



CODEN [USA]: IAJPB

ISSN : 2349-7750

INDO AMERICAN JOURNAL OF PHARMACEUTICAL SCIENCES

SJIF Impact Factor: 7.187

Available online at: <http://www.iajps.com>

Research Article

THE USE OF MATERNAL DIARRHEA AND ANTIBIOTICS IS ASSOCIATED WITH AN ENHANCED RISK OF DIARRHEA IN NON-INFECTED AND HIV-EXPOSED INFANTS IN PAKISTAN

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Article Received: October 2020

Accepted: November 2020

Published: December 2020

Abstract:

Young people who are uninfected and uninsured by HIV are a developing population at particular risk of infection transmission, where preventing intestinal laxity can reduce under-five mortality in sub-Saharan Africa. A remarkable companion (1999-2002) of Pakistani HEU babies followed from birth to one year of age was used. Maternal and neonatal morbidity was observed during monthly visits to centers and unscheduled visits of weakened children. Our current research was conducted at Jinnah Hospital, Lahore from February 2019 to January 2020. The Andersen-Gill Cox model was used to assess maternal, ecological and infant matches for soft stools, moderate to severe stools (MSD, stools with parchedness, stool softness or corresponding clinical affirmation) and delayed/diligent stools (> 7 days) in newborns. Non-HIV-infected newborns (n = 373) experienced an average of 2.08 (96% CI: 1.93, 2.25) scenes of diarrhea, 0.48 (96% CI: 0.41, 0.56) scenes of MSD and 0.36 (96% CI: 0.28, 0.44) scenes of delayed/constant bowel relaxation during their first year. Maternal baby blues is related to the increased danger of newborn intestinal relaxation (hazard ratio [HR]: 2.07; 96% CI: 1.45, 5.07) and MSDs (HR: 3.87; 96% CI: 1.12, 5.58). The use of maternal anti-infective was a risk factor for delayed/impossible bowel relaxation (HR: 1.64; 96% CI: 1.05, 3.56). Newborns living in families with pit toilets were 1.46 (96% CI: 1.18, 1.76) and 1.47 (98% CI: 1.06, 2.15) times more likely to experience bowel relaxation and MSDs, separately, than those with flush toilets. Current selective breastfeeding was defensive against MSDs (HR: 0.31; 96% CI: 0.16, 0.59), compared to newborns who did not receive breast milk. Decreased maternal bowel relaxation may cause a generous decrease in diarrheal moroseness in young children with HEU, expanding towards normal bowel relaxation, which anticipates mediation.

Keywords: Maternal Diarrhea, Antibiotics, Enhanced Risk of Diarrhea, Non-Infected and HIV-Exposed Infants.**Corresponding author:**

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Please cite this article in press Ghumza Farooq et al, *The Use Of Maternal Diarrhea And Antibiotics Is Associated With An Enhanced Risk Of Diarrhea In Non-Infected And Hiv-Exposed Infants In Pakistan*, Indo Am. J. P. Sci, 2020; 07(12).

INTRODUCTION:

Soft stools remain a major cause of fouling and mortality among young people living in sub-Saharan Africa (SSA) [1], accounting for nearly 12% of under-five passages in the region. Mounting evidence indicates that bowel loosening, particularly moderate to severe, and delayed or neglected bowel movements are linked to long-term fatigue, including developmental problems, increased recurrence of other diseases, and poor intellectual development [2]. In 2018, there were more than 2,500,500 young women (15-24 years) in sub-Saharan Africa living with HIV [3]. Large-scale implementation of projects to prevent mother-to-child transmission of HIV (PMTCT) has effectively reduced the danger of HIV transmission to children, causing the development of a population of children exposed to HIV but not infected (HEU). Being conceived or living with an HIV-infected mother can introduce new risk factors for bowel problems, such as regular episodes of bowel problems in the mother and increased use of anti-infection by the maternal family [4]. With 1.5 million newborns with HEU born each year, a decrease in defecation in this particularly small population would add to the global decrease in the burden of defecation. We determined the rate and risk factors for defecation, MSDs, and prolonged/uninterrupted defecation in a partner of babies with HEU [5].

