

A Study to Assess the Association Between Measles Antibody Titres and Nutritional Status of Paediatric Population

Vinay Kumar Yadav¹, Hemant Kumar^{2*}

¹Junior Resident, Upgraded Department of Pediatrics, Patna Medical College and Hospital Patna, Bihar, India

²Associate Professor, Upgraded Department of Pediatrics, Patna Medical College and Hospital Patna, Bihar, India

Received: 08-09-2020 / Revised: 16-11-2020 / Accepted: 24-11-2020

Abstract

Background: Measles is endemic in the Democratic Republic of the Congo (DRC), and 89–94% herd immunity is required to halt its transmission. Much of the World Health Organization African Region, including the DRC, has vaccination coverage below the 95% level required to eliminate measles, heightening concern of inadequate measles immunity. **Material and methods:** This was a cross-sectional study conducted in the Upgraded Department of Pediatric, Pmch, Patna, Bihar, India for 1 year on 1-10 year old children. Total 600 patients were include in this study for finding out the seroprevalence and anti-measles antibody levels, and studying their association with age, gender, as well as nutritional status of these children. Blood samples were tested for presence of measles specific IgG antibodies. **Results:** Majority (68.33%) of the total subjects had been vaccinated against measles. A similar trend was observed in each of the age groups. However, the relationship between age and vaccination status was not found to be statistically significant ($p=0.227$). No statistically significant difference was observed in the baseline characteristics of vaccinated and unvaccinated group except for mean weight for age Z score which was significantly lower in the unvaccinated group ($p=0.019$). % of the total subjects ≤ 5 years old had severe wasting (severe acute malnutrition), while 20.31% had moderate wasting. Severe and moderate stunting was observed in 4.5% and 25.83% of the total subjects. In children > 5 years, 20.71% had severe thinness, 12.14% had thinness and only 2 (0.71%) case was overweight. **Conclusion:** Nutritional status of children has an association with measles antibody titres as well GMT of measles specific IgG antibody, with those with better nutritional status having higher measles antibody titres.

Keywords: Measles, Vaccine, Antibody, Vaccination, Malnourished, Anthropometry.

This is an Open Access article that uses a fund-ing model which does not charge readers or their institutions for access and distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>) and the Budapest Open Access Initiative (<http://www.budapestopenaccessinitiative.org/read>), which permit unrestricted use, distribution, and reproduction in any medium, provided original work is properly credited.

Introduction

Measles is a highly contagious, primarily childhood, viral disease caused by a single stranded RNA paramyxovirus (genus Morbillivirus)[1], and humans are the natural hosts of measles virus. Eighty-nine to ninety-four per cent herd immunity is required to halt measles transmission, and in 2017, there were 173,330 reported measles cases and 109,638 estimated deaths, with a worldwide estimated measles vaccine (MV) coverage of 85%. [2,3]

*Correspondence

Dr. Hemant Kumar,

Associate Professor, Upgraded Department of Pediatrics, Patna Medical College and Hospital Patna, Bihar, India.

E-mail: hemantpmch@gmail.com

This is lower than the 95% coverage required to eliminate measles, and much of the World Health Organization (WHO) African Region, including the Democratic Republic of the Congo (DRC), has even lower coverage than this worldwide average. [4,5] Measles is endemic in the DRC. Currently, DRC gives one routine dose of measles vaccine to children nine months of age, and in outbreak settings, to children as young as six months. Although the WHO states that all countries should include a second routine dose of MV, regardless of national routine coverage level of the first dose, this recommendation has not been implemented in the DRC [4]. Because coverage achieved through healthcare is low in the DRC, attempts are made to reach missed children through Supplementary Immunization Activities (SIA), which Doshi et al. found to be associated with decreased measles incidence. [6] The ability of an infant to seroconvert is age dependent due to level and decay of maternal

antibodies and immunological development; regional differences in seroprevalence have been observed. Expectant mothers in endemic areas may be more likely to have had natural measles infection, resulting in higher measles antibody levels, and so pass on higher levels of measles antibody transplacentally to their infants, resulting in longer lasting protection than would occur in expectant mothers with vaccine-induced antibody.[7,8] Children in measles-endemic regions are also at risk of exposure to measles at an earlier age, and this must be considered when determining ideal age of vaccination.[4] Determining drivers of seropositivity and low vaccine effectiveness (VE) is complex and can depend on immunization program logistical capacity[9], vaccine potency[10] and host factors, particularly immune system robustness as a function of development (age) and nutritional status.[11,12] Measles has occurred even in well-vaccinated populations, raising questions of why adequate protection is not achieved in such groups.[13-17] While vaccination induces humoral and cellular immune responses similar to those caused by natural disease, the resulting antibody levels are lower among those with vaccine - induced versus natural immunity. As measles continues to be inadequately controlled in DRC.[18, 19] Population assessment of measles immunity is needed. Children who have measles early in life have significantly lower mean weights for age than children of the same age who do not develop measles. Despite of the prevalence of malnutrition, and its fatality, scientific research in this field is lacking.

