biochemical laboratory tests, a normal resting ECG, absence of a history of cardiovascular disease, and diabetes in the subject and first-degree relatives. 211 subjects (110 men, 101 women) fulfilled the inclusion criteria. Cases and control subjects were from the same ethnic group. The Chinese in Singapore, including both the cases and controls, are Han Chinese from Southern China, and represent the second and third generation of immigration. Fasting blood was drawn into EDTA and centrifuged at low speed. Plasma was stored at -20°C until shipped on dry ice to Innsbruck. All laboratory analysis was carried out within a maximum of 6 mo after blood had been drawn.

Laboratory procedures. Apo(a) phenotyping was performed by SDS-polyacrylamide gel electrophoresis of plasma under reducing conditions, followed by immunoblotting as outlined (15), with slight modification. In brief, 2 μl plasma was added to 50 μl 5% (wt/vol) SDS, 0.02 ethylmorpholine pH 8.6, 2 μl β -mercaptoethanol, and the solution was heated for 3 min in a microwave oven. 4 μl 1.5% (wt/vol) bromophenol blue in 10% glycerol were added. A 5- μl aliquot was applied to a 6.6% polyacrylamide gel prepared and run according to Neville (28). Immunoblotting was performed as described, using the monoclonal anti-apo(a) antibody 1A² (29) which does not crossreact with plasminogen. A goat anti-mouse peroxidase conjugate (Dako, Copenhagen, Denmark) was used as second antibody.

Lp(a) quantification was performed by a sandwich-ELISA, essentially as described (29), using a rabbit polyclonal affinity purified anti-Lp(a) antibody for coating, and the horseradish peroxidase conjugated monoclonal anti-apo(a) antibody 1A² for detection. Cholesterol and HDL-cholesterol were determined enzymatically using commercial test kits (Boehringer Mannheim Diagnostics, Mannheim, FRG).

Statistical analysis. Standard statistical methods were used throughout, and were implemented using BMDP Stat. Software, Inc. (Los Angeles, CA) (30). Analysis of these data began with a series of

univariate tests to assess the difference of each variable individually between the cases and the control samples from Singapore. These univariate tests were then followed by multivariate analyses to account for the interrelationships among the independent variables as they combine to predict whether an individual one is a case or a control. Routine parametric adjustment of the predicting variables such as Lp(a), and cholesterol levels for the concomitant effects of age and sex, was not carried out. Rather, adjustment for these relationships were incorporated directly into the multivariate analyses (31). Equality of the apo(a) isoform phenotype frequencies among strata was tested, using a likelihood ratio test. Because the distribution of plasma Lp(a) levels in these (see below) and other data (27) is highly nonnormal, nonparametric statistics such as the Kruskal-Wallis test (32, 33) were used to test the equality of the Lp(a) levels among strata. For consistency, the same tests were used to test the equality of the other phenotypes between the cases and controls. Stepwise logistic regression (31, 34) using a maximum likelihood procedure (35) was used to assess the relationship between disease status and the set of interrelated predictive variables. Because of their widely accepted role in influencing CHD, gender and age were always included in the logistic model; they were not part of the hypothesis testing hierarchy. Both forward and backward stepping procedures arrived at consistent results for these data (data not shown).

Results

Apo(a) types and Lp(a) concentrations in a Chinese control population. A description of Lp(a) and lipid levels in the study population are given in Table I, and the distribution of Lp(a) levels in the sample from the Chinese population in Singapore is shown in Fig. 1 A. The mean Lp(a) concentration (8.9 ± 12.9 mg/dl, Table I) is lower in this sample than in the general Caucasian populations (27), and the distribution is highly skewed toward lower levels. Frequencies of apo(a) size isoform phenotypes are given in Table II. Very large isoforms (designated S4) predominate in the Singapore Chinese. The single-band S4 type was, by far, the most common apo(a) isoform phenotype observed in this sample. Taken together, the phenotypes with only high relative molecular mass apo(a) isoforms (S3, S4, 0) had a relative frequency of 91.4%. Phenotypes with at least one smaller isoform (B, S1, S2) were much less common (8.5%). There exists an inverse association of apo(a) isoform size with Lp(a) concentrations in plasma (Table III). Lp(a) levels were significantly different between apo(a) types (Kruskal-Wallis test = 81.6, 8 d.f., $P < 0.001$). Using the R^2 value from the analysis of variance, $\sim 53\%$ of the variation in Lp(a) concentration was explained by the size polymorphism of apo(a).

