

## Profile of Social, Environmental and Biological Correlates in Intellectual Disability in A Resource-Poor Setting in India

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### ABSTRACT

**Background:** Intellectual disability (ID) is a major public health issue in India. Social, environmental and biological factors all contribute to the nation's high rate of ID. **Objective:** We aimed to investigate the distribution, differences and the association of social, environmental and biological factors with different types of ID in a mixed (tribal and non-tribal) population in India. **Materials and Methods:** Secondary data was collected during a community-based rehabilitation project and analyzed with descriptive statistics: Frequency, percentage and  $\chi^2$ . **Results:** Poverty, low levels of parental education and a family history of epilepsy and ID were all associated in both tribal and non-tribal populations ( $P < 0.05$ ). **Conclusion:** The outcome of this study may be helpful in planning public health initiatives that aim to reduce the burden of ID in mixed populations.

**Key words:** Cerebral palsy, determinates, down syndrome, epilepsy, family history, India, intellectual disability, tribal

### INTRODUCTION

In India, Intellectual Disability (ID) has gained more attention than all other developmental disabilities. Across the world, the prevalence of developmental disabilities has been increasing, but the US has seen the prevalence of ID decline because of advancements in public health and healthcare facilities. Among American children, the rate of ID has fallen by 1.5% over the last decade.<sup>[1]</sup> Currently, no one yet knows if the prevalence of ID is also declining in India. However,

developing nations have almost double the prevalence of ID compared to developed nations.<sup>[2]</sup> Such developed nations, like the US, can accurately estimate and predict prevalence trends and their social, environmental and biological correlates. Conversely, India is still struggling to accurately estimate ID prevalence. While estimates of ID in India vary widely, the best estimate suggests that ID affects 1 to 32 persons per 1,000 people, indicating that ID is a serious public-health concern.<sup>[3]</sup>

Several studies have documented aetiological factors of people with ID in India. Genetic disorders, malnutrition, infectious diseases, early- or late-age pregnancy and poor medical care before, during and after birth are the major contributing factors for ID.<sup>[4-7]</sup> Similarly, the major social, environmental and biological determinates of ID are poverty, poor nutrition, lack of awareness regarding preventive measures, illiteracy, poor healthcare facilities and lack of access to health care services. Overall, these factors negatively affect pre,

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peri and postnatal care of an individual with ID.<sup>[8-12]</sup> To develop a culturally sensitive and need-based public health programme, we must first better understand the how these key social, environmental and biological factors influence ID in India.<sup>[13-15]</sup>

### Objective

This study investigated the distribution, differences and association of social, environmental and biological determinates with different levels (severities) of ID.

## MATERIALS AND METHODS

In this study, we used secondary data collected in a community-based rehabilitation (CBR) project sponsored by a non-government organization, Ashagram Trust (AGT). AGT, which offers services related to disability, health and mental illness, is based in the town of Barwani in the state of Madhya Pradesh, India. The CBR was initiated in 1999 with financial help from Action Aid, India. The screening survey was conducted in 2000 using the National Institute for the Mentally Handicapped Developmental Screening Schedule (NIMH-DDS).<sup>[16-18]</sup> The screening survey was administered by the community-based rehabilitation workers under the supervision of experts in intellectual disabilities.

In total, 63,789 people (24,681 tribal and 39,108 non-tribal) were screened. Only children between the ages of 3-18 were screened. All people who the screen identified as having ID were further evaluated by two experts in intellectual disabilities (one of which was the author). The experts administered two common diagnostic tests: The Developmental Screening Test (DST) and the Vinland Social Maturity Scale (VSMS).<sup>[19]</sup> They also performed clinical observation and conducted a parental interview prior to making the diagnosis. All cases were classified on the intelligence quotient (IQ)-based The International Classification of Diseases, 10th Revision (ICD-10) criteria of intellectual disability: Borderline: IQ >70; Mild: IQ, 50-69; Moderate: IQ, 35-49; Severe: IQ, 20-34 and Profound: IQ <20. Secondary conditions were diagnosed on the basis of clinical observations and specific testing.

