INCIDENCE OF ABO AND Rh BLOOD GROUPS IN MADHYA PRADESH*

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The distribution of ABO and Rh blood groups varies in different races and even in Illerent geographical locations for the same race.

Amongst Australian aborigines, Fijaians, Fillipions, Japanese and Chinese, the Rh macter does not seem to exist at all. The incidence in the rest of south-western Asia seems be related to the southern Europe and the Mediterranean basin, where the frequency of r is wally low, and that of R_1 higher than further north (4).

The incidence of Rh negative rates varies in different races and in different countries. is variation is very wide, ranging from 0% in Burmese and Indonesians to 42.25% amongst sques (6).

The distribution of Rhesus factor varies widely in different parts of India. Different were have observed an incidence varying from 0.8 to 12% (3). Khanolkar and Sanghvi lattempted to explain this obvious paradox of varying incidence in the different States by raving attention to social composition of India. Indians include Hindus, Moslems, tribes at the rest consisting of Christians, Sikhs, Buddhists, Jains, Parsees and Jews. Browman (1) ported a disproportionately low incidence of Rh-negatives in group A but the finding has at been confirmed ever since.

MATERIALS AND METHODS

A random sample of 300 healthy medical students of both sexes of age-group 17 to 24 ars was investigated. These students gave a fairly good cross-section of the State, the adission to the Medical College being open to competition on a Statewide scale.

Anti-A and anti-B sera (Slide Method) containing not less than 64 units isohaemglutinins per ml (high titre) of Haffkine Institute, Bombay, were used, for testing the assical (ABO) blood groups.

For Rh factor, 50 cases were studied with anti-Rho' (anti-CD) serum (albumin-agglutia uting) containing not less than 32 units of isohaemagglutinins (high titre) of Haffkine Instiute. Bovine albumin (22% solution, pH 7.5) of the same Institute was used.

aper read at the XII Annual Conference of the Association of Physiologists and Pharmacologists of India (1966).

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0.4 parts of blood from finger were diluted to mark 11 with isotonic saline (0.9%) in a Thoma's pipette to obtain 2% suspension of red cells. One drop of the suspension was mixed with 1 drop of anti-CD serum (incomplete or monovalent agglutinin) and 1 drop of bovine albumin on a slide, rocked and kept in a moist chamber (Patri dish with damp filter paper to lessen the evaporation and drying). The chamber was then placed in an incubator at 37° C for 30 minutes. The results were confirmed under a microscope.

The remaining 250 cases were studied with hyperimmunized anti-D (Rh_o) serum of Indian (human) origin, of high specificity, titre and avidity; it contained saline (complete or bivalent) agglutinin, and was obtained from Bharat Laboratories, Bombay. For this study, rapid slide conglutination technique was used.

In the ABO system A, B and O serve as symbols of phenotype as well as genes. To denote gene frequency, the letters p, q and r are used to indicate the frequency of genes A, B and O respectively. Genotypes can be homozygous (AA, BB, OO) or heterozygous (AO, BO).

In the Rh system, gene corresponding to antigen D is called D (Rh); when D is absent from chromosomes, its place is occupied by the alternate form (allelomotph) called d (rh). An Rh gene is inherited from both father and mother. If gene D is carried by both sperm and ovum, the resulting gene composition (genotype) of the offspring is DD; if the gametes carry D and d respectively, the result is Dd; if both gametes carry d, the result is dd.

To calculate the genotype frequency :

- 1. With two like genes the genotype frequency is the square of the gene frequency.
- 2. With two unlike genes the genotype frequency is twice the product of the two frequencies.

In Rh system: 'DD'='D'2; 'Dd'-2 ('D'×'d'); 'dd'-'d'²; where 'dd' denotes genotype frequency of dd (frequency of phenotype Rh negative).

When only anti-D serum is used the gene frequency will be : $p^{rh}(d) = \sqrt{^{6}dd}$ or \sqrt{rh} negative $qR^{h}(D) 1 - d(p^{rh})$

where p^{rh} denotes frequency of rh, and rh negative denotes frequency of Rh negative individuals, and qR^{h} denotes frequency of Rh.