Material and methods

This was a cross-sectional study conducted in the Upgraded Department of Pediatric, Pmch, Patna, Bihar, India for 1 year on 1-10 year old children, after taking the approval of the protocol review committee and institutional ethics committee. Total 600 patients were include in this study for finding out the seroprevalence and anti-measles antibody levels, and studying their association with age, gender, as well as nutritional status of these children. Blood samples were tested for presence of measles specific IgG antibodies.

Inclusion criteria

Children in the age group of 1 to 12 years.

Exclusion criteria

- Children were received blood or blood components within last 3 months,
- Children received corticosteroid therapy or other immunosuppressive therapy,

- HIV positive children
- Transplant recipients (bone marrow/ solid organ)
- Received of gamma globulins within last 2 months,
- Children on dialysis and are having malignancies.

Methodology

The techniques of measurement described in Cogill's (2003)[20]. Anthropometric Indicators Measurement Guide were followed to make the following measurements. Weight was measured using a portable electronic weighing scale with a weighing capacity from 1 kg to 150 kg in 100g divisions, accuracy +/- 100g. Height: was measured in centimetres to a precision of 0.1cm by a wall mounted tape measuring up to 2 meters. An infantometer was used to measure the length for children less than 2 years of age. The following indices & their z scores were calculated: Body Mass Index (BMI) = $Weight (Kg) / Height (m)^2$. Weight for age: for children less than 10 years of age by W.H.O standard growth chart and zscore was calculated. Height for age: for all children based on W.H.O standard growth chart and z score was calculated. Weight for height: for children less than 5 years based on W.H.O standard growth chart and zscore was calculated. Nutritional status of children was classified on the basis of the WHO Growth Standards, 2006 for 0-60 months; and the WHO Reference, 2007 for 5-19 years

Results

Children 5-18 Years: Overweight: >+1SD (equivalent to BMI 25 kg/m² at 18 years) Obesity: >+2SD (equivalent to BMI 30 kg/m² at 18 years). Thinness: <-2SD. Severe thinness: <-3SD.

Children 0-5 years: Moderate wasting: weight-for-length/ height Z -score -2 to -3 Severe wasting (severe acute malnutrition): weight-for-length/ height Z -score <-3. Overweight: BMI-for-age or weight-for-length/ height Z -score > 2. Obesity: BMI-for-age or weightfor-length/ height Z -score>3. Moderate stunting: length/ height for age Z -score -2 to -3. Severe stunting: length/ height for age Z -score < -3. Blood samples were collected and serums were separated by centrifugation and stored at -22 degree Celsius till the time of assay. Measles specific IgG antibodies were detected by using a commercial IgG ELISA kit (Measles Virus IgG ELISA, IBL International GMBH) in accordance with the manufacturer's instructions

Table 1: Vaccination status of children against measles

Age group (years)	Vaccinated N (%)	Unvaccinated N (%)	Total	P value
1-10	410(68.33)	190(31.67)	600	0.227

Majority (68.33%) of the total subjects had been vaccinated against measles. A similar trend was observed in each of the age groups. However, the relationship between age and vaccination status was not found to be statistically significant (p=0.227)

Table 2: Baseline characteristics of measles vaccinated and unvaccinated children

Characteristics	Vaccinated mean±SD	Unvaccinated mean±SD	P value
Age (years)	7.2±2.8	6.1±2.9	0.31
Weight (kg)	16.8±6.8	16.1±5.7	0.049
Height (cm)	109.4±21.5	106.8±20.1	0.235
BMI (kg/m ²)	15.2±2.7	14.2±2.1	0.287
Weight for age Z score(1-10 years)	-1.6±1.1	-2.3±1.2	0.019
Height for age Z score	-1.4±0.8	-1.8±1.1	0.0677
Weight for height Z score(1-5 years)	-1.6±1.6	-1.5±1.5	0.485
BMI Z Score	-1.6±3.2	-1.7±2.1	0.712

No statistically significant difference was observed in the baseline characteristics of vaccinated and unvaccinated group except for mean weight for age Z score which was significantly lower in the unvaccinated group (p =0.019).