This survey involved a remarkable companion of HIV-infected mothers and their newborns. The parent study recruited incubating HIV-positive pregnant women ³ 32 weeks who were hospitalized at Pakistan National Hospital in Lahore from 2019 to 2020. Our current research was conducted at Jinnah Hospital, Lahore from February 2019 to January 2020. Further information on this partner has already been described. During pregnancy, women were instructed on the care of the baby and the danger of HIV transmission through breast milk before choosing to breastfeed or giving the recipe. Mothers who decided to breastfeed were instructed to breastfeed their babies only for the first 5-8 months. In accordance with public rules in effect at that time, members received short-term treatment with zidovudine for HIV PMTCT and women with severe immunosuppression (CD4 count < 200 cells/ μ L) received cotrimoxazole prophylaxis and were informed about HIV treatment programs. Newborns did not receive additional prophylaxis and mothers did not receive antiretroviral treatment while breastfeeding. Qualification models used for our survey included the birth of a single child or first twin and, in any case, a negative HIV polymerase chain reaction (PCR) test at a subsequent visit with a recorded report. Babies who tested positive for HIV before the age of several months were considered to have perinatal disease and were not included in the survey.

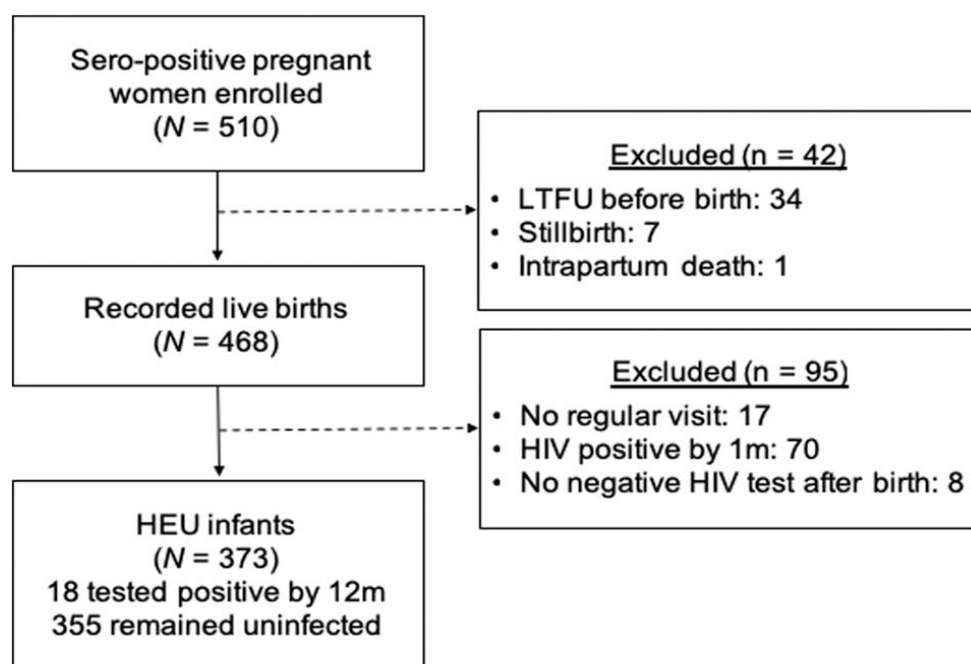
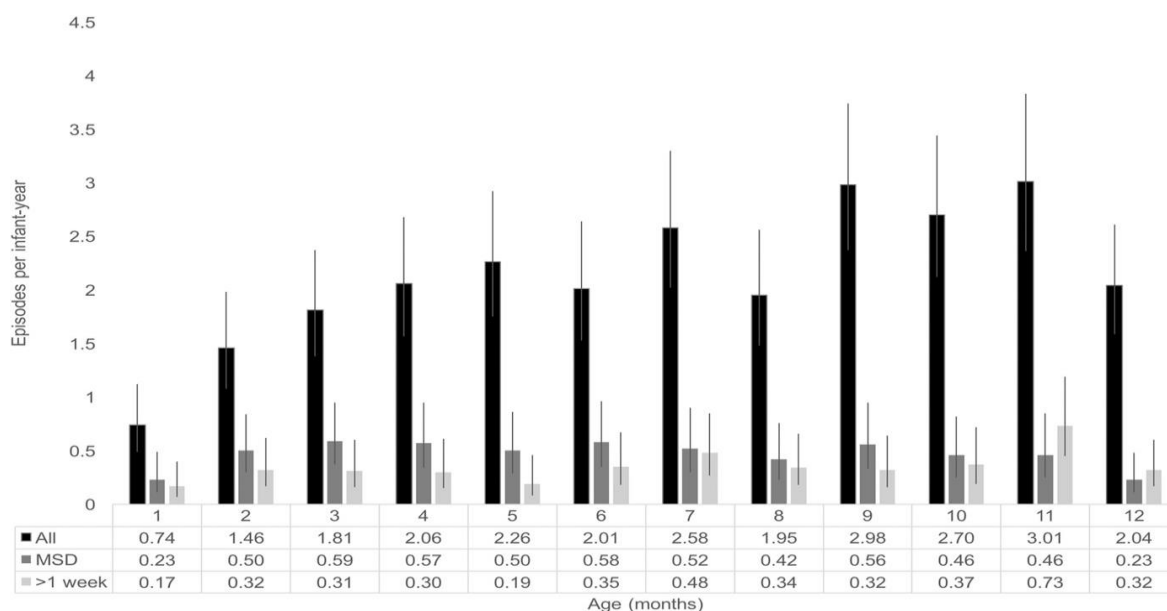
METHODOLOGY:**Figure 1:**