The determinants of ID were classified in to three categories:

1. Social: Socio-economic status, gender, parent education and population type;
2. Environmental: Family history of intellectual disability, mental illness and epilepsy and
3. Biological: Down syndrome, cerebral palsy, epilepsy, behavioural disorders, communication disorders and enuresis.

Regarding the overall population, 63% of people in the Barwani district were living under the poverty level at the time of the survey, but slightly more of the population screened by the CBR was in poverty. The CBR classified families as being either poor or not poor. If families had no source of income other than the seasonal labour, they were considered poor. Those that had brick houses, livestock and agricultural land were considered not poor.

### Statistics

Frequency and percentage of categorical variables were used to describe the data in terms of association or difference of social, environmental and biological factors with different categories of ID. The  $\chi^2$  test statistic was used to observe differences or to measure associations. We also occasionally used  $\chi^2$  with the Yates correction. The Statistical Package for Social Science (SPSS), student version, was used for statistical analyses.

## RESULTS

Contingency tables were formed with each category of ID, and then the frequency of the variable was compared within that category using the  $\chi^2$  statistic. For example, we used the  $\chi^2$  statistic to determine if, in the borderline ID group, the frequency of borderline ID differed in males and females. As shown in Table 1, the distribution of ID in each ID group (borderline, mild, etc.) did not differ between gender or population type (tribal vs. non-tribal). Of the children with ID, children with mild, moderate and severe ID more frequently had impoverished parents, whereas children with borderline and profound ID were more often associated with wealthier parents. Parental (the father's) education differed between all the ID categories. Most children had parents with little or no education. In addition, the child's family history of mental illness was equal among all categories of ID, but severe ID was more highly associated with family history of ID, while mild ID was not associated with family history of ID. History of epilepsy was associated with severe and profound forms of ID, but it was not associated with moderate and mild ID [Table 2]. In terms of secondary conditions, cerebral palsy is highly associated with severe and profound ID, but not with mild and moderate ID. Epilepsy is strongly associated with severe and profound ID, but not with mild ID. Down syndrome is associated with moderate ID; behaviour disorders are associated with all forms except severe and profound ID; and communication disorders are associated with all forms except moderate ID. Enuresis was not associated with any type of ID [Table 3].

## DISCUSSION

We found that the distribution of ID type among the

children we studied was consistent with other studies.<sup>[7]</sup> However, we did not observe a difference in prevalence between males and females, even though ID prevalence is usually higher in males.<sup>[2,20,21]</sup> This inconsistency may have arisen because a substantial percentage (39%) of the people studied were part of the tribal population. Historically, tribal communities have discriminated less on gender differences than their non-tribal counterparts. As such, in certain states in India tribal communities

have a ratio of 1,126 female children to 1,000 male children. In non-tribal communities, the sex ratio is the opposite: Some states in India have only 893 female children per 1,000 male children.<sup>[22-24]</sup>

As in other studies, we also found that low socio-economic status — in terms of poverty and low education levels — is associated with ID.<sup>[2,20,25]</sup> Previous studies have found that a family history of ID and epilepsy are

**Table 1: Socio-economic correlates of intellectual disability (ID): Gender, poverty, population type and parental education**