Table 3: Nutritional status of subjects

Parameter of Nutritional status	Total N (%)	VaccinatedN (%)	Unvaccinated N (%)
Weight for age Z Score (age ≤10yrs)	<-3	110(18.33)	72(17.14)
	-2 to -3	128(21.33)	85(20.24)
	>-2	362(60.34)	255(60.71)
	Total	600	420
Weight for Height Z Score (age≤5yrs)	<-3	50(15.63)	35(15.91)
	-2 to -3	65(20.31)	45(20.45)
	>-2	205(64.06)	140(63.64)
	Total	320	220
Height for Age Z Score	<-3	27(4.5)	18(4.5)
	-2 to -3	155(25.83)	94(23.5)
	>-2	418(69.67)	288(72)
	Total	600	400
BMI for age Z score (age > 5yrs;	<-3	58(20.71)	39(18.57)
	-2 to -3	34(12.14)	24(11.42)
	>-2 to 1	186(66.42)	145(69.04)
	>1	2(0.71)	2(0.95)
	Total	280	210

15.63% of the subjects ≤5 years old had severe wasting (severe acute malnutrition), while 20.31% had moderate wasting. Severe and moderate stunting was observed in 4.5% and 25.83% of the total subjects. In children > 5 years, 20.71% had severe thinness, 12.14% had thinness and only 2 (0.71%) case was overweight.

15.91% of the vaccinated subjects ≤ 5 years old had severe wasting (severe acute malnutrition), while 20.45% had moderate wasting. Severe and moderate stunting was observed in 4.5% and 23.5% of the total subjects. In children > 5 years, 18.57% had severe thinness, 11.42% had thinness and only 2(0.95%) case was overweight.

Amongst unvaccinated subjects, 15% children ≤ 5 years old had severe wasting (severe acute malnutrition), while 20% had moderate wasting. Severe and moderate stunting was observed in 4.5% and 30.5% of the total subjects. In children > 5 years, 27.14% had severe thinness, 14.28% had thinness and no case was overweight.

Table 4: Relationship of measles antibody status with nutritional status of total subjects

Parameter of nutritional status		Antibody status			Total	P value
		Positive N (%)	Negative N (%)	Equivocal N (%)		
Weight for Age z score	<-3	50(45.45)	50(45.45)	10(9.09)	110	0.058
	-2 to-3	77(60.15)	40(31.25)	11(8.60)	128	
	>-2	245(67.68)	67(18.50)	50(13.81)	362	
Height for Age Z score	<-3	13(48.15)	11(40.74)	3(11.11)	27	0.019
	-2 to-3	83(53.55)	60(38.71)	12(7.74)	155	
	>-2	289(69.14)	102(24.40)	27(6.46)	418	
Weight for Height z score	<-3	27(54)	21(42)	2(4)	50	0.577
	-2 to-3	34(52.30)	22(33.85)	9(13.85)	65	
	>-2	105(51.22)	86(41.95)	14(6.83)	205	
BMI Z score (age \geq 5yrs)	<-3	36(62.07)	16(27.59)	6(10.34)	58	0.001
	-2 to-3	20(58.82)	12(35.29)	2(5.88)	34	
	>-2 to 1	159(85.48)	18(9.67)	9(4.84)	186	
	>1	2(100)	0	0	2	

Table 5: Nutritional status wise geometric mean titer (GMT) of measles specific igg antibody of total children

Parameter of nutritional status		GMT (mIU/mL)	P value
W/A z score	> -2SD	889	0.001
	<-2SD to-3SD	687	
	<-3SD	548	
H/A z score	-2SD	1912	0.005
	-2SD to-3SD	685	
	<-3SD	387	
W/H z score	> -2SD	605	0.499
	-2SD to-3SD	587	
	<-3SD	549	

Discussion

Measles vaccination triggers both a cellular and a humoral immune response. Following the activation of T-lymphocytes, B-cells produce measles-specific antibodies. The specific level of immunoglobulin is an indicator of the immune response. Protein energy malnutrition can be a contributory factor for immunodeficiency, thereby diminishing the immunological response to the vaccine. Though many studies reported normal antibody response to measles vaccination in malnourished children, however, specific antibody response was suppressed in severe cases of PEM.

