Assessment of the effectiveness of PMTCT program in eight service delivery points in North Central Nigeria

Obinna Ositadimma Oleribe
Ede Enenche
Deborah Udofia
Ekei Ekom
Princess Ifunanya Osita-Oleribe
Jin Un Kim
Simon David Taylor-Robinson

1Excellence and Friends Management Care Centre (EFMC); Abuja, Nigeria; 2Hepatology Unit, Imperial College London, London, UK

Background: Mother-to-child transmission (MTCT) of HIV is one of the commonest avenues through which infants are infected with HIV. To achieve an HIV-free generation, MTCT of HIV should be eliminated. Nigeria began prevention of mother-to-child transmission (PMTCT) services 13 years ago, but it still contributes to over one-third of global MTCT burden. We set out to explore and define the effectiveness of PMTCT in selected sites in North Central Nigeria.

Methods: We conducted a retrospective secondary data analysis at eight service delivery points in two states. One thousand four hundred and fifty-four mother–infant pair data sets from 2012 to 2016 were extracted and analyzed. Maternal/infant antiretroviral (ART) services, early infant diagnosis (EID), and final outcomes were reviewed to examine the predictors of MTCT of HIV in these centers.

Results: We retrieved 1,454 mother–infant pair data sets. While 89.5% (1,302) of positive pregnant women (PPW) and 92.2% (1,340) of HIV-exposed infants (HEIs) received ART prophylaxis/ARV treatment (ART), 88.4% (1,285) infants were breastfed with 32.5% still receiving breast milk at the time of dry blood spot (DBS) collection. EID PCR positivity rate was 3.5% (range, 0.0%–11.1%). Facility of delivery ($\chi^2=24.99, P<0.00$), mother on ART ($\chi^2=48.8, P<0.00$), mother having received ART prophylaxis ($\chi^2=89.59, P<0.00$), infant having received ART prophylaxis ($\chi^2=58.56, P<0.00$), and baby having received cotrimoxazole ($\chi^2=55.24, P<0.00$) all significantly prevented positive EID results. However, mode of delivery and breastfeeding were not significantly associated with positive EID results.

Conclusion: This study supports PMTCT services as it minimizes the transfer of HIV from infected mothers to HEIs. To eliminate HIV and achieve zero new HIV infections, every HIV-positive pregnant woman should receive ART prophylaxis and should be supported postdelivery to prevent transfer of infection to the newborn. Also, HEIs should receive timely ART and cotrimoxazole prophylaxis.

Keywords: antiretroviral therapy, infectious diseases, prevention of mother-to-child transmission, women, Africa

Background
HIV has remained a major public health challenge, with ~36.7 (range, 34.0–39.8) million people living with HIV, 1.8 (range, 1.6–2.1) million new infections, and 1 million mortalities by the end of 2016.¹ To date, over 35 million lives have been lost to HIV and associated infections globally, with sub-Saharan Africa remaining the most affected region, as it accounts for two-thirds of the global total of new HIV infections and with 36.7 million people living with HIV in 2016.¹ In Nigeria, over 3.1 million people are living with HIV, and the current national HIV prevalence...
is 3.0%, according to the recent national sentinel studies, among pregnant women attending antenatal care.²

HIV-positive woman can infect their babies during pregnancy, childbirth, and/or breastfeeding, and this accounts for >90% of new HIV infections among children.³ For instance, in the absence of any interventions during these stages, infected mothers can transmit the virus to their babies in 15%–45% of cases.⁴ In 2015, there were about 1.8 million children (age, 0–14 years) infected with HIV with 490,000 in West and Central Africa and 260,000 in Nigeria alone.⁵,⁶