These results agree closely with our previous results on an

Table I. Mean Age and Lipid Levels in Chinese CHD Patients and a Control Population Sample

	Controls <i>n</i> = 210	CHD patients <i>n</i> = 162	<i>P</i>
Age yr	37.3 \pm 14.5 (17–75 yr)	57.6 \pm 8.5 (35–81 yr)	<.01
TC mg/dl	215 \pm 50 (99–369 mg/dl)	244 \pm 56 (132–478 mg/dl)	<.01
HDL-C mg/dl	48.0 \pm 13.1 (19–98 mg/dl)	34.0 \pm 9.1 (12–68 mg/dl)	<.01
Lp(a) mg/dl	8.95 \pm 12.98 (0.2–75.0 mg/dl)	20.71 \pm 24.0 (1.0–142.5 mg/dl)	<.01

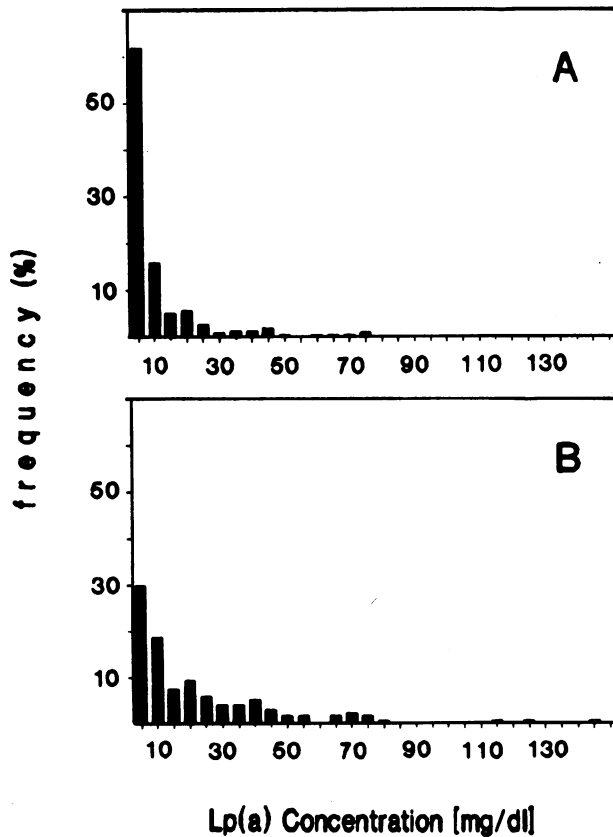


Figure 1. Histograms of Lp(a) level distributions in a Chinese control population (A), and Chinese patients with CHD (B) from Singapore.

independent Chinese sample from Singapore (27). Notably, Lp(a) levels in the Chinese with the S4 isoform are lower than in Caucasians, but levels are higher in Chinese S2 and S3 subjects compared with Caucasians (compare Table III) (27). The variance of Lp(a) levels within apo(a) isoform phenotypes is smaller than in most other groups. Together, this results in a

Table II. Frequencies (%) of Apo(a) Phenotypes in Chinese CHD Patients and a Control Population Sample

Phenotype	Controls (n)*	CHD patients (n)*
B	—	—
S1	1.4 (3)	3.7 (6)
S2	3.3 (7)	6.8 (11)
S3	4.8 (10)	13.0 (21)
S4	56.2 (118)	45.7 (74)
0	16.7 (35)	19.1 (31)
B/S2	—	0.6 (1)
B/S4	—	1.2 (2)
S1/S2	0.95 (2)	—
S2/S3	0.95 (2)	1.9 (3)
S2/S4	1.9 (4)	1.2 (2)
S3/S4	13.8 (29)	6.8 (11)
Total	210	162

* Frequency difference between CHD patients and controls by likelihood ratio test; chi-square = 26.3, 10 d.f., $P < 0.01$.

