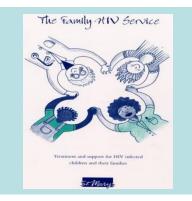
Breast Feeding with HIV in Resource Rich Settings – What do Women Want to Do?

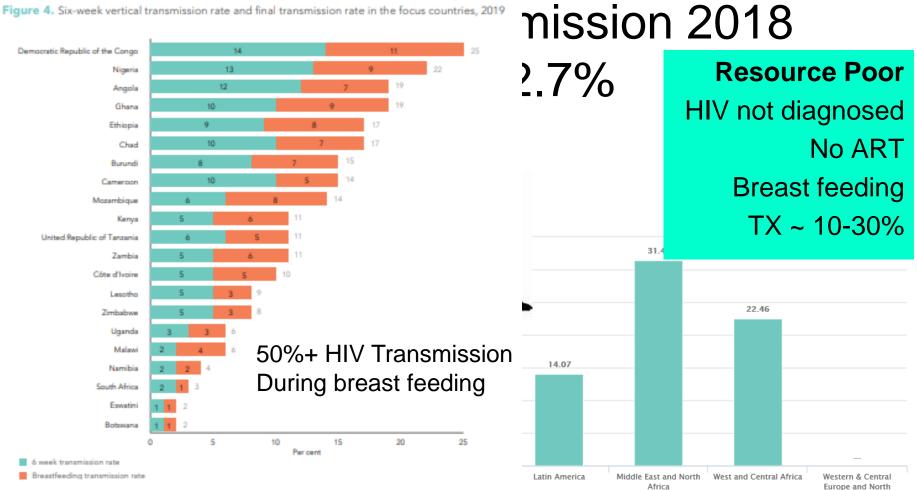


Hermione Lyall Imperial Healthcare NHS Trust 30.9.20 <u>Hermione.lyall@nhs.net</u>



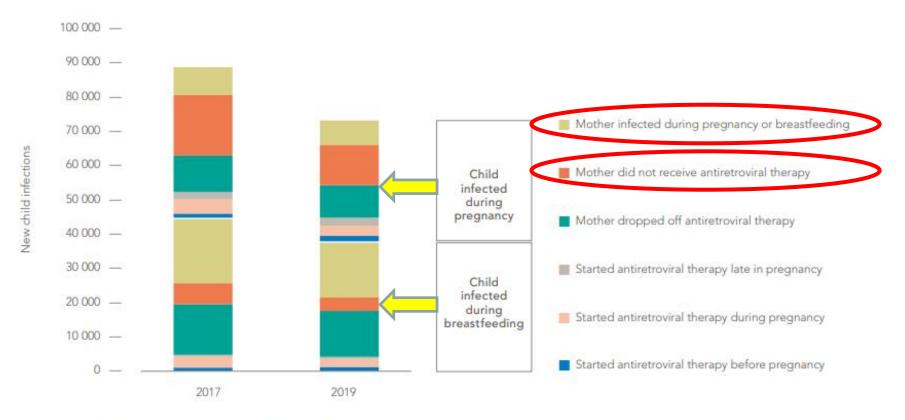
Acknowledgements:

Nell Freeman-Romilly, Pat Tookey, Yvonne Gileece & BHIVA Guidelines Team Claire Thorne, Kate Francis, Helen Peters, Graham Taylor, Paula Seery, Pat Flynne, Angela Colbers, Karoline Aebi-Popp



America

New child infections due to gaps in prevention of vertical transmission, eastern and southern Africa, 2017 and 2019



Source: UNAIDS epidemiological estimates, 2020 (see https://aidsinfo.unaids.org/).

Countries can use gap analysis → For elimination of MTCTNew ways to prevent infection in women & their infants:Protect the HIV negativeTreat the HIV positivePREP in pregnancyART which has:

PREP during breast feeding for the mother for the infant

Vaccination – passive / active for the mother for the infant



highest barrier to resistance

Least side effects

Least teratogenicity

Additional strategies beyond oral ART are required to eliminate Vertical Transmission......

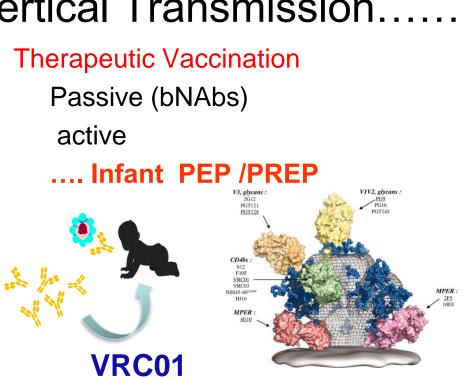
Long acting injectables

Combinations of injectables \rightarrow

Depot contraceptives Depot antipsychotics

etc





Burton et al, 2012

New options for pregnancy & post partum \rightarrow minimise delay to access

Is it safe for me to breast feed my baby?

Scenario 1:

Well woman with HIV, 31 years old First pregnancy - 20 weeks gestation Conceived on ART, still on first line ART VL < 50 for 5 years CD4 count 770



My family expect me to breast feed, I also believe it's the right thing to do -

> I have read the WHO guideline (2016) → it says breast is best for women with HIV what would you advise?

"God will cure me and my child"

Scenario 2:

G 5+0, 32/40 weeks, CD4 - 50, VL - 270,000

Denies HIV, refusing any treatment for herself before delivery for herself at delivery for the infant after delivery wants normal delivery



I want to breast feed – as I did with all my other kids

Risk of transmission to this infant?