ID categories	Gender		$\chi^2$	P-value				
	Male/Female	Yes (number & %) / No (number & %)						
Borderline	male	2 (0.76)	136 (51.90)	0.035	0.567			
	female	3 (1.14)						
Mild	male	40 (15.26)	98 (37.40)	0.189	0.688			
	female	39 (39.88)						
Moderate	male	54 (20.61)	84 (32.06)	0.224	0.702			
	female	45 (17.1)						
Severe	male	32 (12.21)	106 (40.45)	0.242	0.667			
	female	31 (11.83)						
Profound	male	10 (3.81)	128 (40.85)	1.25	0.98			
	female	5 (1.90)						
Poverty (poor vs. not poor)								
Borderline	poor	2 (0.76)	200 (76.33)	0.122*	0.046			
	not poor	3 (1.14)						
Mild	poor	65 (24.80)	137 (52.29)	1.718	0.204			
	not poor	14 (5.35)						
Moderate	poor	78 (29.77)	124 (47.32)	0.257	0.652			
	not poor	21 (8.01)						
Severe	poor	49 (18.70)	153 (58.39)	0.14	1.00			
	not poor	15 (5.72)						
Profound	poor	8 (3.05)	194 (74.04)	5.09	0.05			
	not poor	7 (2.67)						
Type of population (tribal vs non-tribal)								
Borderline	tribal	1 (0.38)	139 (53.05)	0.093*	0.130			
	non-tribal	4 (1.52)						
Mild	tribal	42 (16.03)	98 (37.40)	0.003	1.00			
	non-tribal	37 (14.12)						
Moderate	tribal	57 (21.75)	83 (31.67)	1.096	0.309			
	non-tribal	42 (16.03)						
Severe	tribal	35 (13.35)	105 (40.07)	0.053	0.886			
	non-tribal	29 (11.06)						
Profound	tribal	5 (1.90)	135 (51.52)	2.584	0.119			
	non-tribal	10 (3.81)						
Parental education level								
	Yes/No	None	Primary	Middle	High school	Bachelor		
Borderline	yes	1 (2.3)	0 (0.0)	3 (1.14)	1 (10.38)	0 (0.0)	0.286*	0.000
	no	199 (37.78)	17 (6.48)	18 (6.87)	9 (3.43)	14 (5.34)		
Mild	yes	58 (3.05)	12 (4.58)	6 (2.29)	1 (0.28)	2 (0.76)	0.247*	0.002
	no	142 (54.1)	5 (1.90)	15 (6.72)	9 (3.43)	12 (4.58)		
Moderate	yes	79 (30.1)	4 (1.52)	8 (3.05)	0 (0.0)	8 (3.05)	0.192*	0.040
	no	121 (46.18)	13 (4.96)	13 (4.96)	10 (3.81)	6 (2.29)		
Severe	yes	53 (20.22)	1 (0.38)	4 (1.52)	5 (1.90)	1 (0.38)	0.190*	0.044
	no	147 (56.10)	16 (6.10)	17 (6.48)	5 (1.90)	13 (5.96)		
Profound	yes	9 (3.43)	0 (0.0)	0 (0.0)	3 (1.14)	3 (1.14)	0.267*	0.000
	no	191 (72.90)	17 (6.48)	21 (8.01)	11 (4.19)	11 (4.19)		

\* $\chi^2$  with Yates correction

**Table 2: Environmental correlates of intellectual disability (ID): Family history of mental illness, epilepsy and ID**

ID categories	Mental illness (Yes/No)			$\chi^2$	P-value
	Yes/No	Yes (number & %)	No (number & %)		
Borderline	yes	0 (0.0)	5 (1.90)	0.064*	0.297
	no	46 (17.55)	211 (80.53)		
Mild	yes	11 (4.41)	68 (24.95)	1.031	0.378
	no	35 (13.25)	148 (56.48)		
Moderate	yes	19 (7.25)	80 (30.53)	0.294	0.618
	no	27 (10.30)	136 (51.90)		
Severe	yes	13 (4.96)	51 (19.46)	0.444	0.571
	no	33 (12.56)	165 (62.97)		
Profound	yes	3 (1.14)	12 (4.58)	0.066	0.732
	no	43 (16.41)	204 (77.86)		
Intellectual disability (yes/No)					
Borderline	yes	0 (0.0)	5 (1.90)	0.062*	0.317
	no	43 (16.41)	214 (81.67)		
Mild	yes	3 (1.14)	76 (29.00)	13.12	0.000
	no	143 (54.58)	40 (15.26)		
Moderate	yes	18 (6.87)	81 (30.91)	0.363	0.607
	no	25 (1.21)	138 (52.67)		
Severe	yes	19 (7.25)	45 (172.13)	10.879	0.002
	no	24 (9.16)	174 (66.41)		
Profound	yes	39 (1.14)	12 (4.58)	0.024*	0.699
	no	40 (15.26)	207 (79.00)		
Epilepsy (yes/No)					
Borderline	yes	0 (0)	5 (1.90)	0.077*	0.209
	no	62 (23.66)	195 (74.42)		
Mild	yes	12 (4.58)	67 (25.57)	4.49	0.039
	no	50 (1.90)	133 (50.76)		
Moderate	yes	12 (4.58)	87 (33.20)	11.73	0.001
	no	50 (19.08)	113 (43.12)		
Severe	yes	32 (12.21)	33 (12.59)	28.77	0.000
	no	31 (11.83)	166 (63.35)		
Profound	yes	7 (2.67)	8 (3.05)	0.132*	0.031
	no	55 (20.99)	192 (0/95)		