Interventions to reduce mother-to-child transmission (MTCT) primarily involve antiretroviral (ARV) treatment (ART) for the mother and a short course of ARV drugs for the baby, measures to prevent HIV acquisition in the pregnant woman, and appropriate breastfeeding practices.⁴ Effective prevention of mother-to-child transmission (PMTCT) services require women–infant pair to have access to all relevant interventions.⁷ In September 2015, the WHO released a new guideline that recommended lifelong ART for all pregnant and breastfeeding women living with HIV, commonly called Option B⁴.⁸,⁹

PMTCT of HIV program began in Nigeria in December 2000 with the inauguration of the PMTCT National Task Team (NTT), while actual PMTCT services commenced as a pilot project in July 2002. This is in line with the WHO four-pronged approach.¹⁰ However, since inception, no structured evaluation of the effectiveness of the PMTCT services in Nigeria has been undertaken. Therefore, there is a paucity of data on PMTCT effectiveness of PMTCT programs.¹¹ The purpose of this study was to determine the effectiveness of PMTCT program in selected sites in Nigeria.

Methods

We purposefully selected eight supported sites in two states (Nasarawa and Abuja, Federal Capital Territory [FCT]; Figure 1). This study was conducted in sites supported by Excellence and Friends Management Care Center (EFMC), a non-governmental organization involved in comprehensive HIV health care delivery across a wide range of Nigerian states.

Mothers whose infant details were documented in the registers sufficient for analysis and mothers in care for >6 months were included in this study.

Figure 1 Map of Nigeria showing study locations – Nasarawa and Abuja, FCT.

Abbreviation: FCT, Federal Capital Territory.
We designed a data extraction form (DEF) using Microsoft Excel spreadsheet to extract and record data. Ethical approval was obtained from the Nigerian Institute of Medical Research (IRB/16/354). All data collected were anonymized to remove identifiers and analyzed in aggregates; therefore, patient consent was not required. The training day included review of the data extraction template, data collection processes, use of Excel sheet, and review of entries. Data collection took place between December 2016 and January 2017. One thousand four hundred and fifty-four data sets were extracted from the early infant diagnosis (EID) PCR request and result forms and the child follow-up register across the eight facilities in the two states. Additional information was obtained from Delivery and Maternal registers. The chi-squared test was utilized to test for significance, and data were validated and analyzed using MS Excel, SPSS version 23, and OpenEpi.12,13

Results
A total of 1,454 mother–infant data sets were extracted from the eight health care facilities (seven public and one private) located in Abuja, FCT (7; 87.5%) and Nasarawa (n=1, 12.5%). There were four primary and four secondary level facilities (Table 1), but 1,207 (82.8%) of the extracted data were from secondary level of care.

There were 50.8% female infants (n=738) and 48.3% male infants (n=703), with 0.9% missing gender data (n=13). The average age of the 1,453 (99.9%) infants whose ages were properly included in the data set at the time of dry blood spot (DBS) collection was 11.2±18.1 weeks with a median and mode of 7 and 6 weeks, respectively. Maternal age range was 18–40 years.

One thousand four hundred and forty-one (99.1%) had a documented reason for DBS, with 99.9% (1,440) being the first test for the healthy exposed baby. One thousand three hundred and forty (96.5%) infants received ARV prophylaxis for HIV, 1,285 (92.2%) were breastfed, and 89.1% were breastfed exclusively. Extracted data showed that among those with relevant details, 85.4% were still on breast milk at the time of DBS collection and 91.3% received cotrimoxazole prophylaxis (Table 3).

Table 1 Data extraction details from study facilities in Abuja and Nasarawa

| Facilities | Level of care | No extracted | Frequency (%) |
|------------|---------------|--------------|---------------|
| A          | Secondary     | 116          | 8.0           |
| B          | Primary       | 53           | 3.6           |
| C          | Primary       | 17           | 1.2           |
| D          | Primary       | 51           | 3.5           |
| E          | Secondary     | 264          | 18.1          |
| F          | Secondary     | 100          | 6.9           |
| G          | Secondary     | 727          | 50.0          |
| H          | Primary       | 126          | 8.7           |
| Total      |               | 1,454        | 100.0         |