Table III. Lipoprotein(a) Concentration (mg/dl±SD) in Relation to Apo(a) Type in Chinese Patients with Coronary Heart Disease (CHD) and a Control Population Sample

Phenotype	Controls n = 210*	CHD patients n = 152†
S1	31.7 (28.4)	57.9 (44.8)
S2	44.2 (21.1)	55.4 (24.0)
S3	17.2 (16.4)	29.2 (16.6)
S4	5.7 (7.4)	12.1 (10.2)
0	2.2 (1.8)	3.7 (1.7)
B/S2	—	30.5
B/S4	—	74.7 (5.2)
S1/S2	46.0 (14.1)	—
S2/S3	11.9 (12.2)	88.5 (47.7)
S2/S4	31.5 (12.2)	60.5 (10.6)
S3/S4	10.9 (9.2)	19.2 (9.1)
Total	8.95	20.7

* Kruskal-Wallis Test = 81.63; d.f. 8; $P < 0.0001$.

† Kruskal-Wallis Test = 93.77; d.f. 9; $P < 0.0001$.

relatively larger difference of Lp(a) concentrations between the major types, and in a stronger association of apo(a) type with Lp(a) level.

Relationship between Apo(a) types and Lp(a) levels in Chinese patients with CHD. Lp(a) levels, apo(a) phenotype frequencies, and Lp(a) concentrations in the different apo(a) phenotypes in the Chinese patients with CHD are given in Tables II and III, and the distribution of Lp(a) levels in this sample is shown in Fig. 1 B. The same inverse relationship of apo(a) size with Lp(a) level as in the control population was present in the CHD patients (Table III). On the average, patients with large apo(a) isoforms had lower Lp(a) levels than those with smaller isoforms. The differences in Lp(a) concentrations between apo(a) types were highly significant (Kruskal-Wallis test = 93.8; 9 d.f., $P < 0.001$).

In the patients, the apo(a) size polymorphism explained 65% of the variation in Lp(a) levels. This is higher than in the population sample, and may be due to the increased frequency of those alleles associated with elevated Lp(a) levels in the CHD group (see below).

Thus, Lp(a) levels are mainly determined by apo(a) type in both Chinese controls and patients with CHD. This suggests that the differences in levels are in part due to differences in apo(a) allele frequencies.

Comparison of apo(a) type frequencies and Lp(a) levels between CHD patients and the control population. Lp(a) concentrations are almost twice as high in the CHD patients than in the general population ($P < 0.001$). Mean Lp(a) levels are 8.9 mg/dl in the population sample, and 20.9 mg/dl in patients. Average Lp(a) levels for each phenotype in the CHD patient and control groups are shown in Table III. For each variable, levels are significantly higher in the cases than in the population sample. We next asked whether the effects of the apo(a) types on Lp(a) levels are the same in both groups. Fig. 2 shows the deviation of Lp(a) concentrations from the mean Lp(a) level of the respective group for each of the common apo(a) phenotypes. In patients and controls, phenotypic deviations from group means were virtually identical. These data suggest that the effects of apo(a) types on Lp(a) levels are the same in

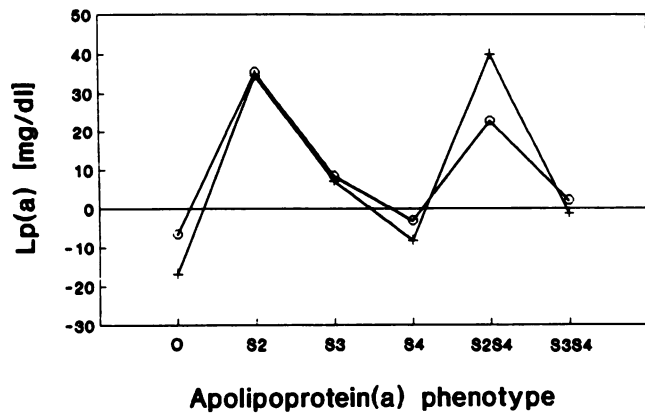


Figure 2. Graphic representation of the deviation of mean Lp(a) concentrations in the common apo(a) types from the respective group means of controls (o), and patients with CHD (+).

both groups. Consistent with this result, using a two-way analysis of variance (36), there is no significant evidence for an interaction effect between apo(a) type and group on Lp(a) levels (data not shown). The central question of this study was whether apo(a) types associated with high Lp(a) levels are more frequent in patients than in the control population. A clear and significant difference in apo(a) isoform frequencies between patients and controls was observed (Table II, $P < 0.01$). Phenotypes associated with high Lp(a) levels in Caucasians and Chinese were almost twice as frequent in the patients. The small B isoform that is associated with the highest Lp(a) levels in all populations studied thus far was only seen in the sample of patients. Therefore, higher Lp(a) levels in the patients may be at least partially attributable to differences in apo(a) phenotype frequencies between patients and controls. However, the differences in Lp(a) levels between CHD patients and the population sample can not be totally accounted for by the differences in apo(a) type frequencies. Notably, Lp(a) levels were also elevated over controls in each of the common apo(a) phenotypes (Table III). This suggests that the relationship between apo(a) type, Lp(a) level, and CHD may be more complex than anticipated.

