Research

Epidemiology of intussusception among infants in Togo, 2015-2018

Enyonam Tsolenyanu1,2,*, Komlatsè Akakpo-Numado3, Djatougbe Eliane Akolly1, Jason Mathiu Mwenda4, Jacqueline Tate6, Amevegbe Boko2, Dadja Landoh6, Komlan Gnassingne3, Yawo Atakouma1, Umesh Parashar5

1Department of Paediatrics, Medical School of Lome, Togo, West Africa, 2Ministry of Health, Togo, 3Department of Paediatrics Surgery, Medical School of Lome, Togo, West Africa, 4The World Health Organization, Regional Office for Africa, Brazzaville, Congo, 5National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, Atlanta, USA, 6The World Health Organization, Country Office, Togo

*Corresponding author
Enyonam Tsolenyanu, Department of Pediatrics, National coordinator for New Vaccines Surveillance, Lome, Togo. tsolenyanu_enyonam@yahoo.fr

Abstract

Introduction: Intussusception is the leading cause of bowel obstruction in infants and young children. We describe the epidemiology and diagnostic and treatment characteristics of intussusception among Togolese infants over a 4-year period.

Methods: We implemented active surveillance among infants younger than 1 year of age admitted with intussusception from 2015 to 2018 at Sylvanus Olympio Teaching Hospital and in 2018 at Campus Teaching Hospital. Brighton Collaboration Level 1 case definition criteria were used to confirm the diagnosis of intussusception.

Results: During four years, 41 cases of intussusception, with an annual range of 8 to 14 cases (median: 10) were reported; and the highest number of cases (89%) was enrolled at Sylvanus Olympio teaching hospital. Intussusception was uncommon in the first 2 months of life, peaked from 5 to 7 months old (63%), with male predominance (63%), and showed no significant seasonality. One third of cases (34%) were transferred to the sentinel surveillance site from another health facility; and the median delay in seeking care was 4 days (range: 0-11) with ≥ 48-hour delay in 59% of cases. Clinical symptoms, ultrasound and surgery were combined to diagnose intussusception in all the cases (100%). The treatment was exclusively surgical, and intestinal resection was common (28/41, 68%). A high case fatality rate (23%) was observed and the average length of hospital stay was 10 days (range: 1-23).

Conclusion: Active surveillance for intussusception in Togo has highlighted exclusive use of surgical therapy; often associated to an intestinal resection with a very high case fatality rate.

This articles is published as part of the supplement Intussusception in African Infants, commissioned by Supported by the Gavi Alliance through the CDC Foundation.

Guest editors:
• Jason Mathiu Mwenda (World Health Organization Office for Africa)
• Umesh Parashar (US Centers for Disease Control and Prevention)
• Jacqueline Tate (US Centers for Disease Control and Prevention)

Available online at: https://www.panafrican-med-journal.com/content/series/39/1/7/full/
Introduction

Intussusception (IS) is defined as invagination of one segment of intestine within a more distal segment [1]. It is the leading cause of intestinal obstruction in children and infants, occurring without any identifiable cause in 90% of cases [2]. Boys are often more affected than girls; and delay in treatment may result death [3]. Globally, the rate of naturally occurring IS in infants younger than 1 year of age was estimated to be 74 cases per 100,000 infants, with a peak incidence among 5–7 month old infants [4]. Data are available on IS worldwide before rotavirus vaccine introduction in immunization schedules [4–10]; but few studies were published from the African region [11–13]. The incidence rate of IS was 56 per 100,000 among infants younger than 1 year of age in South Africa. This incidence ranged from 31 per 100,000 in South Africa to 60 per 100,000 in Zambia among children younger than 2 years of age [4, 14]. In Togo, at least two studies have been published prior to rotavirus vaccine introduction; both at Sylvanus Olympio teaching hospital; but none of them was specifically about epidemiology of the disease. The first was a 6-year retrospective IS study included 37 infants aged 2–13 months; and the second was a cross-sectional prospective study over a 2-year period that included 15 young children aged 4–42 months. Boys were predominant in both studies. The average delay in seeking care was 4 days; with 66% of cases treated after >48 hours. Surgery was the treatment method in over 94% of cases; with a range of 22 to 27% case fatality rate [15, 16].

