

eCommons@AKU

Department of Pathology and Laboratory Medicine

Medical College, Pakistan

January 1994

Pattern of dyslipoproteinemia in selected population of Karachi

Erum Khan Aga Khan University

Ayesha Molla Aga Khan University

Naila Kayani Aga Khan University

Mohammad Khurshid *Aga Khan University,* mohammad.khurshid@aku.edu

Follow this and additional works at: http://ecommons.aku.edu/ pakistan_fhs_mc_pathol_microbiol Part of the <u>Pathology Commons</u>

Recommended Citation

Khan, E., Molla, A., Kayani, N., Khurshid, M. (1994). Pattern of dyslipoproteinemia in selected population of Karachi. *Journal of Pakistan Medical Association*, 44, 165-168. **Available at:** http://ecommons.aku.edu/pakistan_fhs_mc_pathol_microbiol/582

Pattern of Dyslipoproteinemia in Selected Population of Karachi

Pages with reference to book, From 165 To 166

Erum Khan, Ayesha Molla, Naila Kayani, Mohammed Khurshid (Department of Pathology, The Aga Khan University Hospital, Karachi.)

Abstract

Pattern of dyslipoproteinemia was studied ,over a period of 3-1/2 years in 487(394 males, 93 females) cases aged 4 to above 70 years. Type IV hyperlipoproteinemia was most prevalent (36%) followed by type V (15%) (JPMA 44:165,1994).

Introduction

Lipid abnormalities are frequent both in healthy school going children and adults in Pakistan^{1,2}. Studies in neonates and their mothers indicate that diet may be an important aetiological factor for hyper cholesterolaemia in children³. A substantial number of Pakistanis get their first heart attack in the fourth decade of life³ and hypertriglyceridaemia and low LDL has been reported in hypertensive subjects⁴. Lipoprotein phenoytyping is useful in classifying lipid abnormalities and helps in assessing the probable cause of the lipid disorders⁵. The present study was undertaken to see the pattern of dyslipoproteinemia in a selected group of hyperlipidemic patients.

Patients and Methods

All the patients referred to clinical laboratory at The Aga Khan University Medical Centre, for lipoprotein electrophoresis between October, 1989 to March, 1993 were included. Patients were fasting for 12 hours. Five ml. blood was collected in plain tube and serum stored at 4°C. The paragon lipoprotein electrophoresis (LEP) kit P/N 855910 (Beck men Instruments Inc/Diagnostic System Group) was used for electrophoretic separation of lipoproteins in human serum. Phenotypes were described according to Fredrickson's classification⁶.

Results

Patients were divided into different age groups from 0 to greater than 70 years. Phenotypes were identified and mentioned against each age group. Total number along with specific type of hyperlipidemia in individual age groups were studied and their percentages were calculated (Tables I,II and III).

		-						
Age group (years)	Type I	Type II	Type III	Type IV	Type V	Normal pheno-	Total	%
						type		
0-14	0			2	1	6	9	1.81
15-29	8			9	7	6	30	6.04
30-49	45	3	4	114	45	82	293	60.0
50->70	21	2	2	50	21	60	87	31.8
Total of various types	74	5	6	175	74	154		
% of prevalence of various types	15.1	1.0	1.2	35.9	15.1	31.6	487	
Total	487							

Table I. Pattern of dyslipoproteinemia among the total patients

Table II. Pattern of dyslipoproteinemia among the males

Age group	Туре	Туре	Туре	Туре	Туре	Normal	Total	%
	I	п	III	IV	v	pheno-		
						type		
0-14				2		5	7	1.5
15-29				9	5	4	23	5.8
30-49	39	2	4	99	39	63	246	62.2
50->70	13	2	1	38	19	4	118	29.6
Total of	57	4	5	148	63	117	394	
diff. types								
Percentage	14.4	1.0	1.2	37.5	15.9	29.6		
Total	394							

Age group	Туре	Туре	Туре	Туре	Туре	Normal	Total	%
	I	II	III	IV	v	phenoty		
0-14						1	1	1.0
15-29	3				2	2	7	7.4
30-49	6	1		15	6	19	50	50.4
50->70	8		1	12	2	15	40	42.8
Total of various types	17	1	1	27	10	37	93	
% of various types	18.2	1.0	1.0	29.0	10.7	39.7		

Table III. Pattern of dyslipoproteinemia among the females.