In ABO systems : (when anti-A and anti-B sera are used) :

$$p_{A} = \sqrt{\mathbf{O}^{\prime} + \mathbf{A}^{\prime}} - \sqrt{\mathbf{O}^{\prime}}$$
$$q_{B} = \sqrt{\mathbf{O}^{\prime} + \mathbf{B}^{\prime}} - \sqrt{\mathbf{O}^{\prime}}$$
$$r_{O} = \sqrt{\mathbf{O}^{\prime}}$$

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where as
$$A' = p_A^2 + 2p_A r_o$$
; $B' = q_B^2 + 2q_B r_o$; $O' = r_o^2$

where p_A denotes frequency of gene A, q_B denotes frequency of gene B, and r_0 denotes frequency of gene O; and 'O', 'B' and 'A' denote the frequency of individuals (phenotype) of blood group 0, B and A respectively.

The frequency of three allelic genes must total one.

$$q_{\mathsf{A}}+q_{\mathsf{B}}+r_{\mathsf{O}}=\sqrt{\mathbf{O}^{\prime}+\mathbf{A}^{\prime}}+\sqrt{\mathbf{O}^{\prime}+\mathbf{B}^{\prime}}+\sqrt{\mathbf{O}^{\prime}}=1.$$

RESULTS

TABLE I

The Distribution of ABO Blood Groups in Medical Students (Madhya Pradesh)

Sex	di da bashiri 1915 - Jacob Sangara 1925 - Jacob Sangara	No. of Persons Tested	0	A	В	AB	L PECCO
Male		210	63	63	69	15	
Female		90	28	26	27	9	
	TOTAL	300	91	89	96	24	1. 75.6
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TABLE II

Phenotype Percentage Of ABO Blood Groups

Sex	te na emilient california (in)	0	А	В	AB
Male	Callinie	30.00	. 30.00	32.86	7.14
Female		31.11	28.89	30.00	10.00
Temare	TOTAL	30.33	29.67	32.00	8.00

TABLE III

Genotype frequency and Gene Frequency of ABO Blood Groups

Genotype Frequency				Gene Frequency			
'A *	°В'	'O'	'AB'	er.ec p	q	o r	
0.2967	0.3200	0.3033	0.0800	0.229	0.244	0.550	

-					-			
Ε.		-	*	10	-	1		
	10	15		ь.			v	
	1.2	-	-	-				

Distribution o	of Rh Factor	in Madhya Pradesh
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e logid lo (squamig) sh	Total No. of	Rh po	ositive	Rh negative		
Sex	Tested	Number	percentage	Number	percentag	
Male	212	207	97.65	5	2.35	
Female	88	83	94.32	5	5.68	
Total	300	290	96.67	10	3.33	

TABLE V

Genotype Frequecny and gene frequency of Rh Factor

	G	enotype frequency		Gene frequency		
15	DD (Rho Rho)	Dd (Rho rh)	dd (rhrh)	D (Rh)	d (rh)	
	0.670	0.288	0.033	0.818	0.182	

DISCUSSION

Tables VI to IX have been provided to compare the results of the previous workers with those of the present investigation.

IA	BLE	VI	

ABO Blood Groups Distribution in India (Phenotype Percentage)

	the state property	Pu	Punjabis		Indians	Indians	
ikina Manana ang	only solido	Talwar and Sahney (9)	Pathak (5)	Sen <i>et al</i> (7)	Shrivastava et al (8)	Bhargava an Rajani (1966)	
A	Construction of	21.27	22.3	22.22	21.49	29.67	
В		40.36	38.8	37.18	38.28	32.00	
0		28.73	30.6	33.09	33.74	30.33	
AB	0.244	6.49	9.64	8.3	7.40	8.00	

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TABLE VII

Showing Percentage of Rh Negative Individuals in Various Parts of India (After Mittal, (3))