Vaccines are now available against rotavirus, the leading cause of severe diarrhea associated to death among children worldwide [17]. A previously available rotavirus vaccine was associated with IS [18, 19]; thus large clinical trials (~70,000 infants) were conducted in US, Europe, and South America for the currently available vaccines (Rotarix® and RotaTeq®). These pre-licensure data found no evidence of an association between IS and rotavirus vaccines [20, 21]. Clinical trials from two newer rotavirus vaccines (Rotavac® and Rotasil®, which were recently pre-qualified for use by the World Health Organization, were not powered to assess an association of vaccination with IS [22–24]. However, post licensure data from middle and high-income countries such as USA, Mexico, Australia and Brazil suggest a minimally increased IS risk in vaccinated infants [25–33]. Nevertheless, given the magnitude of declines in rotavirus disease and associated mortality, compared with this small increase in the risk of IS, the benefits of rotavirus vaccination outweigh the small increase risk of IS [34–36]. Further, active surveillance for IS in seven lower-income sub-Saharan countries has shown that the risk of IS after administration of monovalent human rotavirus vaccine (Rotarix) was not higher than the background risk of IS [37]. According to the recommendation from the World Health Organization [38], Togo introduced the monovalent human rotavirus vaccine – Rotarix®- in its routine immunization schedule in June 2014 (first dose at 6 weeks old and second dose at 10 weeks old). Evidence of early impact of the vaccine on diarrheal disease magnitude among Togolese children has been observed [39–41]. The current study describes the epidemiology and diagnostic and treatment characteristics of intussusception among Togolese infants.

Methods

Study design

We conducted active surveillance for IS among infants younger than 1 year of age at Sylvanus Olympio Teaching Hospital from January 2015 through December 2018 and at Campus Teaching Hospital from January to December 2018. All infants in this age group that were admitted to one of these surveillance health facilities with a diagnosis of IS during the surveillance period were enrolled. All surgeons in the surveillance health facilities received standardized training on inclusion criteria for IS cases. Case investigation forms were available at each surveillance health facility. One pediatric surgeon coordinated IS surveillance activities. Periodic visits and phone calls were made to surgeons to encourage reporting of cases and to address likely concerns. Additionally, periodic review of surgical and ultrasound registers was conducted to ensure the thoroughness of reporting.

Study sites and population under surveillance

Total population of infants younger than 1 year of age in Togo was more than 280,700; and more than 118,500 (42%) of them lived in the geographical catchment area of the two surveillance health facilities. These health facilities were the main national referral hospitals in Togo; and were both located in Lome, the capital city. They are the only health facilities with paediatric surgeons in their staff. The current IS surveillance coordinator was based at Sylvanus Olympio Teaching Hospital from January 2015 to December 2017 and in January 2018, he moved to Campus Teaching Hospital to inaugurate a paediatric surgical ward at this health facility. No health facilities in the country have the ability to perform enema contrast in children, due to the lack of adequate contrast medium. Often in case of IS, the child was first admitted to pediatric ward; and was only referred to surgery when ultrasound result was in favour of diagnosis.

Case definition

We used Brighton Collaboration Level 1 case definition criteria for the diagnosis certainty of IS. Confirmation of IS by air and/or liquid contrast enema, and/or ultrasound (with confirmed reduction on subsequent ultrasound or enema), and/or at surgery and/or autopsy is classified as Level 1 [1]. As enema contrast medium was not availability during surveillance period, included cases were those confirmed by ultrasound and/or surgical criteria.

Data collection and analysis

The main variables collected by the study were gender, dates of birth, symptom onset, admission to first health facility, admission to surveillance health facility, and disposition, diagnosis certainty definition criteria, diagnostic method(s), treatment method, and outcome at discharge. Collected data were analysed by Epi Info 7.

Disclaimer: the findings and conclusions in this paper are those of the authors and do not necessarily represent the official position of the World Health Organization or those of Centers for disease control and prevention.

Results

Summary of surveillance findings

During IS active surveillance period, 41 cases of IS were reported among infants younger than 1 year of age; 36 (88%) of them were admitted to Sylvanus Olympio National Referral and Teaching Hospital from 2015 to 2018, and 5 (12%) to Campus Teaching Hospital through 2018. A range of 8 to 14 (median: 10) IS cases were reported annually; and 63% (26/41) were boys (Figure 1). A peak of reported cases (26/41, 63%) was observed among 5–7 month old infants (Figure 2). January (19%), 8/41 and December (17%, 7/41) had the highest number of cases (Figure 3).