Maximum number of patients (101) were in the age. group of 40 to 44 years which constitute 20.7% of total population studied. Type IV hyperlipidemia, defined as broad pre-beta band with moderate to markedly increased VLDL in LEP was most prevalent (35.9%) in all age groups. This included 2 of the 6 patients between the ages of 0 to 4 years. Type I hyperlipidemia, defined by the presence of an intense band at application with moderate to markedly increased chylomicrones in the LEP, is most prevalent (48.6%) in age group of 35 to 44 years. Pure hypercholesteremia type IIa, defined as beta lipoproteins in the LEP and grossly elevated with normal triglycerides and chylomicrones, mixed variety type IIb, defined as increased beta and pre-beta lipoproteins in the LEP and type III (defined as intermediate density lipoprotein fraction appearing between beta and pre-beta band cholesterol and triglycerides slightly or grossly elevated in the LEP) were found in very few patients constituting 1 to 1.2% of the total population respectively.

Discussion

The present study done on a very selected group, does not represent the entire population of Karachi. However, it provides an opportunity to see the prevalence of different patterns of dyslipoproteinemia in this population. Lipoprotein phenotyping is useful in classifying lipid abnormalities and in assessing probable cause of underlying disorders. However, a given type may be due to the presence of various genetic abnormalities³. The result of this study indicate high prevalence of hyper triglyceridemia in contrast to the previous study which showed high incidence of hypercholesterolemia⁷. However, previous studies were done on a sample of healthy population selected at random whereas in this study all the patients were referral cases suspected to have hyperlipidemia. Further workup is, hence, strongly emphasized. High incidence of moderate to severe hypertriglyceridemia (type IV, V and I) is alarming because of its association with atherosclerosis, constituting 6% of all the incidence with myocardial infarction⁸. Evidence support the view that hyper triglyceridemia is more commonly associated with obesity, uncontrolled diabetes meilitus, access of alcohol ingestion, renal failure, SLE, lipodystrophy, glycogen storage disease and use of various medications like beta-blockers and oral contraceptives⁴. The result of this study demands a more comprehensive prospective study to evaluate the involvement of secondary factors in development of dyslipoproteinemia and molecular defects (determined by using latest predieator, plasma apolipoproteins, forming the genetic basis of lipid disorders in our population.

References

1. Badnsddin, S.H., Khurshid, M. Molla, A. eta!. Factors associated with elevated serum cholesterol levels in well to do Pakistani school children. J.Trop.Med.Hyg., 1991;94: 124-29.

2. Ibrahim, K.,Zubesi,S.J. and Hussein, N. Bloodlipid patterns in Karachi. J.Pak.Med.As soc., 1981;31:164-71.

3. Badruddin, S.F., Lalani, R., Khurshid, M. et al. Serum cholesterol in neonates and their mothers. Apilotstudy. J.Psk.Med.Assuc., 1990;40:108-9

4. Bano. K.A., Jabeen, M., Din, F. and Haider, z. Bloodlipidprotile inireated anduntreated hypertensive patients. Pak.J.Med.Res., 1984;23:58-62.

5. Schaefer, E.J. and Levy, RI. Psthogenesis and management of lipoprotem disorders. N.Engl.J.Med., 1985;312:1300-10.

6. Stein, E.A. Lipids, lipoprotein and apolipoproteins. In Fundamentals of clinical chemistry.

Philadelphia, NW. Trite Ed. Saunders Co., Philadelphia, PA; 1987; pp.448-81.

7. Molla, A., Msnser, WW.T., Lalani, K. etsl. Blood lipids ins healthy Karachi population. J.Trop.Med. Flyg., 1990;93:295-99.

8. Kottke, BA., Moll, PP. and Micheles, VV. Levels of lipids, lipoprotein and apolipoprotein ins defined population. Mayo Cli. Proc., 1991;66: 1198-1208.