SN	Authors	Centre	Population Surveyed	Total No.	% of Rh Negative
1	Greval and Roy Chowdhary, 1943	Calcutta	Indians	200	10.00
2	Das Gupta, 1944	Calcutta	Indians (62) and	240	10.00
3	Khanolkar and Sanghyi, 1945	Bombay	Indians	100	2.00
4	Wiener et al., 1945	New York	Indians	156	7.10
5	Greval and Roy Chowdhary, 1946	Calcutta	Indians	200	7.85
6	Ranganathan et al, 1946	Madras	South Indians	145	4.14
7	Bird, 1946	Madras	Indians	390	1.10
8	Mazumdar, 1894	Lucknow	Indians	116	5.43
9	Ranganathan et al, 1948	Madras	South Indians	294	8.50
10	Sanghyi and Khanolkar, 1949	Bombay	Koksnath Brahmans	200	3.50
10	Sunghin and Line , so the		Chandarseniys Prabhu	200	10.0
			Vadnagra Brahmans	200	12.00
			Dasasth Brahmans	200	5.00
			Marathas	200	1.50
11	Prasad et al, 1949	London	Indian Students	105	9.50
12	Venkataraman, 1950	100000	Indians	200	7.00
13	Bird et al, 1951	Poona	Indians	408	7.50
14	Rao, 1952	Madras	South Indians	132	3.79
15	Siamons et al, 1953	Madras	Chenchu South Indians	108	6.00
16	Venkatramiah and Rao, 1953	Madras	Indianss	100	8.00
17	Pathak, 1954	Amritsar	Punjabi	227	7.49
18	Mehrotra and Saksena, 1956	Agra	Indians	125	0.80
19	Roy et al, 1959	Calcutta	Bengalees	1435	5.29
20	Pathak, 1959	Amritsar	Punjabis	550	7.27
21	Talwar and Sahney, 1959;	Amritsar	Punjabis	1000	7.30
22	Sen et al, 1959	Calcutta	Bengalees	2000	3.00
23	Anklesaria and Mathur, 1961	Ahmedabad	Indians	1011	5.80
24	ICMR Surveys, 1961	Ludhiana	Indians	1909	7.43
	the state of the state of the state	Nagpur		1066	1.03
		Lucknow	33	3606	1.33
		Bombay	, main and a second of the second of the	4086	1.83
		Trivendrum	"	1797	7.54
		Calcutta I	we beginned by, one to do the second	4448	3.23
		Calcutta II	salvir somt or prives. (3).	2101	1.71
25	Talwar, 1962	Amritsar	"	300	7.00
26	Anand, 1962	Jaipur	Supported in the second of the	1000	2.80
27	Mittal, 1963	Agra	mous control of different populat	3500	1.06
28	Bhargava and Rajani, 1966	Bhopal	to that the truth of disprise conces	300	3.33

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TABLE VIII

Representative Data of Allele Frequency of ABO Blood Groups of Indians

S.	Race	Tested by	No. of	No. of Frequency of Groups							- Lale
INO.	tion)		Tested	0	A	В	AB	. р	Ч	1	PTYN
1.	Hindus	Wiener & Wex- ler (10)	1000	.313	.190	.412	.085	.149	.291	.560	1.000
2.	Sikhs	Bird et al (1951) (Quoted from Wiener Wexler)	600	-	-	(ol <u>n</u> eli activent Chana)	94	.180	.230	. 590	1.000
3.	Bengalees	Sen et al	2200	.330	.222	.371	.074	.162	.257	. 579	0.99
4. In	dians	Bhargava and Rajani (1966)	300	0.303	.297	.320	.080	.229	.244	. 550	1.02

TABLE IX

Representative Data of Genotype Frequency and Gene Frequency of Rh System Among Indians

CNIC	Race	Tested by	No. of People -	Genot	type Frequ	iency	Gene Frequency	
5.190.	(Population)		Tested	DD	Dd	dd	D	d
1.	Bengalees	Sen et al (1959)	2200	.683	.286	.030	.826	.173
2.	Indians	Mittal, 1959	3500	.804	.184	.010	.897	.103
3.	Bengalees	Roy et al 1959	3800				.942	.058
4.	Indians	Bhargava and Rajani, 1966	300	.670	.288	.033	.818	.182

Till now the incidence of Rh factor has not been reported from Madhya Pradesh. We found the percentage of Rh negative persons to be 3.33, which is within the range so far reported in India. Our results with respect to the distribution of ABO blood groups are also in agreement with those of others.

The Rh distribution is not influenced by age, sex, religion, state of pregnancy or habitat of people, whether rural or urban, (5).

Gene frequencies are found out because, first, they afford a more direct way of comparing the blood group content of different populations than do the phenotype frequencies; second, they allow us to test the truth of theories concerning manner of inheritance of the groups; knowing the gene frequency we can calculate the expected frequency of children of different groups ABO and Rh Groups in Madhya Pradesh 179

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from any type of mating; third, they show that a sample contains a reasonable distribution of the groups, thus giving us confidence in our technique.

Gene frequency analysis showed that the observed frequency of Rh positive and negative blood corresponded to the expected frequency.

SUMMARY

The incidence of ABO and Rh blood groups was determined in 300 medical students of both sexes (representing the population of Madhya Pradesh). Anti-D serum (both albuminagglutinating and saline-agglutinating) was used for Rh typing. 3.33% incidence of Rh negative persons was found in this cross-section of population. Phenotype, genotype and gene frequencies have been calculated both for ABO and Rh blood groups.

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