Table 2 DBS collection and maternal ART details of PMTCT clients in study facilities

| Frequency | Percentage |
|-----------|------------|
| DBS Reason for DBS | | |
| First test for healthy exposed baby | 1,440 | 99.9 |
| First test for a sick baby | 1 | 0.1 |
| Total | 1,441 | 100.0 |
| Mother on ART | | |
| Yes | 1,302 | 91.0 |
| No | 67 | 4.7 |
| Unknown | 62 | 4.3 |
| Total | 1,431 | 100.0 |
| Time when ART was commenced | | |
| Description | | |
| ART started before pregnancy | 357 | 30.1 |
| ART started during pregnancy | 445 | 37.5 |
| ART started after pregnancy | 4 | 0.3 |
| Unknown | 380 | 32.1 |
| Total | 1,186 | 100.0 |
| Mother receiving ART | | |
| Yes | 1,166 | 93.8 |
| No | 23 | 1.9 |
| Unknown | 54 | 4.3 |
| Total | 1,243 | 100.0 |
| ART regimen for mothers | | |
| AZT/3TC/sdNVP in labor | 30 | 2.6 |
| AZT/sdNVP in labor | 2 | 0.2 |
| Triple regimen | 1,118 | 96.3 |
| Triple regimen | 1 | 0.1 |
| Unknown | 6 | 0.5 |
| None | 3 | 0.3 |
| Total | 1,160 | 100.0 |

Abbreviations: ART, antiretroviral treatment; DBS, dry blood spot; PMTCT, prevention of mother-to-child transmission; AZT, zidovudine; 3TC, lamivudine; sdNVP, single dose nevirapine.
The first EID HIV positivity rate was 3.5% as 47 of 1,339 infants tested positive to HIV, and the second EID HIV positivity rate was 1% among the exposed. Positivity rate for first EID ranged from 0.0% in three facilities to 11.1% in one facility ($\chi^2=24.99, P<0.00$). Mothers who were on ART were statistically less likely to have HIV-infected infants ($\chi^2=54.71, P<0.00$). Also, infants of mothers who received ARV prophylaxis ($\chi^2=97.49, P<0.00$), infants who received ARV prophylaxis ($\chi^2=67.44, P<0.00$), infants who were exclusively breastfed ($\chi^2=14.07, P<0.00$), and infants who received cotrimoxazole prophylaxis ($\chi^2=55.97, P<0.00$) were statistically less likely to be HIV infected when compared to the rest (Table 4).

Most deliveries took place in the primary health care center (61.5%), followed by secondary facilities (23.1%) and maternity homes (14.9%). Others had their babies at their private homes (0.8%). The place of delivery ($\chi^2=1.49, P=0.68$) was statistically not significantly related to EID results. From the total birth, 95.5% were delivered through spontaneous vaginal delivery (SVD). The mode of delivery ($\chi^2=6.62, P=0.01$) was significantly associated with positive EID results.

The reasons for second EID included repeat test after cessation of breastfeeding ($n=133, 85.8\%$) and repeat test to confirm the original result ($n=12, 7.7\%$). Other reasons included first testing for the healthy exposed baby and follow-up for breastfeeding children ($n=3, 0.9\%$ each) and repeat testing because of technical problems with the first test ($n=2, 1.3\%$). The rest included first testing for a sick baby and/or to confirm positive PCR.

In the final outcome results analysis, those who became sick or died increased from 1.3% to 2.7% between the 6 and 12 months postdelivery.

### Discussion

DBS for HIV PCR was performed on all infants, and in 99.9% of these infants, DBS was the first test for a healthy exposed baby. The rest were tests for sick babies. The high DBS performance rate is a pointer to quality PMTCT services, as all children from both booked and unbooked mothers were tested for HIV at delivery or soon afterward. Thus, the number of sick infants who were tested for the first time, simply because they were sick, was a very small proportion of the total. However, the majority of the repeat tests was needed after cessation of breastfeeding (85.8%) and to confirm the first PCR test (7.7%). This result may be skewed because of lost to follow-up of most infants and their infected mothers. However, this will need further studies for proper elucidation.