Figure 2:



RESULTS:

The accompanying parent enlisted 510 HIV-infected pregnant women and registered 469 live births of singleton twins or firstborns (six second-born twins were banned). Of these 469 babies, 73 were prenatally infected with HIV, 17 had died before the main visit, and eight had not been registered as HIV-negative at a normal visit (Figure 1). Mothers of infants whose development was interrupted before the main visit had a slightly higher mean log VL (method contrast = 0.43 log₁₀ duplicate/mL; 96% CI: 0.07, 0.79), but were not extraordinary in practice for some other attributes at the baseline. The additional 377 HEU infants were selected for this examination, of whom 359 were not infected during the main year and 19 therefore contracted HIV at a mean age of 181 days (interquartile range [IQR]: 90-358; Figure 1), whose visits after the last negative HIV test were recorded with a blue pencil. At enrollment, most mother-child

sets (86%) reported living in overcrowded families and using pit toilets (53%). The average age of mothers at the time of enrolment was 25 years (IQR: 22-28), and 42 percent were over school age. Previously, the mothers' average CD4 count was 448 cells/ μ L (IQR: 318-624) and the average HIV rate was 5.9 log₁₀ double/mL (IQR: 4.2-6.2); in addition, 16% of the mothers were considered undernourished (MUAC < 23.5 cm). During pregnancy, 6% of mothers revealed at least one running scene and 20% reported using antitoxins. Baby blues, 23% of the mothers already detailed the runs, of which 127 complete scenes were recorded (rate: 0.43 scenes/year). Most women (79%) detailed the use of antimicrobials for treatment in all cases once for a total of 818 events, and 14% of women reported the use of cotrimoxazole for prophylaxis, showing extreme immunosuppression (Table 1).

Table 1:

| Variable | Received multivitamins | | | Received vitamin A | | |
|---|------------------------|---------------|---------|--------------------|---------------|---------|
| | Yes (n=450) | No (n=424) | p-value | Yes (n=443) | No (n=431) | p-value |
| Mothers | | | | | | |
| Age (years) | 24.7 (4.6) | 24.7 (4.8) | NS | 24.6 (4.7) | 24.8 (4.7) | NS |
| Education level (n[%]) | | | NS | | | NS |
| None | 33 (7.3) | 30 (7.1) | | 28 (6.3) | 35 (8.1) | |
| Primary education, 1–4 years | 21 (4.7) | 20 (4.7) | | 16 (3.6) | 25 (5.8) | |
| Primary education, 5–8 years | 344 (76.4) | 339 (80.0) | | 348 (78.6) | 335 (77.7) | |
| Secondary education, 8 years | 52 (11.6) | 35 (8.3) | | 51 (11.5) | 36 (8.4) | |
| CD4 ⁺ cell count (n[%]) ^b | | | NS | | | NS |
| 0–199 cells/ μ l | 50 (11.9) | 52 (12.9) | | 50 (12.0) | 52 (12.8) | |
| 200–499 cells/ μ l | 241 (57.1) | 224 (55.7) | | 241 (57.7) | 224 (55.2) | |
| \geq 500 cells/ μ l | 131 (31.4) | 126 (31.3) | | 127 (30.4) | 130 (32.0) | |
| CD4 ⁺ cell count (cells/ μ l) | 420.4 (196.4) | 419.4 (206.1) | NS | 413.3 (196.5) | 426.8 (205.7) | NS |
| CD8 ⁺ cell count (cells/ μ l) | 731.7 (319.6) | 741.5 (309.2) | NS | 734.0 (310.0) | 739.1 (319.3) | NS |
| ESR (mm/h) | 57.3 (34.7) | 59.7 (36.5) | NS | 57.1 (34.7) | 59.8 (36.5) | NS |
| Haemoglobin (g/dl) | 9.4 (1.6) | 9.6 (1.7) | NS | 9.4 (1.7) | 9.5 (1.7) | NS |
| Vitamin A (μ g/dl) | 24.1 (8.8) | 25.2 (10.9) | NS | 24.5 (9.1) | 24.7 (10.5) | NS |
| Vitamin E (μ mol/dl) | 9.8 (2.8) | 9.7 (2.8) | NS | 9.9 (2.7) | 9.6 (2.9) | NS |
| Children | | | | | | |
| Male sex (n[%]) ^b | 218 (48.8) | 222 (52.7) | NS | 220 (50.1) | 220 (51.3) | NS |
| Birth weight (g) | 3080 (471) | 2970 (518) | 0.001 | 3027 (505) | 3025 (498) | NS |
| Birthweight <2500 g (n[%]) ^b | 30 (7.5) | 54 (14.2) | 0.002 | 42 (10.6) | 42 (11.0) | NS |
| Gestational age <37 weeks (n[%]) ^b | 104 (23.2) | 108 (25.5) | NS | 111 (25.1) | 101 (23.4) | NS |
| HIV+ at 6 weeks of age (n[%]) ^b | 69 (15.3) | 51 (12.0) | NS | 71 (16.0) | 49 (11.4) | 0.05 |

