### JOURNAL OF OBSTETRICS AND GYNAECOLOGY OF INDIA

# THE EFFECT OF CONSANGUINEOUS MARRIAGES ON CONGENITAL MALFORMATIONS OF CHILD

M. S. SASTRY • S. N. PANCHBHAI

### SUMMARY

Eighty cases of consanguineous (30 families) and non-consanguineous (50 families) marriages were studied in and around Nagpur and full details about the nature of birth defect in children were collected. It was found that 30% children have birth defects of some type or other in consanguineous marriages whereas it was 37.13% in non-consanguineous marriages. The same way 17 families (56.67%) were affected due to consanguineous marriages but 74% families were affected in non-consanguineous marriage. The above observation indicates that consanguineous marriage is not a major factor

causing birth defects.

### INTRODUCTION

A recent survey by ministry of social welfare (Government of India) indicates that 2% of Indian population are mentally retarded or abnormal. The percentage of physically handicapped is still more. So the need of the hour is to give a serious thought to find out the reason for all types of congenital malformations and to find a remedy for the same.

In this paper emphasis is given to find out the effect of consanguineous marriages on the state of health of the offspring vis-a-vis nonconsanguineous marriages.

Dept. of Zoology, Nagpur University, Nagpur. Accepted for Publication on 23.04.1993.

# MATERIAL AND METHODS

Since consanguineous marriages are more common in Muslim community, among South Indians and Maharashtrians more concentration was given to study the state of health of children of the above community in and around Nagpur.

Different social welfare organization were approached who are doing a yeoman service for the rehabilitation of physically handicapped and mentally handicapped children. From them the parental address of the affected children were collected and were interviewed to observe whether the marriage was consanguineous or not.

For the sake of interview the questions

# THE EFFECT OF CONSANGUINEOUS MARRIAGES ON CONGENITAL

573

asked to parents were as follows. (1) Their name, age, when married. (2) No. of children male and formula	Sl. No. of families		Abnormal	Normal
<ul><li>(2) No. of children, male and female.</li><li>(3) Whether normal or abnormal.</li></ul>	C11		2	list.
(4) If abnormal (i) Type of abnormality	C12	1	4	
(ii) Sex		1		1
(iii) Age and Name	C13	2		2
(5) The most important question asked was	C14	2	1	1 '
about the relation of parents-whether con-	C15	1	1	
sanguineous or non-consnaguineous.	C16	1	1	-
million and second	C17	1	1	
The parents of such children were inter-	C18	2	-	2
viewed in their houses and school premises and	C19	1		1
hey were very co-operative in providing all	C20 C21 C22	2 3 2	1	2 2 2
the necessary details. Whenever possible with				
the kind permission of the parents, the pho-				
tograph of the children was taken.	C23	2	*	2
Printman	C24	4		4
OBSERVATIONS	C25	2	1	1
Eighty (80) families was studied out of	C26	2	1	1
which thirty (30) were consanguineous with a	C27	1	1	
total number of sixty (60) children. Fifty (50)	C28	2	1	1
families were non-consanguineous with one	C29	2		1
hundred and thirty four (134) children.	C30	2		0
The result of the observations is as follows :	Total 30	60	18	42
Summed up data of concentrational families	N1	3	0	3
Summed-up data of consanguineous families	N2	4	2	2
Sl. No. Total Abnormal Normal of families Children	N3	4	1	3
	N4	4	1	3
	2004			
C1 4 1 3	N5	3	1	2
C2 2 1 1	N6	1	-	1
C3 2 1 1	N7	2	0	2
	N8	1		1
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$		2		2
C4 2 1 1 C5 2 1 1	N9			
C4 2 1 1	N10	2		2
C4         2         1         1           C5         2         1         1           C6         1         1	N10 N11	2 2	_	2
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	N10	2	_	
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	N10 N11	2 2		2

18

Abbreviation C for consanguineous marriage. Abbreviation N for non-consanguineous marriage.

# JOURNAL OF OBSTETRICS AND GYNAECOLOGY OF INDIA

		Abnormal	
of families	Children	-0.1 a 1/2	-0-20
N15	2	0	2
N16	2	1	1
N17	. 3	1	2
N18	2	1	1
N19	2	0	2
N20	5	2	3
N21	5	2 -	3
N22	1		0
N23	1	1	0
N24	2	2	0
N25	4	2	2
N26	2	1	1
N27	1	1	0
N28	1	1	0
N29	3	1	2
N30	3	1	2
N31	4	1	3
N32	3	1	2
N33	1	1	0
N34	3	1	2
N35	2	1	1
N36	3	1	2
N37	2	1	1
N38	1	1	0
N39	2	1	1
N40	- 3	1	2
N41	2	1	1
N42	2	0	2
N43	3	1	2
N44	8	8	0
N45	2	2	0
N46	2	1	1
N47	2	1	1
N48	4	1	3
N49	9	1	8
N50	3	1 .	2
50	134	50	84

# ESULT AND DISCUSSION

Out of the Eighty (80) families studied, hirty families were consanguineous and rest fifty families were non-consanguineous.

Following is the comparative study of birth defects among consanguineous and non-consanguineous marriages.

rative Study guineous families ation I: families 30 60 Children 18 nal Ξ. 30% age = ation II : families = 30 d families 17 56.67% age of affected families = nsancuneous families ation I : families 50 bildren 134 nal 50 age of Abnormality 37.31% = vation II : families 50 d families 37 age of affected families = 74% USION m the above observations it is quite

vident that birth defects are pronounced in consanguineous as well as non-consanguine-

ous marriages, which testifies that consanguineous marriage is not a major factor causing birth defects.

Our observations are in accordance with the observations of Harrod et al. (1984), who evaluated two unrelated infants at 14 and 27 months born to non-consanguineous normal parents. Both the children had severe birth defects. Schinzel and Ltischji (1984), recorded a severely microcephalic, quadripelagic child without extracranial anomalies born to a young non-consanguineous couple. The report of Jaffe et al. in Israel in 1988 also confirms our observation. Buttins et al. in 1989, reported an apparently new MCA-MR syndrome with facial dysmorphism, microcephaly, myopia and Dandy-Walker malformation in three severally mentally ratarded sibling born to normal, non-consanguineous parents. Our observations are further confirmed by the study of Navarrete et al. 1991.

# REFERENCES

- 1. Buttins M., J. P. Fryns, H. Van Den Berghe : Clin. Genet. : 36, 451, 1989.
- Harrod M., J. M. Fridman, G. Gurratino, R. M. Pauli, L. O. Langer : Jr. Am. J. Med. Genet. : 18 (2), 311, 1984.
- Jaffe J., S. Borochowitz : Clin. Genet. : 33 (1), 33, 1988.
- Navarrete C. R., R. Penaloza, F. Salamanca : Clin. Genet. : 40 (1), 29, 1991.
- Schinzel A., M. Litschgi : J. Med. Genet. : 21 (5), 355, 1984.

Mar man fille patients and any inter any in a real patient with the second second second and any other and a second secon

#### WOTT SUBJECT TOWN

and the second s

And a second sec

and the second s

#### HOULDIN DAY PUTATE

the second secon