As mentioned in previous studies, ART services, ARV prophylaxis, exclusively breastfeeding, and cotrimoxazole prophylaxis protected the HIV-exposed infants (HEIs) from infection before, during, and after birth, and this is statistically significant.\(^3\) The calculated positivity rate from this study of 3.5% is well within the acceptable limits.\(^4\) This is similar to the finding from a South African study\(^11\) but lower than a similar Nigerian study in 15 supported sites in five states, where the positively rate was found to be 5.3%. Other reported studies from South Africa and Kenya recorded positivity rates of 8.8% and 10%, respectively.\(^14,15\)

On an individual basis, some sites had no HIV-positive children due to high-quality PMTCT services, while two centers had 8.0% and 11.0% positivity rates, respectively. These positivity rates were far higher than the expected <5% positivity rate, and the difference between centers was statistically significant. Additional review showed that the majority

### Table 3: Infant breastfeeding, ARV, and cotrimoxazole prophylaxis pattern for exposed children in supported sites

| Response                        | Frequency | Percentage |
|---------------------------------|-----------|------------|
| Infant prophylaxis              |           |            |
| Yes                             | 1,340     | 96.5       |
| No                              | 40        | 2.9        |
| Unknown                         | 9         | 0.6        |
| Total                           | 1,389     | 100.0      |
| Infant breastfed                |           |            |
| Yes                             | 1,285     | 92.2       |
| No                              | 107       | 7.7        |
| Unknown                         | 1         | 0.1        |
| Total                           | 1,393     | 100.0      |
| Mode of breastfeeding           |           |            |
| Exclusive feeding               | 1,133     | 89.1       |
| Mixed feeding                   | 139       | 10.9       |
| Total                           | 1,272     | 100.0      |
| Still breastfeeding at the time of DBS |   |      |
| Yes                             | 472       | 85.4       |
| No                              | 42        | 7.6        |
| Unknown                         | 39        | 7.0        |
| Total                           | 553       | 100.0      |
| Infant cotrimoxazole for prophylaxis |         |            |
| Yes                             | 1,092     | 93.1       |
| No                              | 28        | 2.4        |
| Unknown                         | 53        | 4.5        |
| Total                           | 1,173     | 100.0      |

**Abbreviations:** ARV, antiretroviral drug; DBS, dry blood spot.
of the HIV-positive cases were women who were “unbooked”. Booking refers to the initial registration for antenatal care and PMTCT services. Therefore, the majority came into the facility to deliver for the first time and did not receive ARV prophylaxis as a consequence.

Regarding breastfeeding, the latest WHO guidelines recommend the national authorities to promote one infant practice among mothers with HIV, either exclusive breastfeeding while ARV drugs are provided or avoiding all breast milk.\textsuperscript{16,17} The breastfeeding rate (92.2\%) was higher than findings from other African studies,\textsuperscript{11} which may have been linked to cultural practices that support breastfeeding in Nigeria. However, PMTCT may have helped to improve exclusive breastfeeding to 89.1\% and cotrimoxazole prophylaxis to 91.3\% among the surveyed community against 17\% and 13\% exclusive breastfeeding rate in 2013 and 2008, respectively.\textsuperscript{16,17} Studies have also shown that environmental factors such as access to piped water, electricity, gas, and