We next calculated the odds ratios for Lp(a) levels and apo(a) types. These ratios represent the relative odds of having CHD or not having CHD for those with increased Lp(a) levels, or certain apo(a) types, respectively. A value of one indicates that there is no association between CHD and Lp(a) levels, or apo(a) types, respectively. Values significantly greater than one measure the degree to which the odds of disease are increased when the risk factor is present. First, subjects were categorized into those with high and low Lp(a) levels. The histogram of Lp(a) concentrations in the controls show a natural break at 30 mg/dl (Fig. 1). This value also happens to correspond to the 90th percentile. The odds ratio for being in the CHD group for subjects with Lp(a) > 30 mg/dl vs those < 30 mg/dl was 3.975 (Pearson chi-square 20.844, d.f. 1; $P < 0.0001$). Second subjects were divided into those carrying at least one B, S1, or S2 allele vs those without any of these alleles. The odds ratio for subjects with an allele for one of the low molecular weight isoforms in the CHD group was 1.946 (Pearson chi-square 4.211, d.f. 1, $P < 0.05$). Thus, subjects with certain apo(a) alleles have a twofold increased risk for CHD.

Total cholesterol, HDL-cholesterol, Lp(a) level, and apo(a) type as predictors of CHD. Total cholesterol concentrations were 215 ± 50 mg/dl in the population sample, and 244 ± 56 mg/dl in the CHD patients, which is significantly higher ($P < 0.01$). Total cholesterol concentrations in the Chinese control subjects were unexpectedly high and identical to those in Western societies. The reason for this is unknown. Cholesterol levels are, however, typically higher in Singapore Chinese than in Mainland Chinese (37). This probably reflects differences in lifestyle. Whatever the reason for the high cholesterol levels in this particular group, cholesterol levels were even higher in patients with CHD. HDL-cholesterol was significantly lower in the patients (34 ± 9 mg/dl) than in the population (48 ± 13 mg/dl). Thus, the classical lipid risk factors were associated with CHD in the Chinese from Singapore. Stepwise logistic regression was used to assess the independent contribution of the laboratory variable to the prediction of CHD. Because of their widely accepted contribution, sex and age were assumed to be significant predictors, and were automatically incorporated into the logistic risk function. HDL-cholesterol, Lp(a) levels, and total cholesterol, in order, were all independent and significant predictors of disease (Table IV). Apo(a) type was not a significant predictor of disease as long as Lp(a) levels were in the model ($P = 0.106$). However, if Lp(a) levels were excluded, Lp(a) type was a significant predictor of CHD ($P = 0.03$). This result is consistent with a relationship where apo(a) type determines Lp(a) level, and Lp(a) level predisposes to disease. Thus, the important conclusion from the logistic regression analysis is that apo(a) genotype is a significant predictor of disease, and is an independent predictor of CHD after considering the traditional risk factors such as age and cholesterol.

Discussion

This is the first study which firmly establishes a relationship between genetic apo(a) isoforms, Lp(a) levels, and CHD. The comparison between Singapore Chinese patients with CHD, and a Chinese control population, yielded four major results: (a) Lp(a) levels are significantly higher in CHD patients than in the population; (b) Apo(a) types associated with high Lp(a) levels are more frequent in patients with CHD; (c) Lp(a) levels are higher in each common phenotype in the CHD patients; and (d) stepwise logistic regression analysis indicates that apo(a) type is a significant predictor of CHD, independent of total cholesterol and HDL-cholesterol, but not independent of Lp(a) levels.