\* $\chi^2$  with Yates correction

often associated with the ID population, supporting the aetiology of a genetic basis for ID.<sup>[26-29]</sup> Furthermore, we found that epilepsy, cerebral palsy and Down syndrome also correlate with ID, constituting a list of biological factors that influence ID. Down syndrome is one of the known genetic causes of ID, and it might be more prevalent in the Indian population because of consanguineous marriages,<sup>[30,31]</sup> which are very common among Muslims and Hindus and also in tribal communities in certain states. This kind of relationship makes offspring more likely to inherit ID, increasing the prevalence of ID in the population.<sup>[27,29,31,32]</sup>

Similarly to Down syndrome, epilepsy, cerebral palsy, behavioural problems and communication disorders strongly correlate with ID. Epilepsy and cerebral palsy

**Table 3: Biological correlates of intellectual disability (ID): Cerebral palsy, epilepsy, down syndrome, behavioural disorders, communication disorders and enuresis**

ID categories	Cerebral palsy (Yes/No)			$\chi^2$	P-value
	Yes/No	Yes (number & %)	No (number & %)		
Borderline	yes	0 (0)	5 (1.90)	0.094*	0.128
	no	82 (31.29)	175 (0.38)		
Mild	yes	1 (0.38)	78 (29.77)	47.44	0.000
	no	81 (30.91)	102 (38.93)		
Moderate	yes	14 (5.34)	85 (32.44)	21.78	0.000
	no	68 (25.95)	95 (36.25)		
Severe	yes	51 (19.40)	13 (4.96)	92.22	0.000
	no	31 (11.90)	167 (63.74)		
Profound	yes	15 (5.72)	0 (0.0)	0.343*	0.000
	no	67 (25.57)	180 (68.70)		
Epilepsy (Yes/No)					
Borderline	yes	0 (0.0)	5 (1.90)	0.077*	0.209
	no	62 (23.66)	195 (74.42)		
Mild	yes	3 (1.14)	76 (29.00)	24.71	0.000
	no	59 (22.51)	124 (47.32)		
Moderate	yes	18 (6.87)	81 (30.91)	2.64	0.134
	no	44 (16.79)	119 (45.41)		
Severe	yes	32 (12.21)	32 (12.21)	32.51	0.000
	no	30 (11.45)	168 (64.12)		
Profound	yes	9 (3.43)	6 (2.53)	0.206*	0.001
	no	53 (20.22)	194 (74.04)		
Down syndrome (Yes/No)					
Borderline	yes	0 (0.0)	5 (1.90)	0.039*	0.528
	no	19 (23.66)	238 (90.83)		
Mild	yes	5 (1.90)	74 (28.24)	0.149	0.801
	no	14 (5.34)	169 (64.50)		
Moderate	yes	12 (4.58)	87 (0.43)	5.61	0.026
	no	7 (2.67)	156 (59.54)		
Severe	yes	1 (0.38)	63 (24.04)	0.124*	0.044
	no	18 (6.87)	180 (68.70)		
Profound	yes	1 (0.38)	14 (53.43)	0.006*	0.928
	no	18 (6.87)	229 (87.40)		
Behavioural disorders (Yes/No)					
Borderline	yes	2 (0.76)	3 (1.14)	0.149*	0.015
	no	212 (80.91)	45 (17.17)		
Mild	yes	70 (26.71)	9 (3.43)	3.62	0.081
	no	144 (54.96)	39 (14.88)		
Moderate	yes	73 (27.86)	26 (9.92)	6.707	0.013
	no	141 (53.81)	22 (8.39)		
Severe	yes	56 (21.37)	8 (3.05)	1.917	0.196
	no	158 (60.30)	40 (15.26)		
Profound	yes	13 (4.96)	2 (0.76)	0.032*	0.607
	no	201 (76.71)	46 (17.55)		
Communication disorders (Yes/No)					
Borderline	yes	1 (0.38)	4 (1.52)	0.148*	0.018
	no	179 (68.32)	78 (29.77)		
Mild	yes	32 (12.21)	47 (17.93)	41.81	0.000
	no	148 (56.48)	35 (13.35)		