9 yr old, 7 yr old, 5, yr old, 2 yr old – where are they, have they been tested?



WHO Guideline on HIV & Infant feeding 2016

WHO recommends lifelong ART for everyone from the time they are first diagnosed with HIV infection.

This WHO guideline is intended mainly for countries with high HIV prevalence where diarrhoea, pneumonia and under-nutrition are common causes of infant mortality.

However, it may also be relevant to settings with a low prevalence of HIV depending on the background rates and causes of infant and child mortality



GUIDELINE UPDATES ON HIV AND INFANT FEEDING



WHO Guidelines for Infant feeding 2016

		UPDATES ON HIV AND INFANT FEEDING	
Clinical Scenarios	WHO guidance for women with HIV	The duration of lossetfixeding and support from health services to improve fixeding practices among mothers living with Hill	
For how long should mothers with HIV breast feed?	 Mothers living with HIV should breastfeed for at least 12 months and may continue breastfeeding for up to 24 months or longer, if → <i>(same as the general population)</i> Has access to lifelong ART and HIV care Exclusively breastfeeds for the first 6 months Introduce appropriate complementary foods after 6 months and continue breastfeeding Only stop once a nutritionally adequate and safe diet without breast milk can be provided 		
If a mother does not exclusively breastfeed: is mixed feeding with ART better than no breastfeeding at all?	ART also reduces the risk of HIV transmission in mixed feeding Although exclusive breastfeeding is recommended - when on ART, mixed reason to stop breastfeeding	I feeding is not a	
Is a shorter duration of planned breastfeeding with ART better than no breastfeeding at all?	Any duration of breastfeeding is better than never initiating breastfeeding	at all	

Breastfeeding advice for women with HIV living in the UK

British HIV Association guidelines for the management of HIV in pregnancy and postpartum 2018 (2020 third interim update)

BHIVA獭

 In the UK and other high-income settings, the safest way to feed infants born to women with HIV is with formula milk, as there is on-going risk of HIV exposure after birth.

1C

1D

- Abstaining from breastfeeding can have financial and psychological repercussions for women, requiring support from the HIV MDT.
- Women advised not to breastfeed should be provided with free formula to minimise vertical transmission of HIV.
- Women not breastfeeding their infant by choice, or because of viral load >50, should be offered cabergoline to suppress lactation.
 1D

https://www.bhiva.org/file/5f1aab1ab9aba/BHIVA-Pregnancy-guidelines-2020-3rd-interim-update.pdf

Supporting women who choosing to breastfeed in the UK

Women who are fully suppressed on ART with good adherence and choose to breastfeed should be supported to do so.

They should be informed about the low risk of transmission of HIV through breastfeeding in this situation and the requirement for extra maternal and infant clinical monitoring. 1D

Maternal ART (rather than infant pre-exposure prophylaxis PrEP) is advised to minimise HIV transmission through breastfeeding and safeguard the woman's health. 1D

Women who choose to breast feed should fulfil the following criteria:

- A fully suppressed HIV viral load (for as long a period as possible, but certainly during the last trimester of pregnancy)
- A good adherence history
- Strong engagement with the perinatal MDT

• Prepared to attend for monthly clinic review & HIV viral load tests for themselves and their infant during and for 2 months after stopping breastfeeding

Risk Factors for HIV transmission & Breast Feeding



Viral Load CD4 count HIV sero-conversion during BF **Mastitis** Cracked nipples **Duration of BF** Mixed feeding Infant oral thrush

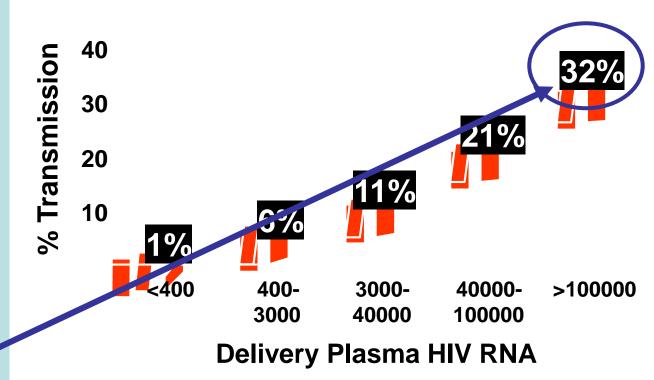
Major Risk Factors for MTCT

Maternal

Plasma viral load CD4 count Advanced HIV Delivery Premature delivery Mode of delivery Duration of rupture of **Membranes** Birth canal infection

Breast feeding -

- No ART



Blattner W et al. WITS study, 1990-1999. XIII AIDS Conf, July 2000, Durban S Africa (LBOr4)

Learning from African Studies on ART and Breastfeeding





Prevention of HIV Transmission from Breastfeeding in Africa H. Coovadia - Plenary abst 13 CROI 2007

NOT breast feeding is unsafe in developing countries

Early cessation of breastfeeding (<6 months) reduces HIV transmission but increases morbidity and mortality in infants born to HIV positive African women

Continued breast feeding reduces morbidity and mortality in HIV infected infants in Africa

Balancing the risk of: breast milk HIV transmission versus – early weaning – malnutrition – gastroenteritis - death Duration & Pattern of Breastfeeding & Postnatal HIV Transmission : Pooled Analysis from West & South African Cohorts