Description of diagnostic and treatment characteristics

The diagnostic methods were multiple: clinical symptoms (child admission for acute abdominal pain, vomiting, associated with mucous and bloody stools), ultrasound and surgery in all the cases (100%). Air or liquid enema and autopsy were not performed in any case. Thirty-four percent (14/41) of cases were transferred to the surveillance health facility from another health facility. A range of 0 to 7 (median: 3 days) interval occurred from admission to the first health facility to transfer to the surveillance health facility. The overall care seeking median delay was 4 days (range: 0–11) from symptom onset; and 59% (24/41) of the cases were admitted to the surveillance health facility only 48 hours or more after the symptom onset. All IS cases were treated by surgical operation; with an intestinal resection in 68% (28/41) of cases. Nine cases died for a case fatality rate was 22% (9/41). Data on duration of hospital stay was available in 78% (32/41) of cases, and ranged from 1 to 23 days; with 10-day average length of stay (Table 1).

Methods

Study design

We conducted active surveillance for IS among infants younger than 1 year of age at Sylvanus Olympio Teaching Hospital from January 2015 through December 2018 and at Campus Teaching Hospital from January to December 2018. All infants in this age group that were admitted to one of these surveillance health facilities with a diagnosis of IS during the surveillance period were enrolled. All surgeons in the surveillance health facilities received standardized training on inclusion criteria for IS cases. Case investigation forms were available at each surveillance health facility. One pediatric surgeon coordinated IS surveillance activities. Periodic visits and phone calls were made to surgeons to encourage reporting of cases and to address likely concerns. Additionally, periodic review of surgical and ultrasound registers was conducted to ensure the thoroughness of reporting.

Study sites and population under surveillance

Total population of infants younger than 1 year of age in Togo was more than 280,700; and more than 118,500 (42%) of them lived in the geographical catchment area of the two surveillance health facilities. These health facilities were the main national referral hospitals in Togo; and were both located in Lome, the capital city. They are the only health facilities with paediatric surgeons in their staff. The current IS surveillance coordinator was based at Sylvanus Olympio Teaching Hospital from January 2015 to December 2017 and in January 2018, he moved to Campus Teaching Hospital to inaugurate a paediatric surgical ward at this health facility. No health facilities in the country have the ability to perform enema contrast in children, due to the lack of adequate contrast medium. Often in case of IS, the child was first admitted to pediatric ward; and was only referred to surgery when ultrasound result was in favour of diagnosis.

Case definition

We used Brighton Collaboration Level 1 case definition criteria for the diagnosis certainty of IS. Confirmation of IS by air and/or liquid contrast enema, and/or ultrasound (with confirmed reduction on subsequent ultrasound or enema), and/or at surgery and/or autopsy is classified as Level 1 [1]. As enema contrast medium was not availability during surveillance period, included cases were those confirmed by ultrasound and/or surgical criteria.

Data collection and analysis

The main variables collected by the study were gender, dates of birth, symptom onset, admission to first health facility, admission to surveillance health facility, and disposition, diagnosis certainty definition criteria, diagnostic method(s), treatment method, and outcome at discharge. Collected data were analysed by Epi Info 7.

Disclaimer: the findings and conclusions in this paper are those of the authors and do not necessarily represent the official position of the World Health Organization or those of Centers for disease control and prevention.

Results

Summary of surveillance findings

During IS active surveillance period, 41 cases of IS were reported among infants younger than 1 year of age; 36 (88%) of them were admitted to Sylvanus Olympio National Referral and Teaching Hospital from 2015 to 2018, and 5 (12%) to Campus Teaching Hospital through 2018. A range of 8 to 14 (median: 10) IS cases were reported annually; and 63% (26/41) were boys (Figure 1). A peak of reported cases (26/41, 63%) was observed among 5–7 month old infants (Figure 2). January (19%), 8/41 and December (17%, 7/41) had the highest number of cases (Figure 3).

Description of diagnostic and treatment characteristics

The diagnostic methods were multiple: clinical symptoms (child admission for acute abdominal pain, vomiting, associated with mucous and bloody stools), ultrasound and surgery in all the cases (100%). Air or liquid enema and autopsy were not performed in any case. Thirty-four percent (14/41) of cases were transferred to the surveillance health facility from another health facility. A range of 0 to 7 (median: 3 days) interval occurred from admission to the first health facility to transfer to the surveillance health facility. The overall care seeking median delay was 4 days (range: 0–11) from symptom onset; and 59% (24/41) of the cases were admitted to the surveillance health facility only 48 hours or more after the symptom onset. All IS cases were treated by surgical operation; with an intestinal resection in 68% (28/41) of cases. Nine cases died for a case fatality rate was 22% (9/41). Data on duration of hospital stay was available in 78% (32/41) of cases, and ranged from 1 to 23 days; with 10-day average length of stay (Table 1).
Table 1: overview of intussusception cases among infants younger than 1 year of age at Sylvanus Olympio Teaching Hospital and Campus Teaching Hospital, 2015 - 2018, Lome, Togo