Our study also gives conflicting results. We have taken four parameters to study nutritional status: weight for age, height for age, weight for height and BMI. A highly statistically significant relationship ($p=0.002$) was observed between BMI Z scores (in subjects aged \geq 5yrs) and seropositivity, with higher seropositivity being noted in children with higher BMI z scores. Similarly, a statistically significant relationship ($p=0.019$) was observed between height for age Z scores and seropositivity, with higher seropositivity being noted in children with higher height for age z scores. However, no significant relationship was observed between seropositivity and either weight for height z scores or weight for age z

scores. According to weight for age, in well-nourished children seropositivity was 79% with GMT 889mIU/ml, moderately malnourished (wasting) 68% with GMT 687 mIU/ml and in severely malnourished (severe wasting) 68% with GMT 548 mIU/ml. It was not found statistically significant with p value 0.05 for seropositivity but significant with p value 0.0001 for antibody levels. According to Height for age in well-nourished children seropositivity was 79% with GMT 657 mIU/ml, moderately malnourished (stunting) seropositivity was 70% with GMT 469 mIU/ml and in severely malnourished (severe stunting) seropositivity 45% with GMT 341mIU/ml. P value noted was 0.005 which is significant. In weight for length/height both seropositivity and GMT were found insignificant in well-nourished, moderately malnourished as well as severely malnourished subject (P value:0.499)

In a study in Nigeria by Ifekwunigwe et al.[21], the geometric mean titer in subjects whose nutritional status was normal (>90% of median weight for age), mildly (75 to 90%), moderately (60 to 75%), severely (<60%) malnourished were 7.5, 8.8, 7.9, and 7.9, respectively. So, malnutrition did not affect the children ability to develop adequate immune response to measles. In another study by Dao et al.[22], seroconversion was not associated with anthropometric indices. McMurray et al.[23]found that the children's nutritional status had no effect after vaccination. All the children have equal immunological response with respect to nutritional status. Mean hemagglutination-inhibition titres are slightly reduced in all nutritional groups 14 months after vaccination. Smedman et al.[24], Halsey et al.[25], Ekunwe et al.[26] found good antibody response in children which were not severely malnourished. Similarly Lyamuya et al.[27]found there were no significant differences in measles antibody levels with regard to variations in nutritional status. Our study is not only showing antibody response in moderately nourished children but also in severely nourished children. Some studies reported seroconversion rates at least as high in malnourished as in well-nourished children because it is cell mediated immunity that is suppressed not the humoralimmunity.[28,29]

Delayed antibody response to measles vaccine was seen in malnourished children[30]. Similar to our study, there was one study which demonstrated that stunting is associated with low antibody response.[31] In the same study, apart from severe stunting, severe wasting was also associated with lower antibody response, an observation which was not observed in our study. Idris et al.[32] found decreased antibody

titre in children with Kwashiorkar. Hafez et al. found decrease humoral response to measles vaccine[33]. So, it was seen that malnourished children in the community can be safely and effectively vaccinated against measles. But some studies showing good antibody response and some showing poor antibody response. The mechanisms behind the immunological response are still inadequately understood. More researches are needed in this field to come to any conclusion.

Conclusion

Nutritional status of children has an association with measles antibody titres as well GMT of measles specific IgG antibody, with those with better nutritional status having higher measles antibody titres.

References

1. De Vries R, Duprex W, De Swart R. Morbillivirus infections: an introduction. *Viruses* 2015;7(2):699–706.
2. Thompson K. Evolution and use of dynamic transmission models for measles and rubella risk and policy analysis. *Risk Anal* 2016;36(7):1383–403.
3. Dabbagh A et al. Progress toward regional measles elimination – worldwide, 2000–2017. *MMWR Morb Mortal Wkly Rep* 2018;67(47):1323–9.
4. World Health Organization. Measles vaccines: WHO position paper – April 2017. *Wkly Epidemiol Rec* 2017;17(92):205–28.
5. Centers for Disease Control and Prevention. Progress toward measles elimination-African Region, 2013–2016. *Morb Mortal Wkly Rep (MMWR)* 2017;66(17):436–43.
6. Doshi R et al. The effect of immunization on measles incidence in the Democratic Republic of Congo: results from a model of surveillance data. *Vaccine* 2015;33(48):6786–92.
7. Cutts F, Grabowsky M, Markowitz L. The effect of dose and strain of live attenuated measles vaccines on serological responses in young infants. *Biologicals* 1995;23(1):95–106.
8. Leuridan E et al. Early waning of maternal measles antibodies in era of measles elimination: longitudinal study. *Bmj* 2010;340:c1626.
9. Doshi R et al. Field evaluation of measles vaccine effectiveness among children in the Democratic Republic of Congo. *Vaccine* 2015;33(29):3407–14.
10. Adu FD et al. Low seroconversion rates to measles vaccine among children in Nigeria. *Bull World Health Organ* 1992;70(4):457–60.
11. Carney JM et al. Cell-mediated immune defects and infection: a study of malnourished hospitalized children. *Am J Dis Children* 1980;134(9):824–7.