| Table 4 Cross tabulation of EID results with various independent variables |
|-----------------------------|-------|-------|-------|-----|
|                            | Yes   | No    | Unknown | Total | Chi square |
| **EID vs mother on ART**    |       |       |         |       |            |
| HIV negative                | 1180  | 51    | 48      | 1279  | 54.71; P<0.00 |
| HIV positive                | 27    | 11    | 6       | 44    |            |
| Total                       | 1207  | 62    | 54      | 1323  |            |
| **EID and mother received ART prophylaxis** |       |       |         |       |            |
| HIV negative                | 1061  | 14    | 39      | 1114  | 97.49; P<0.00 |
| HIV positive                | 24    | 8     | 7       | 39    |            |
| Total                       | 1085  | 22    | 46      | 1131  |            |
| **EID and infant on ART prophylaxis** |       |       |         |       |            |
| HIV negative                | 1216  | 30    | 7       | 1253  | 67.44; P<0.00 |
| HIV positive                | 28    | 9     | 2       | 39    |            |
| Total                       | 1244  | 39    | 9       | 1292  |            |
| **EID and breastfeeding**   |       |       |         |       |            |
| HIV negative                | 1158  | 98    | 1       | 1257  | 0.04; P<0.98 |
| HIV positive                | 37    | 3     | 0       | 40    |            |
| Total                       | 1195  | 101   | 1       | 1297  |            |
| **EID and mode of breastfeeding** |       |       |         |       |            |
| HIV negative                | 1025  | 121   | 1146    | 116  | 14.07; P<0.00 |
| HIV positive                | 25    | 11    | 36      | 36    | X\textsubscript{c}y^{2} =12.13; P<0.00 |
| Total                       | 1050  | 132   | 1182    |      |            |
| **EID and mode of delivery** |       |       |         |       |            |
| HIV negative                | 55    | 2     | 57      | 66    | 6.62; P<0.01 |
| HIV positive                | 1     | 0     | 1       | 1     |            |
| Total                       | 56    | 2     | 58      |      |            |
| **EID and baby on cotrimoxazole prophylaxis** |       |       |         |       |            |
| HIV negative                | 999   | 20    | 48      | 1067  | 55.97; P<0.00 |
| HIV positive                | 22    | 7     | 1       | 30    |            |
| Total                       | 1021  | 27    | 49      | 1097  |            |
| **EID and place of delivery** |       |       |         |       |            |
| HIV Negative                | 1     | 17    | 73      | 26    | 1.49; P<0.68 |
| HIV Positive                | 0     | 1     | 4       | 0     |            |
| Total                       | 1     | 18    | 77      | 26    |            |
| **EID and facility**        |       |       |         |       |            |
| Chi Square                  | 24.99 |      | P<0.00  |      |            |

**Abbreviations:** ART, antiretroviral treatment; EID, early infant diagnosis.
paraffin for fuel, limit the implementation of the WHO/United Nations Children’s Fund guidelines. This results in inappropriate infant-feeding choices such as mixed feeding and lower infant HIV-free survival.

While the place of delivery was not statistically significantly associated with MTCT, the mode of delivery (cesarean section vs SVD) was statistically significant ($P<0.00$). Improved tracking of HEIs is needed in PMTCT programs, where access to EID is still limited as the lowest prevalence of infant HIV infection or death was observed to occur among children completing the cascade.

In this secondary data analysis, adherence, economic costs, the mothers’ behavior during highly active antiretroviral treatment (HAART), and the child feeding program during the therapy were not evaluated to ascertain how they could have affected the overall efficacy of ARV services in PMTCT. These are possible issues to be explored in future studies. Finally, one of the strengths of this study is that the result is free of self-reporting bias (including social desirability bias) seen in other studies where women were asked to self-report their ART use.

Despite the effectiveness of PMTCT services, pediatric HIV infection is still common in Nigeria with the country contributing >30% to the global burden of the disease. Several nations in North America and Europe including Cuba have virtually eliminated pediatric AIDS, but to achieve this in Nigeria, there is the need to periodically assess the effectiveness of the PMTCT program and, based on the findings, modify the strategies and processes in place.