Becquet R, et al PLoS ONE 4(10): e7397. 2009

N = 1195 infants, not perinatally infected, & breast fed **No maternal post natal ART**

	18 month HIV infection risk
Less than 6 months BF	3.9% (2.3-6.5)
More than 6 months BF	8.7% (6.8 – 11.0)

Exclusive BF very similar to predominantly BF (only other liquids)

Solids in first 2 months of life 2.9 fold (1.1-8.0) ↑ risk of HIV

For breast feeding mothers advise \rightarrow exclusive BF & **NO** early solids

West Africa - Ditrame-Plus and South Africa – Vertical Transmission study

Feeding and mother *not on ART* Risk of postnatal HIV transmission

Becquet R, et al PLoS ONE 4(10): e7397. 2009

Estimated postnatal risk of transmission:

Overall risk: 9.0/100 child-years

(95% CI 6.2–11.7)

Exclusive breastfeeding: 9.0/100 child-years

(95% SI, 6.0-12.1)

Predominant breastfeeding: 8.5/100 child-years

(95%SI, 1.2–18.1)

Breastfeeding plus solids: 41.2/100 child-years

(95%SI, 1.1–74.5)

Breast feeding mother no ART - infant PEP ANRS 12174 - PROMISE Pre-EP

Nagot et al Lancet. 2016 Feb 6;387(10018):566-73. doi: 10.1016/S0140-6736(15)00984-8.

RCT 1500 M-I pairs BurkinaFaso, South Africa, Uganda, Zambia

HIV-uninfected infants at day 7 - born to mothers not eligible for ART Exclusive breastfeeding until 26th week of life Cessation of breastfeeding at a maximum of 49 weeks

Randomised to: infant PEP - Lamivudine or Lopinavir/ritonavir

Primary endpoint \rightarrow HIV-1 Tx - day 7 - 50 weeks of age

Secondary endpoints \rightarrow safety (including resistance, adverse events and growth) & HIV-1-free survival until 50 weeks.

Infant Pre-EP

2009 -2012 enrolled1273 infants \rightarrow analysed 1236615 \rightarrow lopinavir-ritonavir621 \rightarrow lamivudine

17 HIV-1 infections (8 lopinavir/rit versus 9 lamivudine)

50 week cumulative HIV-1 infection rate - no difference

Lopinavir/rit 1.4% (95% CI 0.4–2.5) Lamivudine 1.5% (0.7–2.5)

Clinical / biological severe adverse events - no difference

- Lopinavir/rit 251 (51%) grade 3–4 events
- Lamivudine 246 (50%) grade 3–4 events

Nagot et al Lancet. 2016 Feb 6;387(10018):566-73. doi: 10.1016/S0140-6736(15)00984-8.

Breast Feeding not on ART & Risk of HIV Transmission

Risk of HIV Transmission to the uninfected Infant	Mother in Africa Not on ART	Mother in Africa Not on ART	
after birth	Breast Feeding for 6 months exclusively	Breast Feeding for 6 months exclusively	
	Choldonvery	Infant on daily Pre-EP	African data
Duration of		Pre-EP trial (3TC/LPV/r)	
Breast feeding	Ditrame / VTS	PROMISE trial (NVP)	
6 months	3.9% (2.3-6.5)	PROMISE 0.3% (0.1-0.6)	
12 months	8.7% (6.8 – 11.0)*	PROMISE 0.6% (0·4–1.1)	
		Pre-EP 1.5% (0.7–2.5)	

*Each additional month of BF beyond 6 mths of age \rightarrow 1% risk of HIV (95%CI, 0.5–1.7)

Postnatal HIV transmission in breastfed infants of HIV-infected women on ART- meta-analysis

Bispo S et al. Journal of the International AIDS Society 2017, 20:21251

Reviewed studies 2005 to 2015 – 11 studies selected

All mother advised to exclusively breast feed for 6 months

Outcomes: overall & postnatal HIV Tx at 6, 9, 12, 18 months:

Overall 6 months Tx rate: 3.54% (95% CI: 1.15–5.93%) Overall 12 months Tx rate: 4.23% (95% CI: 2.97–5.49%)

Postnatal 6 months Tx rate:1.08 (95% CI: 0.32–1.85)Postnatal 12 months Tx rate:2.93 (95% CI: 0.68–5.18)

ART mostly provided for PMTCT and did not continue beyond 6 months postpartum No study provided data on mixed feeding & transmission risk



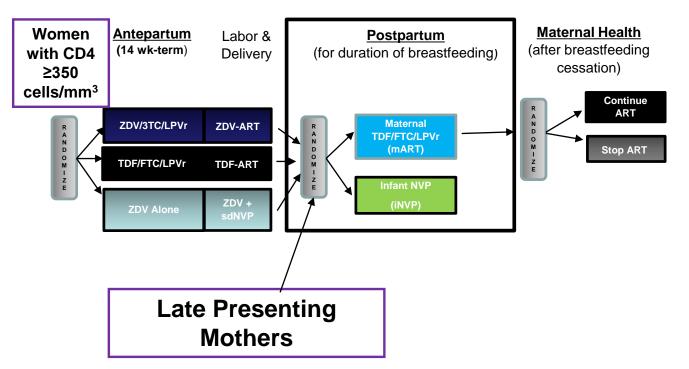
Association of Maternal Viral Load and CD4 Count with Perinatal HIV-1 Transmission Risk during Breastfeeding in the PROMISE Postpartum Component