| Items                                      | N=41 | %   |
|--------------------------------------------|------|-----|
| **Surveillance sites**                     |      |     |
| Sylvanus Olympio Teaching Hospital (2015-2018) | 36   | 87.8|
| Campus Teaching Hospital (2018)            | 5    | 12.2|
| **Infant transferred from another health facility** |      |     |
| Yes                                        | 14   | 34.1|
| No                                         | 27   | 65.9|
| **Number of day(s) between admission to first health facility and admission to surveillance health facility** |      |     |
| 0-1 day                                    | 6    | 42.9|
| 2-3 days                                   | 6    | 42.9|
| 4-7 days                                   | 2    | 14.2|
| Not applicable                             | 27   | -   |
| **Number of day(s) between symptoms onset and admission to surveillance health facility** |      |     |
| 0-1 day                                    | 17   | 41.5|
| 2-3 days                                   | 12   | 29.3|
| 4-7 days                                   | 11   | 26.8|
| >7 days                                    | 1    | 2.4 |
| **Infant meet Brighton level 1 definition criteria for intussusception** |      |     |
| Yes                                        | 41   | 100.0|
| No                                         | 0    | 0.0 |
| **Diagnosis of intussusception**           |      |     |
| Clinical symptoms*, ultrasound and surgery | 41   | 100.0|
| Clinical symptom, ultrasound, enema        | 0    | -   |
| Clinical symptom, ultrasound, autopsy      | 0    | -   |
| **Treatment of intussusception**           |      |     |
| Surgery                                    | 41   | 100.0|
| Spontaneous reduction                      | 0    | 0.0 |
| **Intestinal resection**                   |      |     |
| Yes                                        | 28   | 68.3|
| No                                         | 13   | 31.7|
| **Number of days between admission and disposition from surveillance health facility** |      |     |
| 1-5 days                                   | 7    | 21.9|
| 6-10 days                                  | 13   | 40.6|
| 11-15 days                                 | 5    | 15.6|
| 16-23 days                                 | 7    | 21.9|
| Missing data                               | 9    | -   |
| **Outcome at discharge**                   |      |     |
| Discharge home                              | 31   | 77.5|
| Death                                      | 9    | 22.5|
| Missing data                               | 1    | -   |

*child admission for acute abdominal pain, vomiting, associated with mucous and bloody stools.
Discussion

Our findings compared to available previous data on IS disease in Togo, confirmed also male predominance. Data from previous studies did not allow any comparison for likely change in age distribution [15, 16]. The very small number of intussusception cases identified precluded us from evaluating the possible association between rotavirus vaccine administration and occurrence of IS. Moreover, compared to previous data in Togo, we did not observe any improvement regarding seeking health care delay, treatment method, and case fatality rate; although the problem were already raised since the last two studies published respectively in 2004 and 2012 [15, 16]. Several publications on IS disease had already mentioned this poor management of cases in developing countries, particularly in Africa. Surgery was the common treatment method in Africa and was used in over 76% of cases with highest case fatality rates up to 34% [4, 11-14, 36]. Efforts can be made to improve IS case management in Africa as it is in high-income countries. In high-income countries, seeking health care delay is shortened, non-invasive methods are used to reduce IS in most cases, and surgery is required in less than 30% of cases resulting in dramatically low case fatality rates less than 1% [4]. In some middle-income countries even if surgery is still often use, they are improving in reduction of case fatality rate [4, 6-9]. This example can also be used in Africa with strengthen more over hospitals technical capacity in intensive care before and after surgical therapy [16].

Our study has some limitations that should be noted. Due to inadequacy of technical training in regional hospitals, IS surveillance is ongoing only in national referral hospitals located at the capital city. This limitation is related to a lack of radiological technology system, especially in regional hospitals but also at national referral hospitals. Enema reduction was not performed in children, because of lack of adequate contrast medium for this age even at the national referral hospitals. Ultrasound is performed only at national referral hospitals and pediatric surgeons are not available at regional hospitals. Some infants with IS may die, especially in rural zones far from the capital city, without any identification of the cause of the death. Another limitation of the study is the small population of Togo. It does not allow an evaluation of association between IS and rotavirus vaccine at the local level.