To accelerate PMTCT uptake and ensure that no woman or child is left behind, health care workers should initiate and facilitate provider-initiated and sustained patient counseling (PISC). PISC has five basic components: 1) every pregnant woman should be tested for HIV (and given their results); 2) PPW should be placed on ART or ARV prophylaxis once they are identified during pregnancy or in labor; 3) HEIs should be placed on ARV prophylaxis immediately after birth or at first appearance at a health care facility; 4) all HEIs should receive cotrimoxazole prophylaxis as long as it is required; and 5) HEI should not be breastfed if possible. But if breastfeeding is necessary, exclusive breastfeeding should be encouraged.

**Limitations**

PMTCT services in Nigeria are fraught with multiple challenges. In this study, there were missing information from incomplete documentation, lost to follow-up, and delay in result delivery to the facility and to caregivers. Dates were not properly entered and thus were difficult to analyze. This is not unexpected as previous studies have documented the various difficulties encountered (including personnel and infrastructure requirements) with implementing PMTCT and EID services in resource-limited settings, such as Nigeria, which make proper monitored, effective counseling and quality service delivery difficult, if not impossible. Thus, despite the substantial benefits of PMTCT and EID to HIV-infected and HIV-uninfected infants, their families, and programs providing PMTCT services, access to quality services is hindered by these limitations. Only eight sites in two states were analyzed for this study, and therefore, the results are not representative of Nigeria. Although the findings are instructive and should guide programming, there is a need for a wider national survey using nationally representative samples. There will also be the need to compare implementing partners’ results, states, and facilities outcomes to identify key challenges and barriers toward eliminating MTCT of HIV in Nigeria.

**Conclusion**

Although services have been fully decentralized in Nigeria to primary health care facilities, large number of clients (82.2%) still visits the secondary-level facilities, as shown in this study. Improving infrastructures and human resources at the secondary level of care will facilitate the provision of quality and sustainable PMTCT services to pregnant women and supervisory support to linked PHCs. This will culminate in better outcome for all HIV-PPW and HEIs.

To ensure MTCT of HIV is eliminated and impact on the infant minimized, there is a need to make greater efforts to reach all mothers, provide care to all PPW, and screen all HEIs early enough to provide care and treatment to the infected individuals.

**Acknowledgments**

We acknowledge the excellent support of all staff and members of Excellence and Friends Management Care Center (EFMC) and management; staff of all supported PMTCT facilities in Abuja and Nasarawa; and Institute of Human Virology, Nigeria (IHVN) for technical assistance in the implementation of this project. SDT-R is grateful to the NIHR Biomedical Facility at Imperial College London for infrastructure support. This study was funded in part by the Wellcome Institutional Strategic Support Fund at Imperial College London.
Author contributions
All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; gave final approval of the version to be published; and agree to be accountable for all aspects of the work.

Disclosure
The authors report no conflicts of interest in this work.