10th Workshop on HIV Pediatrics July 20-21, 2018 Amsterdam, the Netherlands

Patricia M. Flynn, MD, Taha E Taha, MD, Mae Cababasay, MS, Kevin Butler, MS, Mary Glenn Fowler, MD, Lynne M. Mofenson, MD, Maxensia Owor, MD, Susan Fiscus, PhD, Lynda Stranix-Chibanda, MD, Anna Coutsoudis, PhD, Devasena Gnanashanmugam, MD, Nahida Chakhtoura, MD, Katie McCarthy, MPH, Cornelius Mukuzunga, MD, Bonus Makanani, MD, Dhayendre Moodley, PhD, Teacler Nematadzira, MD, Bangani Kusakara, MD, Sandesh Patil, MD, Tichaona Vhembo, MD, Raziya Bobat, MD, Blandina T Mmbaga, MD, Maysseb Masenya, MD, Mandisa Nyati, MD, Gerhard Theron, MD, Helen Mulenga, MD, David E. Shapiro, PhD and the PROMISE Study Team

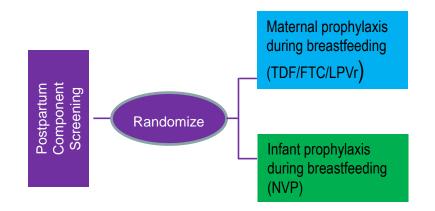
Background

 Increased maternal viral load (MVL) and decreased CD4 cell counts (CD4) have been associated with increased risk of perinatal and postnatal HIV-1 transmission



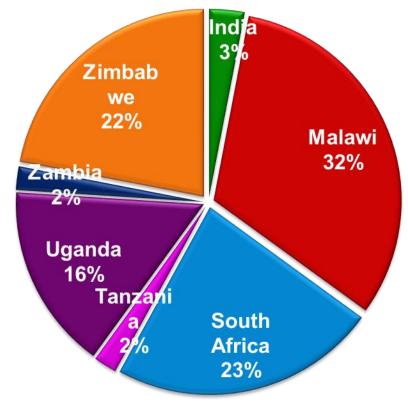
PROMISE – Postpartum Component

- Eligible mother-infant (M-I) pairs (maternal CD4 ≥ 350 cells/mm³ or country-specific level and infant HIV-1 NAT negative and > 2 kg) were randomized at 6 – 14 days postpartum to a maternal three-drug ART regimen (TDF/FTC/LPVr preferred, mART) arm or infant nevirapine (iNVP) arm
- Infants in the mART arm also received NVP for 6 weeks
- Late-presenting women were enrolled during labor or within 6 days of delivery after assuring the mother's CD4 count met eligibility criteria and infant had a negative HIV-1 NAT



PROMISE – Postpartum Component

- 2,431 M-I pairs were randomized at 6-14 days postpartum to mART (n=1,220) or iNVP (n=1,211) at 14 sites in 7 countries
- 95% of the mothers had been enrolled in the antepartum component (42% ZDV and 53% mART)
- Randomized regimens were continued until 18 months postpartum, unless stopped earlier due to cessation of breastfeeding, infant HIV-1 infection, or toxicity



Methods

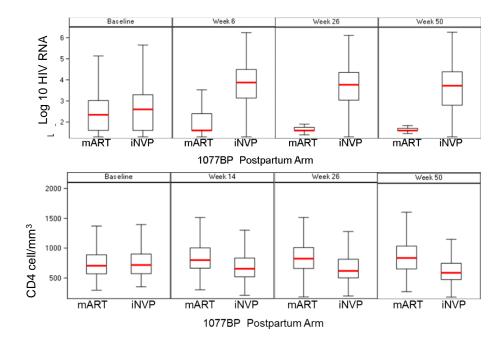
• Study Evaluations

Maternal Viral Load	Entry (6-14 days postpartum), weeks 6, 14, 26, and 50 postpartum
Maternal CD4	Entry, weeks 14, 26, 38, 50 postpartum
Infant HIV-1 NAT	Entry (6-14 days postpartum), week 6, every 4 weeks until week 26, then every 12 weeks

- Infant infection was defined as a positive HIV-1 NAT at any two post-entry timepoints
- The associations of baseline and time-varying MVL (<1000 or ≥1,000 copies/ml) and CD4 (< 500 or ≥ 500 cells/mm³) with transmission risk were assessed using proportional hazards regression models and adjustment for randomization to the mART arm during the antepartum component of PROMISE

Analysis

 For analyses using time-varying MVL and CD4, each treatment arm was analyzed separately because the post-randomization visits showed little overlap between the two arms with respect to MVL and CD4 cell count



Results

Daseline MVL and CD4 cell count by Treatment Arm			
	mART	iNVP	
	n=1,220	n=1,211	
Baseline Maternal Viral			
<u>Load</u>			
< 1,000 copies/mL	911 (75%)	814 (67%)	
≥ 1,000 copies/mL	309 (25%)	397 (33%)	
Baseline CD4 count			
< 500 cells/mm³	162 (13%)	170 (14%)	
≥ 500 cells/mm³	1,058 (87%)	1,041 (86%)	

• Baseline MVL (p=0.11) and CD4 cell count (p=0.51) were not significantly associated with infant HIV-1 transmission

Results, continued

- •Time-varying MVL was significantly associated with infant HIV-1 infection in the mART arm but not in the iNVP arm
- •Time-varying CD4 was significantly associated with infant HIV-1 infection in the mART arm but not in the iNVP arm
- •Adjusting for whether or not the mother was randomized to the mART arm in the antepartum component of PROMISE component did not change these findings