**Table 3: (Continued)**

ID categories	Yes/No	Cerebral palsy (yes/no)		$\chi^2$	P-value
		Yes (number & %)	No (number & %)		
Moderate	yes	68 (25.95)	31 (11.83)	0.000	1.00
	no	112 (42.74)	51 (19.48)		
Severe	yes	64 (24.42)	0 (0.0)	38.58	0.000
	no	116 (44.24)	82 (31.29)		
Profound	yes	15 (5.72)	0 (0.0)	0.164*	0.007
	no	165 (62.97)	82 (31.29)		
Enuresis (Yes/No)					
Borderline	yes	0 (0)	5 (1.90)	0.047*	0.444
	no	27 (10.30)	230 (87.78)		
Mild	yes	7 (2.67)	72 (27.48)	0.255	0.825
	no	20 (7.63)	163 (62.21)		
Moderate	yes	11 (4.19)	88 (33.58)	0.112	0.835
	no	16 (6.10)	147 (56.10)		
Severe	yes	9 (34.35)	55 (20.99)	1.29	0.247
	no	18 (6.87)	180 (68.70)		
Profound	yes	0 (0.0)	15 (5.72)	0.083*	0.176
	no	27 (10.30)	220 (83.96)		

\* $\chi^2$  with Yates correction

can both increase the severity of ID and put people at greater risk of mortality.<sup>[33-35]</sup> Within these coexisting disorders, Trisomy 21 in Down syndrome was most associated with ID; generalized tonic clonic seizures in epilepsy were most associated with ID and ataxic, then athetoid cerebral palsy were most associated with ID. The most common communication disorders seen in children with ID were delayed language, articulation and voice-related disorders. In ID, delayed language often arises because of lack of appropriate stimulation, but voice-related disorders may arise from excessive crying and vocal abuse; articulation disorders likely come from faulty learning. Regarding enuresis, bed wetting is a common behavioural problem for children with ID; it comes from skill deficits, lack of control and a response to fear and punishment. Clinical interviews revealed that several children — especially around age 10 with mild and moderate ID who had acquired control over urine in past — pathologically lost their urine control later on and were diagnosed with enuresis.

To combat its increasing rate of ID, similar public health initiatives are needed in India, at two different levels.<sup>[36-38]</sup> First, India must understand the magnitude and relationship of the social, environmental and biological determinates of ID; second, they must develop appropriate prevention and health promotion plans that are sensitive to different races, religion and geographically diverse populations.

### Strengths and limitations

This is the first study that describes the profile and

distribution of various determinates of ID in a tribal population. Logistic regression likely would have offered a better description of the data and reduced potential multi-collinearity effects.

## CONCLUSION

This study demonstrates that socioeconomic, environmental and biological factors are associated with certain categories of ID. This understanding may help professionals develop better rehabilitation plans for people with ID and help parents and communities learn about the preventive aspects of ID. The findings of this study also provide insight into the problems and associated factors that public health professionals, governments and non-government agencies face when developing a need-based public health plan.