Table IV. Result of a Stepwise Logistic Regression to Predict CHD Status among Chinese in Singapore

Variable*	Coefficient	(SE)	Chi square	P value
Constant	2.22	1.15	—	—
Gender	0.672	0.427	—	—
Age	0.116	0.0174	—	—
HDL-cholesterol	0.162	0.0253	58.5	<0.001
Lp(a)	0.0513	0.0141	20.1	<0.001
Total cholesterol	0.00933	0.00367	7.17	<0.01

* The constant term along with gender and age were forced into the logistic function. The other variables are presented in the order in which they were entered.

A large number of studies have demonstrated an association of plasma Lp(a) concentration with CHD (3–6, 38–41). Most of these have investigated Caucasian populations, but two were performed on Japanese (6, 42). With the only exception of one small-nested prospective case-control study (43), all studies relating Lp(a) levels to CHD were retrospective case-control studies. Such studies may be biased, since the event itself may alter the factor under study. Even prospective studies may be biased in a situation where there is no single event, but rather the longlasting process of atherosclerosis. Such a problem does not exist if the factor under study is a genetic polymorphism that is not changed by disease or throughout life. Lp(a) levels are widely believed to be largely genetically controlled. However, there is also evidence that several nongenetic factors including postmenopause (44), diabetes mellitus (45, 46), and end-stage renal disease (24) influence Lp(a) concentrations in plasma, and some of them are associated with an increased risk for atherosclerosis (24, 44, 47–49). One report claims that Lp(a) behaves like an acute phase reactant (26). Therefore, Lp(a) concentration cannot be considered an unchangeable, genetically controlled phenotype. Apo(a) phenotypes, however, are a true polymorphism that is not changed by disease or throughout life. Relating apo(a) isoforms to disease, therefore, would definitely show that the apo(a) gene which determines apo(a) type, and influences Lp(a) level, indeed contributes to susceptibility to CHD.

Apo(a) isoforms were originally designated according to their relative mobility, compared with apo B-100 in SDS-polyacrylamide gel electrophoresis, and six isoforms were distinguished (1, 13). Meanwhile, improved separation techniques have enabled us to distinguish more than 10 apo(a) isoforms by SDS-PAGE and immunoblotting (16, 27), and more than 20 apo(a) size alleles have been demonstrated by pulsed field gel electrophoresis (20) (H. G. Kraft, S. Köchl, and G. Utermann, unpublished). These alleles and isoforms differ by the number of kringle 4 repeats in the apo(a) gene and protein, respectively. Higher resolution and better separation of isoforms does not, however, affect the linear relationship between apo(a) isoform size and Lp(a) concentration. The same inverse association between apo(a) isoform size and Lp(a) levels was obtained in studies based on 6 isoforms (13–15, 27, 50), 11 isoforms (16), 19 alleles (20), and 26 alleles (H. G. Kraft, S. Köchl, and G. Utermann, unpublished). To keep the conditions between the different population samples comparable, we used the same methodology and the same apo(a) isoform standards throughout (27). Isoforms were designated according to our original nomenclature (1, 13). Isoforms that did not comigrate exactly with one of the isoform standards were binned with the closest respective isoform in the standard (51). (The apparent molecular weight of apo(a) isoforms were as follows: F, < 400 kD; B, ~ 460 kD; S1, ~ 520 kD; S2, ~ 580 kD; S3, ~ 640 kD; S4, > 700 kD.)

The results from this study clearly show that apo(a) type frequencies are significantly different between Chinese CHD patients and controls. This is due to a higher frequency of isoforms B, S1, and S2 in the Chinese patients. These isoforms, which are associated with elevated Lp(a) levels in all Caucasian and Asian populations, were almost twice as frequent in the CHD patients, compared with the control population. Apo(a) types were also determined in a second independent set of Chinese controls ($n = 189$) and CHD patients ($n = 192$) from Singapore. (Mean age of the CHD-patients in this set was

57.7±8.5 yr). In this second and independent sample, apo(a) type frequencies were also significantly different between patients and controls (likelihood-ratio chi-square $P < 0.01$, Table V). As in the first set, it was the low relative molecular mass isoforms (B, S1, S2) associated with high Lp(a) levels that were significantly overrepresented, and almost twice as frequent in the CHD patients (Table V). No Lp(a) level information was available for this set (samples had been stored for more than 12 mo at -20°C , which prohibits reliable determination of Lp(a) concentrations but not apo(a) phenotyping with our assays). Therefore, no other data from this set were included here. The phenotyping data from this independent set does, however, strongly support our major conclusion from the first set, namely, that alleles at the apo(a) gene locus contribute to the risk for CHD.