	Hazards Ratio (95% Confidence Interval)		
	mART	iNVP	
Time- varying MVL	13.96 (3.12-62.45)	1.04 (0.20-5.39)	
Time- varying CD4	0.18 (0.03-0.93)	0.38 (0.08-1.77	

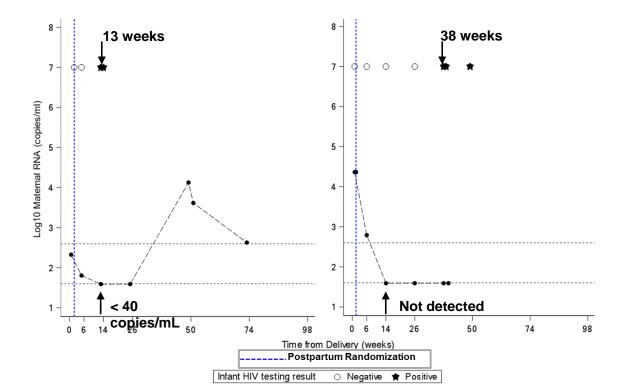
Infant HIV-1 Infections

There were seven infants with HIV-1 infection in each treatment group

	mART	iNVP
HIV-1 infections	n=7	n=7
Median infant age at first positive HIV-1 NAT	38 weeks (range, 13-50 weeks)	26 weeks (range, 6-74 weeks)
MVL closest and prior to first positive infant HIV-1 NAT	Not detected – 52,002 copies/mL	815 – 153,963 copies/mL

Results, continued

 In the mART arm there were two infected infant cases where MVL was undetectable or < 40 copies/mL in assessments prior to first positive infant HIV-1 NAT



Conclusions

- In the iNVP arm, time-varying MVL and CD4 were not significantly associated with HIV-1 transmission during breastfeeding. However, in the mART arm, increased MVL and decreased CD4 during breastfeeding were associated with increased risk of infant HIV-1 infection
- Two infant transmissions were observed following periods of MVL that were < 40 copies/mL
- These data emphasize the importance of adherence to mART in controlling MVL and preventing infant HIV-1 infection
- iNVP could be considered in situations with documented poor maternal ART adherence

Breastfeeding with maternal antiretroviral therapy or formula feeding to prevent HIV postnatal mother-to-child transmission in Rwanda

"AMATA" study

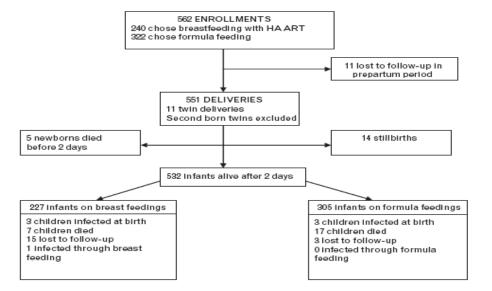
Cécile Alexandra Peltier^a, Gilles François Ndayisaba^a, Vilippe Lepage^b, Johan van Griensven^a, Valériane Leroy^c, Crristine Omes Pharm^a, Patrick Cyaga Ndimubanzi^a, Olivier Courteille^a and Vic Arendt^d

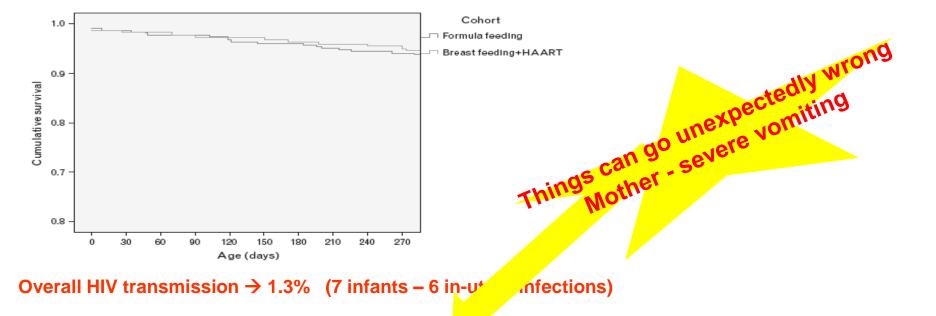
Non randomised Interventional cohort study:

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BF + ART for 6 months
∨
Formula feeding
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All received ART from 28wks

9mth cumulative risk of HIV transmission rate & HIV free survival





One infant in the BF group infected at 3-7 months

9 month post natal infection risk with BF 0.5% (95% CI – 0.1-3.4%; p =0.24) 9 month cumulative mortality:

3.3% in BF group (95% CI – 1.6 – 6.9%)

5.7% in FF group (95% CI - 3.6 - 9.2%) (p = 0.2)

HIV free survival at 9 months:

95% in BF group (95% CI – 91-97%) 94% in FF group (95% CI – 91-96%) (p = 0.66)

Europe - Breast Feeding on ART & Risk of HIV Tx

Risk of HIV Transmission to the uninfected Infant after birth Duration of Breast feeding	Mother in Europe On ART (long term) Formula Feeding	Mother in Europe On ART (long term) Breast Feeding	Mother in Africa On ART (most ART only for 6 months) Breast Feeding for 6 months exclusively then adding complementary foods Meta-analysis*	Mother in Africa On ART (long term) Breast Feeding for 6 months exclusively then adding complementary foods PROMISE Trial
6 months	0	No data	1.08% (0.32-1.85)	0.3% (0.1-0.8)
12 months	0	No data	2.93% (0.68-5.18)	0.7% (0.3-1.4)
18 months	0	No data	No data	0.7% (0.3-1.4)
24 months	0	No data	No data	0.7% (0.4-1.5)