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## REFERENCES

- Center for Disease Control and Prevention (CDC). Developmental Disabilities Increasing in US. Specific Developmental Disabilities in US Children Aged 3 to 17 Years; 2014. Available from: [http://www.cdc.gov/features/dsdev\\_disabilities/](http://www.cdc.gov/features/dsdev_disabilities/) [Last accessed on 2014 June 6].
- Maulik PK, Mascarenhas MN, Mathers CD, Dua T, Saxena S. Prevalence of intellectual disability: A meta-analysis of population-based studies. *Res Dev Disabil* 2011;32:419-36.
- Girimaji SC, Srinath S. Perspectives of intellectual disability in India: Epidemiology, policy, services for children and adults. *Curr Opin Psychiatry* 2010;23:441-6.
- Jain S, Chowdhury V, Juneja M, Kabra M, Pandey S, Singh A, et al. Intellectual disability in Indian children: Experience with a stratified approach for etiological diagnosis. *Indian Pediatr* 2013;50:1125-30.
- Aggarwal S, Bogula VR, Mandal K, Kumar R, Phadke SR. Aetiologic spectrum of mental retardation & developmental delay in India. *Indian J Med Res* 2012;136:436-44.
- Sasanfar R, Haddad SA, Tolouei A, Ghadami M, Yu D, Santangelo SL. Paternal age increases the risk for autism in an Iranian population sample. *Mol Autism* 2010;1:2.
- Bhawalkar JS, Aswar NR, Wahab SN. A study of some etiological factors and morbid conditions in mentally handicapped children. *Indian J Med Sci* 1997;51:35-40.
- Parnes P, Cameron D, Christie N, Cockburn L, Hashemi G, Yoshida K. Disability in low-income countries: Issues and implications. *Disabil Rehabil* 2009;31:1170-80.
- Emerson E. Poverty and people with intellectual disabilities. *Ment Retard Dev Disabil Res Rev* 2007;13:107-13.
- Krahn GL, Hammond L, Turner A. A cascade of disparities: Health and health care access for people with