12. Neumann CG et al. Immunologic responses in malnourished children. *Am J Clin Nutr* 1975;28(2):89–104.
13. Rosen JB et al. Outbreak of measles among persons with prior evidence of immunity, New York City, 2011. *Clin Infect Dis* 2014;58(9):1205–10.
14. Haralambieva IH et al. Variability in humoral immunity to measles vaccine: new developments. *Trends Mol Med* 2015;21(12):789–801.
15. Avramovich E et al. Measles outbreak in a highly vaccinated population – Israel, July–August 2017. *MMWR Morb Mortal Wkly Rep* 2018;67(42):1186–8.
16. Cherry J et al. Urban measles in the vaccine era: a clinical, epidemiologic, and serologic study. *J Pediatr* 1972;81(2):217–30.
17. Cherry JD, Zahn M. Clinical characteristics of measles in previously vaccinated and unvaccinated patients in California. *Clin Infect Dis* 2018;67(9):1315–9.
18. Mancini S et al. Description of a large measles epidemic in Democratic Republic of Congo, 2010–2013. *Confl Health* 2014;8(9)
19. Gerard S, Kyrousis E, Zachariah R. Measles in the Democratic Republic of Congo: an urgent wake-up call to adapt vaccination implementation strategies. *Public Health Action* 2014;4(1):6–8.
20. Cogill B. Anthropometric indicators measurement guide. Revised 2003.
21. Ifekwunigwe AE, Grasset N, Glass R, Foster S. Immune responses to measles and smallpox vaccinations in malnourished children. *The American journal of clinical nutrition*. 1980 Mar 1; 33(3):621-4.
22. Dao H, Delisle H, Fournier P. Anthropometric status, serum prealbumin level and immune response to measles vaccination in Mali children. *Journal of tropical pediatrics*. 1992 Aug 1; 38(4):179-84.
23. McMurray DN, Rey H, Casazza LJ, Watson RR. Effect of moderate malnutrition on concentrations of immunoglobulins and enzymes in tears and saliva of young Colombian children. *The American journal of clinical nutrition*. 1977 Dec 1;30(12):1944-8.
24. Smedman L, Silva MC, Gunnlaugsson G, Norrby E, Zetterstrom R. Augmented antibody response to live attenuated measles vaccine in children with *Plasmodium falciparum* parasitaemia. *Annals of tropical paediatrics*. 1986 Jun; 6(2):149-53.
25. Halsey NA, Boulos R, Mode F, Andre J, Bowman L, Yaeger RG, Toureau S, Rohde J, Boulos C. Response to measles vaccine in Haitian infants 6 to 12 months old: influence of maternal antibodies, malnutrition, and concurrent illnesses. *New England journal of medicine*. 1985 Aug 29;313(9):544-9.
26. Ekunwe EO. Malnutrition and seroconversion following measles immunization. *Journal of tropical pediatrics*. 1985 Dec 1;31(6):290-1.
27. Lyamuya EF, Matee MI, Aaby P, Scheutz F. Serum levels of measles IgG antibody activity in children under 5 years in Dar-es-Salaam, Tanzania. *Annals of Tropical Paediatrics: International Child Health*. 1999 Jun 1;19(2):175-83.
28. Bhaskaram P, Madhusudhan J, Radhakrishna KV, Reddy V. Immune response in malnourished children with measles. *Journal of tropical pediatrics*. 1986 Jun 1;32(3):123-6.
29. Kimati VP, Loreto K, Munube GM, Kimboi F. The problem of measles virus response with reference to vaccine viability, age, protein energy malnutrition and malaria in the tropics. *Journal of tropical pediatrics*. 1981;27(4):205-9.
30. Powell GM. Response to live attenuated measles vaccine in children with severe kwashiorkor. *Annals of tropical paediatrics*. 1982 Sep;2(3):143-5.
31. Waibale P, Bowlin SJ, Mortimer EA, Whalen C. The effect of human immunodeficiency virus-1 infection and stunting on measles immunoglobulin-G levels in children vaccinated against measles in Uganda. *International journal of epidemiology*. 1999 Apr 1;28(2):341-6.
32. Idris S, El Seed AM. Measles vaccination in severely malnourished Sudanese children. *Annals of tropical paediatrics*. 1983 Jun;3(2):63-7.
33. Hafez M, Aref GH, Mehareb SW, Kassem AS, El-Tahhan H, Rizk Z, Mahfouz R, Saad K. Antibody production and complement system in protein energy malnutrition. *The Journal of tropical medicine and hygiene*. 1977 Feb;80(2):36-9.

Conflict of Interest: Nil

Source of support: Nil