What we learn from African Studies on ART and Breastfeeding

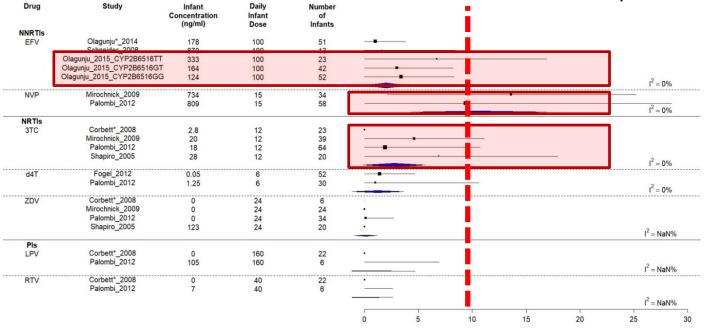




Maternal ART / infant ART & Breast feeding MTCT ~ 0.7%

Infant Exposure to ARVs in Breast milk - low

Waitt and Bonnett, CROI 2018 Abstract 55 Updated from Waitt JAC 2015



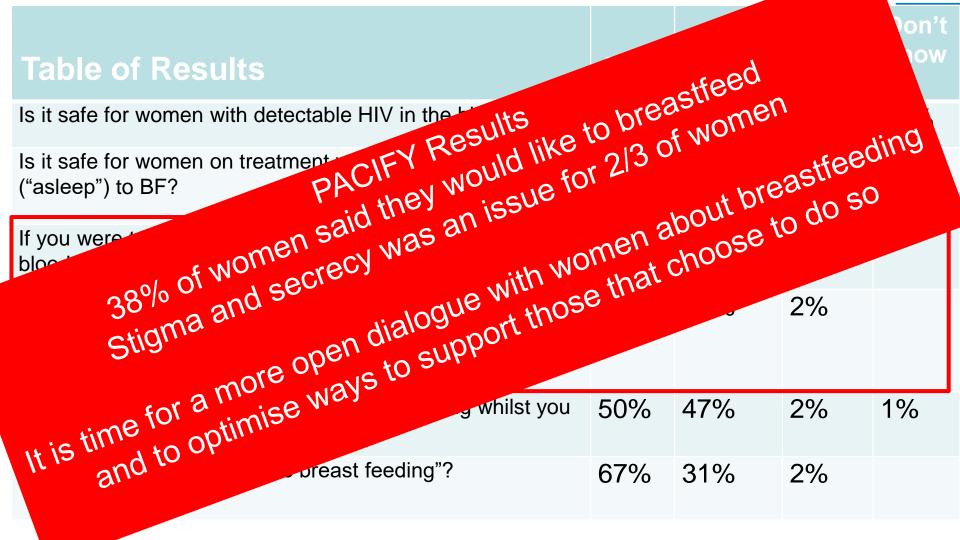
as percentage of recommended paediatric dose

TFV (given as maternal TDF): Infant concentrations not detectable DTG ~ 1% infant mg/Kg dose

PACIFY – Survey of Women's attitudes to BF with HIV in the UK (Farai Nyatsanza, HIV Women's Meeting, Seattle 2019)

Women recruited from 12 UK clinics June 2017–June 2018 94 women responded to our questionnaire (either pregnant or up to 3 months post partum) 69% of participants were Black African Median age 36 years (range 20-44) Median CD4 count was 618 cells/mm³ 92% had undetectable HIV viral load at delivery (1 had a HIV viral load of 268 copies/ml, 7 no data)

Table of Results	Yes	Νο	No answer	Don't know
Have healthcare workers discussed BF with you?	89%	9%	2%	
Would you like more information on BF with HIV?	48%	51%	1%	
Have you ever had to lie about your reasons for not breast feeding?	66%	22%	12%	
Have any friends, family or members of your community ever questioned you about your reasons for not breast feeding?	62%	27%	12%	
Living with HIV, would you like to breast feed your child?	38%	48%	7%	6%







How many women are breast feeding?

NSHPC conducts **national surveillance** of all pregnancies to women living with HIV in the UK/Ireland

Enhanced surveillance: data collected by phone for **all reported cases of planned/supported breastfeeding** with paediatric and maternity respondents and included:

- Reasons for wanting to breastfeed
- Whether the woman's partner and GP knew her HIV status
- Duration of breastfeeding
- Whether any mixed feeding occurred before 6 months of age
- Details of maternal and infant test results during breastfeeding
- Maternal cART during breastfeeding
- Infant confirmatory antibody tests (18-24mths)