- intellectual disabilities. *Ment Retard Dev Disabil Res Rev* 2006;12:70-82.
11. Emerson E, Graham H, Hatton C. The measurement of poverty and socio-economic position in research involving people with intellectual disability. *Int Rev Res Ment Retard* 2006;32:77-108.
  12. Durkin M. The epidemiology of developmental disabilities in low-income countries. *Ment Retard Dev Disabil Res Rev* 2002;8:206-11.
  13. Graham H. Intellectual disabilities and socioeconomic inequalities in health: An overview of research. *J Appl Res Intellect Disabil* 2005;18:101-11.
  14. Leeder SR, Dominello A. Health, equity and intellectual disability. *J Appl Res Intellect Disabil* 2005;18:97-100.
  15. Aylward GP. Environmental influences on the developmental outcome of children at risk. *Infants Young Children* 1990;2:1-10.
  16. Madhwan T, Menon DK, Kalyan M, Narayan J, Subbarao TA. Mental retardation, A manual for village rehabilitation workers, National Institute for Mentally Handicapped, Hyderabad: New Era Printpacks; 1988.
  17. Robertson JM, Hatton C, Emerson E. (2009) The Identification of Children with or at Significant Risk of Intellectual Disabilities in Low and Middle Income Countries: A Review. Center for disability Research, CeDR Research Report, 3, Lancaster University, UK, 2009. Available from: [http://eprints.lancs.ac.uk/27956/1/CeDR\\_2009-3\\_Identifying\\_Children\\_with\\_ID\\_in\\_LAMI\\_Countries.pdf](http://eprints.lancs.ac.uk/27956/1/CeDR_2009-3_Identifying_Children_with_ID_in_LAMI_Countries.pdf) [Last cited on 2014 June 6].
  18. Arya S. Screening of pre-school children for early identification of developmental disabilities in rural area. *Indian J Clin Psychol* 1991;18:65-70.
  19. Kishore MT, Basu A. Early developmental and behavioural indicators of autism. *J Indian Assoc Child Adolesc Ment Health* 2014;10:93-109.
  20. Boyle CA, Boulet S, Schieve LA, Cohen RA, Blumberg SJ, Yeargin-Allsopp M, *et al.* Trends in the prevalence of developmental disabilities in US children, 1997-2008. *Pediatrics* 2011;127:1034-42.
  21. Christianson AL, Zwane ME, Manga P, Rosen E, Venter A, Downs D, *et al.* Children with intellectual disability in rural South Africa: Prevalence and associated disability. *J Intellect Disabil Res* 2002;46:179-86.
  22. Bushra B. Backward tribal pockets show healthy sex ratio. Hyderabad: The Times of India; 2013. Available from: <http://timesofindia.indiatimes.com/city/hyderabad/Backward-tribal-pockets-show-healthy-sex-ratio/articleshow/19854175.cms> [Last cited on 2014 June 6].
  23. Balgir RS. Impact of Gender Bias on Health and Nutrition of the Tribal Women In Relation To Dynamics of Development in India. *Internet Journal of Biological Anthropology* [internet]. 2009;3: [about 6p]. Available from: <http://ispub.com/IJBA/3/1/8573> [Last cited on 2014 June 6].
  24. Indiaonlinepages.com [homepage on the internet] Population of India, Punjab's population 2014. Available from: <http://www.indiaonlinepages.com/population/punjab-population.html> [Last accessed on September 12, 2014].
  25. Chaudhari S, Otiv M, Chitale A, Hoge M, Pandit A, Mote A. Biology versus environment in low birth weight children. *Indian Pediatr* 2005;42:763-70.
  26. Jazayeri R, Saberi SH, Soleymanzadeh M. Etiological characteristics of people with intellectual disability in Iran. *Neurosciences (Riyadh)* 2010;15:258-61.
  27. Ropers HH, Hamel BC. X-linked mental retardation. *Nat Rev Genet* 2005;6:46-57.
  28. Madhavan T, Narayan J. Epilepsy and mental retardation. *Indian J Psychiatry* 1992;34:12-7.
  29. Madhavan T, Narayan J. Consanguinity and mental retardation. *J Ment Defic Res* 1991;35:133-9.
  30. Korenberg JR, Chen XN, Schipper R, Sun Z, Gonsky R, Gerwehr S, *et al.* Down syndrome phenotypes: The consequences of chromosomal imbalance. *Proc Natl Acad Sci U S A* 1994;91:4997-5001.
  31. Bittles A. Consanguinity and its relevance to clinical genetics. *Clin Genet* 2001;60:89-98.
  32. Hussain R, Bittles AH. The prevalence and demographic characteristics of consanguineous marriages in Pakistan. *J Biosoc Sci* 1998;30:261-75.
  33. Strømme P, Mangelsdorf ME, Shaw MA, Lower KM, Lewis SM, Bruyere H, *et al.* Mutations in the human ortholog of *Aristaless* cause X-linked mental retardation and epilepsy. *Nat Genet* 2002;30:441-5.
  34. Strauss D, Cable W, Shavelle R. Causes of excess mortality in cerebral Palsy. *Dev Med Child Neurol* 1999;41:580-5.
  35. Durkin MV, Kavaggia EG, Pendleton E, Neuhäuser G, Opitz JM. Analysis of etiologic factors in cerebral palsy with severe mental retardation. I. Analysis of gestational, parturitional and neonatal data. *Eur J Pediatr* 1976;123:67-81.
  36. Carmeli E, Imam B. Health promotion and disease prevention strategies in older adults with intellectual and developmental disabilities. *Front Public Health* 2014;2:31.
  37. Cooper SA, Melville C, Morrison J. People with intellectual disabilities: Their health needs differ and need to be recognised and met. *Br Med J* 2004;329:414-5.
  38. Verma IC, Bijarnia S. The burden of genetic disorders in India and a framework for community control. *Community Genet* 2002;5:192-6.

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