Conclusion

This active surveillance for IS in Togo has outlined the exclusive use of surgery as IS therapy method, often associated to an intestinal resection with a very high case fatality rate. Continued enhanced surveillance is needed to identify risk factors for intestinal resection and death.

What is known about this topic

• Delay in seeking care, intestinal resection is common, and the case fatality rate is high.

What this study adds

• No significant seasonality, children in 5-7 age-group are more affected, and surgery is the only method of treatment.

Competing interests

The authors declare no competing interests.

Authors’ contributions

All authors have read and agreed to the final version of this manuscript.

Acknowledgment

The authors acknowledge participants to this study, GAVI Alliance that provided financial support to WHO/AFRO for supporting new vaccines surveillance, and the Ministry of Health and national expanded program on immunization for their guidance and leadership in implementing the surveillance in Togo.

References

1. Bines EJ, Kohl SK, Forster J, Zanardi LR, Davis RL, Hansen J et al. Acute intussusception in infants and children as an adverse event following immunization: case definition and guidelines of data collection, analysis, and presentation. Vaccine. 2004 Jan 26;22(5-6):569-74.
2. Yakan S, Caliskan C, Makay O, Deneci A-G, Korkut M-A. Intussusception in adults: Clinical characteristics, diagnosis and operative strategies. World J Gastroenterol. 2009 Apr 28;15(16):1985-9.

3. Parashar UD, Holman RC, Cummings KC, Staggs NW, Curns AT, Reingold AL, CM et al. Trends in intussusception-associated hospitalizations and deaths among US infants. Pediatrics. 2000 Dec;106(6):1413-21.

4. Jiang J, Jiang B, Parashar U, Nguyen T, Bines J, Patel MM. Childhood intussusception: a literature review. PLoS One. 2013 Jul 22;8(7):e68482.

5. Imai N, Parashar U. Assessment of post licensure safety of rotavirus vaccines, with emphasis on intussusception. J Infect Dis. 2009 Nov 1;200 Suppl 1:S282-90.

6. Muhsen K, Kasem E, Efraim S, Goren S, Dani Cohen, Ephros M. Incidence and risk factors for intussusception among children in northern Israel from 1992 to 2009: a retrospective study. BMC Pediatrics. 2014 Aug 31;14:218.

7. Singh V, Kamath V, Kumar V, Prasad R, Saluja T et al. Retrospective surveillance for intussusception in children aged less than five years at two tertiary care centers in India. Vaccine. 2014 Aug 11;32 Suppl 1:A95-8.

8. Srinivasan R, Kumar CPG, Naaraayan SA, Jehangir S, Jeromie S. Intussusception hospitalizations before rotavirus vaccine introduction: Retrospective data from two referral hospitals in Tamil Nadu, India. Vaccine. 2018 Dec 14;36(51):7820-7825.

9. Samad L, Cortina-Borja M, El Bashir H, Sutcliffe AG, Marven S, Claire E. Childhood intussusception: a literature review. PLoS One. 2013 Jul 22;8(7):e68482.

10. Fernandes EG, Leshem E, Patel M, Flannery B, Guedes Pellini AC, Torres FJ et al. Post marketing surveillance of intussusception following mass introduction of the attenuated human rotavirus vaccine in Mexico. Pediatr Infect Dis J. 2012 Jul;31(7):736-44.

11. Butterly JP, Danchin MH, Lee KJ, Carlin JB, McIntyre PB, Elliott EJ et al. Intussusception associated with rotavirus vaccine administration: post-marketing surveillance in the National Immunization Program in Australia. Vaccine. 2011 Apr 5;29(16):3061-6.

12. Haper B, Patel M, Izuiriti HS, James Baggs, Paul Gargiullo, Eric Weintraub et al. Post licensure monitoring of intussusception after RotaTeq vaccination in the United States, February 1, 2006, to September 25, 2007. Pediatrics. 2008 Jun;121(6):1206-12.

13. Belongia EA, Irving SA, Shui IM, Kullendorf D, Lewis E, Yin R et al. Real-time surveillance to assess risk of intussusception and other adverse events after rotavirus, bovine-derived rotavirus vaccine. Pediatr Infect Dis J. 2010 Jan;29(1):1-5.