2012-2019: UK - 7187 livebirths to HIV diagnosed women

1.9% (135/7187) reported planned and/or supported to BF

13% (18/135) BF more than one infant

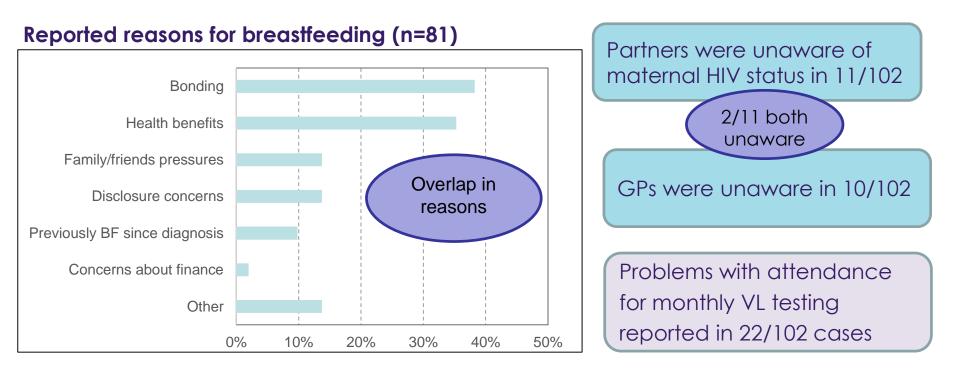
93% (125/135) women diagnosed before pregnancy83% (112/135) women born abroad

Median age at delivery was 35yrs (IQR: 31,40)

UC

Results

Enhanced data collection has been carried out for 102 supported BF cases to date:





Breastfeeding had stopped in 90/102, 3/102 not known (LTF)

Wide range of duration of BF: ranged from 1 day- 2 years Median duration: 7wk (IQR: 3, 16)

Reasons for stopping BF:

plan to stop (36), mastitis (3), VL rebound (4), problems latching (6), hospitalisation of mother and/or infant (2)

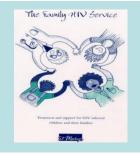
Mixed feeding before 6 months of age \rightarrow reported in 10/90 cases Mastitis \rightarrow reported in 2 cases where breastfeeding continued

2015-16 - one postnatal transmission likely due to covert BF -Edin orted BF cases so for... 2013-10-one posmarar manshole throughout pregnancy by a woman who was undetectable throughout pregnancy

L

Highlights the importance of HARM REDUCTION & 🗾 reported at least 1 detectable VL

the importance of the as possible with 4/90: atory antiber brights the importance of the as breakt



Breast feeding experiences of mothers with HIV from two UK centres

Paula Seery^{1,2}, Hermione Lyall^{1,2}, Moira Marks¹, Sophie Raghunanan¹, Caroline Foster¹, Waheed Khan², Sarah Dermont² ¹Imperial College Healthcare NHS Trust, ²Chelsea & Westminster Hospital NHS Foundation Trust

Background: 8 women breast fed (BF) 10 babies between 2012-2017 in two London Hospitals

Methods: A retrospective record review of BF mothers and babies

Results:

Women: aged 35-46yrs (median 38.25yrs), 6/8 Black African.

7 on ART from conception, all CD4 counts >500, all VL <50 at delivery.

Babies: 3 vaginal delivery, 7 elective/ emergency section. Median gestation 38+3. All babies received 4 weeks Zidovudine PEP.

Problem s	Total	Mastitis	Diarrhoe a +/- vomiting	↓Infant weight	Abnor mal LFTs	Other
Mother Baby Age	3	1 FF \rightarrow BF \rightarrow MF <1 week	1 BF \rightarrow FF \rightarrow MF <1 week			1 Urticaria BF 6 weeks
Baby Baby Age	5		1 BF	${}^{1}_{\text{BF}} \rightarrow \text{MF}$	1 BF	2 Baby reluctant to stop BF
			19 months	<1 week	2-7 months	>12 < 22 months

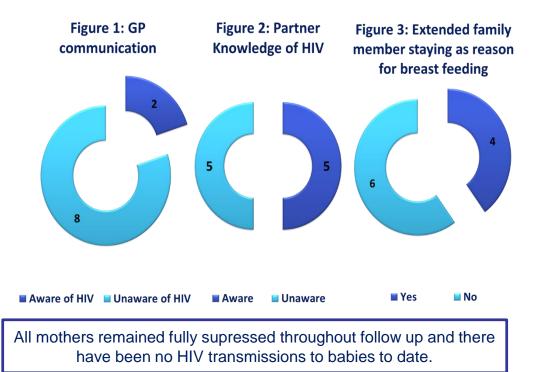
Breastfeeding : 3 babies BF for <1 week (2/3 pasteurised EBM)

- 7 babies, average BF duration → 33 weeks (range 4 – 95 wks)
- 3 babies were exclusively BF 4 mixed fed
- → 3/4 introducing formula for clinical reasons (table)

1 baby was covertly breast fed (mixed fed)

Monitoring: 6 babies additional blood tests to routine follow up (1-19 extra tests) Mothers \rightarrow on average 6 extra VL tests

One mother & baby travelled abroad unmonitored for 6 months



Conclusion: Women who BF enjoyed many benefits but also faced challenges.

A higher proportion than expected had not informed partners, family, or primary HCPs about their HIV.

Raising concerns BF could be part of maintaining "the secret"

Breastfeeding Advice as Harm Reduction

"People will make more health-positive choices if they have access to adequate support, empowerment, and education".

Levison, J., Weber, S. and Cohan, D. (2014). Breastfeeding and HIV-Infected Women in the United States: Harm Reduction Counseling Strategies. Clinical Infectious Diseases, 59(2), pp.304-309.

Patient Information on HIV and Breastfeeding

Which simplifies complex (and changing) information

╋

Accounts for patient's wishes

╋

Persuasively guides patients towards the safest approach

The Safer Triangle

No virus Only breastfeed if your HIV is undetectable. Happy tums

Only breastfeed if both you and your baby are free from tummy problems

Healthy breasts for mums Only breastfeed if your breasts and nipples are healthy with no signs of injury or infection.