NS: not significant.

^a Data are the mean (SD) unless otherwise stated.^b Data were not available for some mothers/children for these variables, therefore the denominators are not the same as the column totals.**Table 2:**

| Cell population | HIV RNA $\geq 5 \log_{10}$ (n = 6) [median (IQR)] | HIV RNA < 5 \log_{10} (n = 57) [median (IQR)] | p-value* |
|-------------------------|---|---|----------|
| CD8 T-cells | | | |
| counts (cells/ μ l) | 1585 (1080–1702) | 1112 (71–1447) | 0.14 |
| activated (%) | 25.8 (11.2–40.3) | 8.8 (3.3–18.3) | 0.09 |
| naive (%) | 54.3 (42.8–67.8) | 79.8 (67.6–88.2) | 0.04 |
| memory (%) | 54.6 (30.5–75.8) | 21.9 (14.3–40.3) | 0.014 |
| CD4 T-cells | | | |
| counts (cells/ μ l) | 2594 (2174–3185) | 2291 (1955–3028) | 0.337 |
| activated (%) | 5.3 (4.0–7.8) | 3.5 (2.1–6.5) | 0.557 |
| naive (%) | 70.1 (60.4–74.8) | 70.3 (60.6–78.9) | 0.79 |
| memory (%) | 19.8 (15.4–21.7) | 23.9 (13.2–45.8) | 0.574 |

Analysis is based on n = 63 infants with maternal HIV RNA measures available.

*P-value from kruskal-wallis test.

DISCUSSION:

In this partner of HEU babies, runs, including MSD and loose stools, occurred regularly during the first year of life [6]. Hazard factors fluctuated according to the type of soft stool, with diarrhea primarily related to the likelihood of irresistible introduction (type of latrine, swarming and maternal diarrhea), while risk

factors for MSDs included the likelihood of irresistible presentation (type of latrine and maternal diarrhea) and factors related to the child's ability to fight off contamination (high maternal leukemia and breastfeeding) [7]. Delayed/diligent loose bowel was linked to low financial status (maternal education); also, maternal use of antimicrobials, a possible marker

of the mother's general well-being. The rate of excretion of this HEU associate was slightly lower than the 3.6 scenes/day-year detailed for youth in Nairobi during this period [8]. The lower rates of liquid stool in newborns in our review, compared to other assessments distributed simultaneously, may be influenced by the finding of liquid stool during monthly facility visits, as opposed to the more regular home visits used in different surveys. Because of the longer time periods between the different pieces of information, mothers may have simply detailed more extreme episodes of loose stool [9]. Our review found a higher rate of bowel relaxation in babies aged about a few months, consistent with the maximum frequency (6-9 months) previously detailed in the HEU and other cohorts. Our accomplice had comparable delayed/industrialized rates of intestinal laxity; furthermore, MSDs were studied late, with a rate of onset of MSDs almost indistinguishable from that reported in western Pakistan, a place where maternal HIV is prevalent, at 0.56 scenes/year [10].

CONCLUSION:

In summary, we found that both family and natural variables anticipated both intestinal laxity and MSD, while selective breastfeeding; furthermore, a low maternal LV was remarkably defensive for MSD, and maternal training and the baby blues that the mother uses as an anti-toxin seemed significant for delayed/stunted loose bowel. With variable hazard factors between all intestinal slackening and this is just the tip of the iceberg, severe types of intestinal slackening in babies may have particular components for these diarrheal conditions. Recognizable evidence of the mother's involvement in diarrhea and the use of an anti-infective may provide an opportunity to recognize high-risk babies and intercede to prevent or, conversely, treat babies with diarrhea. Direct transmission of mediation to HIV-infected parent figures during standard site visits, including instruction on indications of extreme illness in young people, empowerment of care-seeking behaviors, breastfeeding support, and oral rehydration salts and zinc, can cause a significant decrease in morbidity and mortality of newborns with diarrheal disease.

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