We believe that these clear results were obtained because we selected the Chinese population from Singapore for study. Measured variability at the apo(a) gene locus (apo[a] isoforms) explains only a fraction of the inter individual variation in plasma Lp(a) levels. Plasma Lp(a) concentrations, in turn, contribute to the risk of CHD. Therefore, it will be more difficult to detect a difference in apo(a) isoform frequencies between CHD patients and controls, compared with differences in Lp(a) levels. In the Chinese controls, a large fraction of the variability in Lp(a) levels is predicted by variation at the apo(a) locus (R^2 from the analysis of variance 0.54).

Logistic regression analysis demonstrated that apo(a) types are a predictor of CHD, independent of age, sex, total cholesterol, and HDL cholesterol. The odds ratio for Chinese subjects with isoforms B, S1, and S2 suggest that their risk to suffer from CHD at a mean age of 57 yr is approximately twice that of subjects in which these isoforms are absent. It is important to note that odds ratios were higher for Lp(a) levels than for apo(a) types, and that apo(a) type was only a predictor for disease when Lp(a) levels were omitted from the logistic regression analysis. This strongly suggests that the effect of the apo(a) gene on disease is mediated by Lp(a) level, and not directly by apo(a) type. This implies that apo(a) phenotyping is not recommended for CHD risk assessment in practical medicine. Lp(a) concentrations are sufficient, and superior to isoform determination for the prediction of CHD risk. In this context, another observation is noteworthy. Lp(a) levels in CHD patients were elevated within each of the common apo(a) phenotypic classes. On the surface, this result may seem counterintuitive if one

Table V. Frequencies (%) of Subjects with Low and High *M_r* Isoforms in the Two Independent Sets of Chinese Patients with Coronary Heart Disease (CHD) and Controls from the General Population

	Set 1		Set 2	
	Controls	CHD	Controls	CHD
Isoforms	$n = 210$	$n = 162$	$n = 189$	$n = 193$
B/S1/S2*	8.5	15.9	9.5	18.7
S3/S4/0†	91.5	84.1	90.5	81.3
Likelihood ratio				
chi-square	$P < 0.01$		$P < 0.01$	

* All subjects with either of the isoforms B, S1, and S2.

† All subjects with only S3 or S4 isoforms or with Null type.

assumes that the elevated average Lp(a) level in the patients was attributable to an increased frequency of certain apo(a) isoforms. However, it does not contradict our hypothesis that differences in apo(a) isoform frequencies contribute to Lp(a) level differences, and, in fact, might be expected if one considers some features of the Lp(a)/apo(a) system. Two points are relevant to this discussion. First, the relationship between apo(a) type and Lp(a) level is not strict (1, 50). An isoform of the same size may be associated and cosegregated with high Lp(a) levels in one, and with low levels in another family (H. G. Kraft, S. Köchl, Ch. Sandholzer, and G. Utermann, unpublished); second, apo(a) alleles have an additive effect on Lp(a) levels (1, 14, 50). Thus, there may be more true homozygotes among the patients, and also more subjects with isoforms that are associated with levels in the higher range of the respective type. Both phenomena would result in an increase of the type-specific Lp(a) levels in CHD patients over controls. Thus, finding both a higher frequency of apo(a) isoforms B, S1, and S2 that are associated with high Lp(a) levels in the general population, and a higher mean concentration of Lp(a) in a given phenotype, is compatible with the postulated relationship between apo(a) isoforms, Lp(a) levels, and the risk for CHD. This does not exclude the possibility that other factors, genetic or non-genetic, that are unrelated to the apo(a) locus, might also contribute to the elevated Lp(a) levels in CHD patients. Mutations at the LDL-receptor locus that cause familial hypercholesterolemia have been shown to result in elevated Lp(a) levels (23). Other yet unidentified genes may also raise Lp(a) concentrations and increase the risk for CHD. Thus, the elevated type-specific Lp(a) levels in the CHD patients may well result from the operation of both mechanisms.

We conclude that Lp(a) may be an important predictor of CHD in Singapore Chinese, and that high Lp(a) concentrations in this population are genetically determined.

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