BHIVA Pregnancy guidelines 2018 Two Patient Leaflets BHIVA

1 – for all pregnant women with HIV: 'General information on Feeding Your Baby'

2- for women with HIV who want to breastfeed: *'HIV and Breastfeeding Your Baby'*

HIV and breastfeeding your baby

The safest way for a mother living with HIV in the UK to feed her baby is to bottle feed using formula milk.

If you are on treatment with an undetectable viral load and choose to breastfeed your baby we can help you make it as safe as possible for your baby, but it will not be as safe as using formula. Until we know more about the safety of breastfeeding on antiretroviral therapy, our careful guidance will give your baby his or her best chance of remaining HIV free while being breastfed. Always protect your baby using 'The Safer Triangle' below:

No virus

If the HIV virus in your blood is detectable, there will be HIV in your breast milk, and HIV will enter your baby's body on feeding. You should only breastfeed your baby if your HIV is undetectable.

Happy tums

Diarhoes and vomiting show that a tummy is irritated. If your baby's tummy is irritated it may be more likely that HIV will cross into the blood seam and infect your baby. If your tummy is initiated you may not absorb your HIV medication properly. Only breastfeed if both of you have a happy tummy.

Healthy breasts for mums

There maybe HIV in your breast milk if your nipples are cracked or bleeding, or if you have thrush, develop an infection or have mastitis. Only breastfeed if your breasts are healthy.

The Safer Triangle means:

No Virus + Happy Tums + Healthy Breasts for Mums

Only breastfeed if your HIV is undetectable AND both you and your baby are free from tummy problems AND your breasts and nipples are healthy with no signs of infection.

If HIV virus becomes detectable in your blood: Stop breastfeeding and start using formula milk. Do not use breast milk you have expressed and stored. Feed your baby using formula only until you have spoken with your HIV clinic.

If your baby has diarrhoea or vomiting: Feed your baby with formula milk only. Keep feeding your baby using formula milk even after their tummy is healed.

If you have diarrhoea or vomiting, or your breasts have an injury or infection: Stop breastfeeding and feed your baby with formula milk OR use breast milk that you expressed more than 2 days (48 hours) before your tummy or breast problem began. If your baby has formula milk while you are ill, continue feeding your baby formula milk only. If your baby did not receive formula milk you may return to breastfeeding 2 days (48 hours) after your breast problem is healed. If you had tummy problems you must contact your HIV clinic before breastfeeding.

with HIV formula eir babies:

come HIV positive

If 100 mothers with HIV breas while having an undetectal

1-2 babies may be

Testing for HIV in infants born to breast feeding mothers with HIV on ART

Formula Fed Infant 4 (5 if high risk) blood tests

Birth HIV PCR High risk infants - additional week 2-3 HIV PCR

Week 6 HIV PCR (off PEP) Week 12 HIV PCR (off PEP)

Loss of HIV antibody at 18 months

Breastfed Infant 4 + X monthly blood tests

Birth HIV PCR Clinical review & monthly HIV PCR when Breast feeding

Week 4 HIV PCR (off BF) Week 8 HIV PCR (off BF)

Loss of HIV antibody at 18 months

Back to our 2 women – breast feeding?



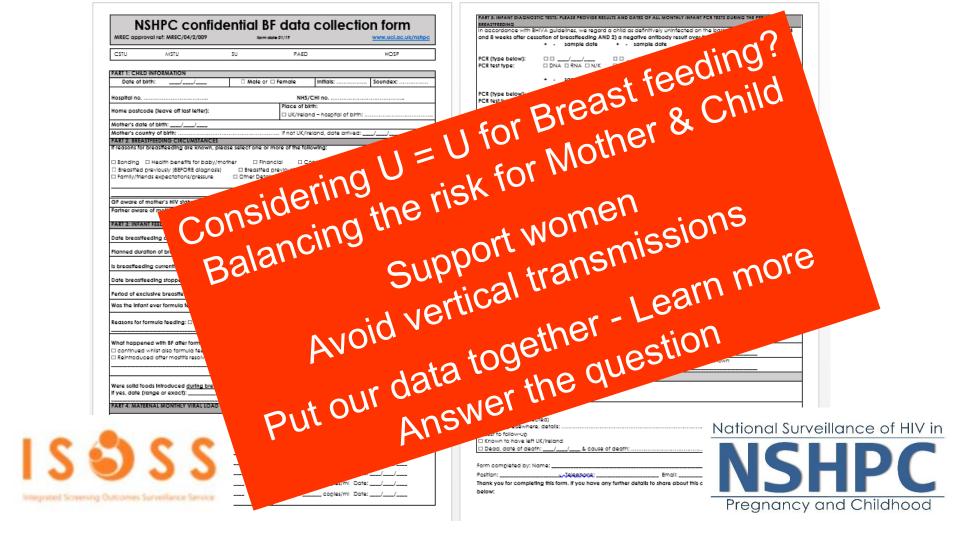
Scenario 2

CD4 - 50, VL - 270,000 Does not believe in HIV Not on ART

FF ~ 15% risk of HIV Tx BF ~ 30% risk of HIV Tx

Not engaging with MDT, unlikley to comply with ART, \rightarrow antenatal SC referral

We **not** would support BF → SC referral



Thank You