References
1. United Nations AIDS. Fact sheet – Latest statistics on the status of the AIDS epidemic. [Updated 2017]. Available from: www.unaids.org/en/resources/fact-sheet. Accessed September 11, 2018.
2. Federal Ministry of Health (FMOH). 2014 National HIV Sero-prevalence Sentinel Survey among Pregnant Women Attending Antenatal Clinics in Nigeria. Department of Public Health National AIDS/STI Control Programme. FMOH. Abuja Nigeria. 2015.
3. de Cock KM, Fowler MG, Mercier E, et al. Prevention of mother-to-child HIV transmission in resource-poor countries: translating research into policy and practice. JAMA. 2000;283(9):1175–1182.
4. World Health Organization (WHO). Mother-to-child transmission of HIV. Available from: http://www.who.int/hiv/topics/mtct/en/. Accessed March 2, 2017.
5. UNICEF. Global Summary of HIV epidemic among Children (0-14 years), Western and Central Africa, 2015. Available from: https://data.unicef.org/wp-content/uploads/2015/12/2016_West-and-Central-Africa.xlsx. Accessed March 2, 2017.
6. UNICEF. Nigeria PMTCT Fact Sheet. Available from: http://www.unicef.org/aids/files/Nigeria_PMTCTFactSheet_2010.pdf. Accessed March 2, 2017.
7. Padian NS, McCoy SI, Karim SS, et al. HIV prevention transformed: a new prevention research agenda. Lancet. 2011;378(9787):269–278.
8. World Health Organization (WHO). Guideline on when to start antiretroviral therapy and on pre-exposure prophylaxis for HIV; September 2015. [cited March 2, 2017]. Available from: http://www.emtct-iatt.org/wp-content/uploads/2015/09/WHO-Guidelines-on-When-to-Start-ART-and-PrEP-September-2015.pdf.
9. WHO. Global Guidance on Criteria and Processes for Validation: Elimination of Mother-to-Child Transmission of HIV and Syphilis. World Health Organization; 2014. http://apps.who.int/iris/bitstream/10665/112858/1/9789241505888_eng.pdf?ua=1&ua=1.
10. World Health Organization (WHO). PMTCT strategic vision 2010-2015: preventing mother-to-child transmission of HIV to reach the UNGASS and Millennium Development Goals. Available from: http://www.who.int/hiv/pub/mtct стратегического плаана.pdf.
11. Goga AE, Dinh TH, Jackson DJ, et al. First population-level effectiveness evaluation of a national programme to prevent HIV transmission from mother to child, South Africa. J Epidemiol Community Health. 2015;69(3):240–248.
12. UNICEF. Available from: http://www.unicef.org/aids/files/Nigeria_PMTCTFactsheet_2010.pdf. Accessed March 2, 2017.
13. OpenEpi [homepage on the Internet]. Available from: http://www.openepi.com/RbyC/RbyC.htm. Accessed March 2, 2017.
14. Coetzee D, Hildebrandt K, Boullé A, Draper B, Abdullah F, Goemaere E. Effectiveness of the first district-wide programme for the prevention of mother-to-child transmission of HIV in South Africa. Bull World Health Organ. 2005;83(7):489–494.
15. Ngemu EK, Khayeka-Wandawwa C, Kweka EJ, Choge JK, Anino E, Oyoo-Ooko E. Effectiveness of option B highly active antiretroviral therapy (HAART) prevention of mother-to-child transmission (PMTCT) in pregnant women. BMC Res Notes. 2014;7(1):52.
16. The World Health Organization. Updates on HIV and Infant Feeding. Available from: apps.who.int/iris/bitstream/handle/10665/246260/9789241549707-eng.pdf?sequence=1. Accessed April, 2018.
17. Ciaranello AL, Park JE, Ramirez-Avila L, Fredberg KA, Walensky RP, Leroy V. Early infant HIV-1 diagnosis programs in resource-limited settings: opportunities for improved outcomes and more cost-effective interventions. BMC Med. 2011;20;9(1):59.
18. Doherty T, Chopra M, Jackson D, Goga A, Colvin M, Persson LA. Effectiveness of the WHO/UNICEF guidelines on infant feeding for HIV-positive women: results from a prospective cohort study in South Africa. AIDS. 2005;83(7):1791–1797.
19. Ahoua L, Ayikoru H, Gnauck K, et al. Evaluation of a 5-year programme to prevent mother-to-child transmission of HIV infection in Northern Uganda. J Trop Pediatr. 2015;69(3):240–248.
20. Chi BH, Tih PM, Zanolini A, et al. Implementation and operational research: reconstructing the PMTCT cascade using cross-sectional household survey data: the PEARL study. J Acquir Immune Defic Syndr. 2015;70(1):e5–9.
21. Mselati P. Improving access to mother-to-child transmission (PMTCT) programs in Africa: an ongoing process. In: Liamputtong P, editor. Women, Motherhood and Living with HIV/AIDS. The Netherlands: Springer; 2013: 177–187.