14. Shui IM, Baggs J, Patel M, Parashar UD, Rett M, Belongia EA et al. Risk of intussusception following administration of a pentavalent rotavirus vaccine in US infants. JAMA. 2012 Feb 8;307(6):598-604.

15. Weintraub ES, Baggs J, Duffy J, Vellozzi C, Edward Belongia A, Stephanie Irving et al. Risk of intussusception after monovalent rotavirus vaccination. N Engl J Med. 2013 Feb 28;368(9):821-30.

16. Yih WK, Liu TA, Kullendorf D, Belongia EA, Irving S, Klein NP et al. Intussusception risk after rotavirus vaccination in U.S. infants. N Engl J Med. 2014 Feb 6;370(6):513-9.

17. Gnanasinghe K, Abarchi H, Akapo-Numado G. Problems posed by les invaginations intesnales aiguës du nourrisson au CHU-Tokin de Lome à propos de 37 cas. J Afr Chir Digest. 2004;4(1):33-7.

18. Boume MA, Akapo-Numado GK, Gnanasinghe K. Prise en charge and pronostic des invaginations intestinales aigus de l’enfant au CHU-Tokin de Lome (Togo). J Afr Chir Digest. 2012; 12(1):1274-80.

19. Tate JE, Parashar UD. Rotavirus vaccines in routine use. Clin Infect Dis. 2014 Nov 1;59(9):1291-301.

20. Murphy TV, Gargiullo PM, Lieu TA, Kulldorff M, Belongia EA, Irving S, Klein NP et al. Safety of a heat-stable antigen vaccine in the Expanded Programme of Immunization. Journal of Tropical Pediatrics. 2016;62(1):8.

21. Gnanasinghe K, Abarchi H, Akapo-Numado Gk. Les invaginations intesnales aiguës du nourrisson au CHU-Tokin de Lome à propos de 37 cas. J Afr Chir Digest. 2004;4(1):33-7.

22. Bhandari N, Rongsean-Chandola T, Bavadkar A, Jacob John, Kalpana Antony, Sunita Tanjane et al. Efficacy of a monovalent human-bovine (116E) rotavirus vaccine in Indian infants: a randomised, double-blind, placebo-controlled trial. Lancet. 2014 Jun 21;383(9935):2135-43 Epub 2014 Mar 12.

23. Vasikari T, Matson OD, Denney P, et al. Safety and efficacy of a pentavalent human-bovine (WC3) reassortant rotavirus vaccine. N Engl J Med. 2006 Jan 5;354(1):23-33.

24. Ruiz-Palacios GM, Pérez-Schael I, Velázquez FR. Safety and efficacy of an attenuated oral rotavirus vaccine against severe rotavirus gastroenteritis. N Engl J Med. 2006 Jan 5;354(1):11-22.

25. Bhandari N, Rongsean-Chandola T, Bavadkar A, Jacob John, Kalpana Antony, Sunita Tanjane et al. Efficacy of a monovalent human-bovine (116E) rotavirus vaccine in Indian infants: a randomised, double-blind, placebo-controlled trial. Lancet. 2014 Jun 21;383(9935):2135-43 Epub 2014 Mar 12.

26. Vasikari T, Matson OD, Denney P, et al. Safety and efficacy of a pentavalent human-bovine (WC3) reassortant rotavirus vaccine. N Engl J Med. 2006 Jan 5;354(1):23-33.

27. Ruiz-Palacios GM, Pérez-Schael I, Velázquez FR. Safety and efficacy of an attenuated oral rotavirus vaccine against severe rotavirus gastroenteritis. N Engl J Med. 2006 Jan 5;354(1):11-22.

28. Bhandari N, Rongsean-Chandola T, Bavadkar A, Jacob John, Kalpana Antony, Sunita Tanjane et al. Efficacy of a monovalent human-bovine (116E) rotavirus vaccine in Indian infants: a randomised, double-blind, placebo-controlled trial. Lancet. 2014 Jun 21;383(9935):2135-43 Epub 2014 Mar 12.

29. Vasikari T, Matson OD, Denney P, et al. Safety and efficacy of a pentavalent human-bovine (WC3) reassortant rotavirus vaccine. N Engl J Med. 2006 Jan 5;354(1):23-33.

30. Ruiz-Palacios GM, Pérez-Schael I, Velázquez FR. Safety and efficacy of an attenuated oral rotavirus vaccine against severe rotavirus gastroenteritis. N Engl J Med. 2006 Jan 5;354(1):11-22.