



HIV and Contraception, Preconception and Reproductive Health

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Statewide Clinical HIV Update
June 17, 2016





Disclosure

No relevant financial disclosures





Objectives

- Review options for ART for reproductive capable HIV infected women.
- Describe contraceptive options for women living with HIV.
- Explain specific consideration related to hormonal contraception and antiretroviral treatment
- Discuss reproductive strategies for HIV-infected individuals including pre-exposure prophylaxis (PrEP).
- Discuss issues with postpartum retention in HIV care.





Case 1- Sheila

- 27 yo female newly diagnosed with HIV after testing at STD clinic
- CD4 586, PVL 21,000 copies/mL
- No children, active with one sexual partner, occasional condom use.





Do you discuss pregnancy plans with HIV(+) women in your care?

- 1. At intake only
- 2. Once a year
- 3. Only if patient initiates
- 4. At every visit
- 5. Never





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Introduction

Recommendations in these guidelines are based on scientific evidence and expert opinion and are rated using the system below:

Strength of Recommendation	Quality of Evidence
A: Strong	I: One or more randomized trials with clinical outcomes and/or validated laboratory end
B: Moderate	points
C: Optional	II: One or more well-designed, nonrandomized trials or observational studies with long-term clinical outcomes
	III: Expert opinion

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Science: There is evidence that individual components of preconception care work:

and

- ☐ Rubella vaccination
- ☐HIV/AIDS screening
- ☐ Management control of:
 - **□** Diabetes
 - Hypothyroidism
 - **□**PKU
 - **□**Obesity

- Folic acid supplements
- Avoiding teratogens:
 - Smoking
 - Alcohol
 - Oral anticoagulants
 - Accutane





Preconception Counseling and Care

Purpose:

- Prevention of unintended pregnancies.
- Optimization of maternal health prior to pregnancy.
- Prevention of perinatal transmission.
- Prevention of HIV transmission to an uninfected partner while trying to conceive.





ART for HIV-infected women of childbearing age

- A regimen's effectiveness.
- A woman's hepatitis B status.
- Teratogenic potential of the drugs in the cART regimen. (Efavirenz* -pregnancy testing)
- Possible adverse outcomes for the mother and fetus.





Case 2—Roberta

- □ 30 year-old woman tested HIV+ positive during her recent pregnancy and started HIV treatment with Complera (tdf/ftc/rilpivirine)
- □ CD4 (T-cells) have improved on treatment and her viral load is undetectable
- Infant is 4 months old and HIV-uninfected
- ☐ Plan:
 - Renew medications today, check labs before she returns for a check up in 3 months.
 - Encourage adherence
 - Remind to use condoms





Case 2—Roberta...

☐ You ask about contraception.

- ☐She previously used oral contraceptives and asks about restarting them.
- ☐ How do you counsel her?





Preconception Counseling and Care

Recommendations

 Discuss childbearing intentions with all women of childbearing age on an ongoing basis throughout the course of their care (AIII).

 Provide information about effective and appropriate contraceptive methods to reduce the likelihood of unintended pregnancy (AI).







Preconception Counseling and Care

- HIV infection does not preclude the use of any contraceptive method (AII). However, drug-drug interactions between hormonal contraceptives and cART should be taken into account.
 - Interactions between some ARVs and hormonal contraceptives may lower contraceptive efficacy.





Condoms

 The one method that protects against STDs and provides contraception

How do your patients feel about using male condoms? Female condoms?





Condoms

- However, 15% failure rate in preventing pregnancy
- Many couples (even serodiscordant couples=one partner HIV+ and one partner HIV-) use condoms off and on, rather than always
- So, a second method is recommended





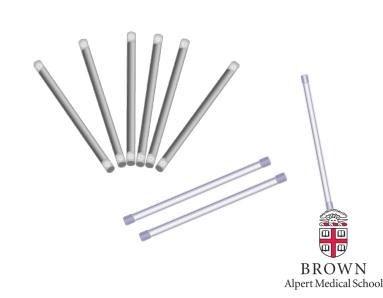


Hormonal Contraceptives

- Combined oral contraceptive pills (COCs)
- Progestin-only oral contraceptive pills (POPs)
- Injectables (Depo-Provera/DMPA)
- Implants (Norplant, Jadelle, Sinoplant, Implanon)







Drug Interactions between ARVs and Hormonal Contraceptives (CIII)

NNRTIs: See Guidelines Table 3

ARV Drug	Recommendation for Combined Hormonal Methods and Progestin- Only Pills	Recommendation for DMPA	Recommendation for Etonogestrel Implants
EFV	Use alternative or additional contraception	No additional contraceptive needed	Use alternative or additional contraception
ETR	No additional contraceptive needed	No additional contraceptive needed	No additional contraceptive needed
NVP	Consider alternative contraceptive, or barrier + oral hormonal methods	No additional contraceptive needed	Consider alternative contraceptive, or barrier + implant
RPV	No additional contraceptive needed	No additional contraceptive needed	No additional contraceptive needed

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Drug Interactions between ARVs and Hormonal Contraceptives (CIII)

RTV-Boosted Pls: See Guidelines Table 3

ARV Drug	Recommendation for Combined Hormonal Methods and Progestin- Only Pills	Recommendation for DMPA	Recommendation for Etonogestrel Implants
ATV/r	Use alternative or additional contraception	No additional contraceptive needed	Consider alternative contraceptive, or barrier + implant
DRV/r	Use alternative or additional contraception	No additional contraceptive needed	No additional contraceptive needed
FPV/r	Use alternative or additional contraception	No additional contraceptive needed	Consider alternative contraceptive, or barrier + implant
LPV/r	Use alternative or additional contraception	No additional contraceptive needed	Consider alternative contraceptive, or barrier + implant

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Drug Interactions between ARVs and Hormonal Contraceptives (CIII)

Pls without RTV: See Guidelines Table 3

ARV Drug	Recommendation for Combined Hormonal Methods and Progestin- Only Pills	Recommendation for DMPA	Recommendation for Etonogestrel Implants
ATV	No additional contraceptive needed •OC should contain ≤30 µg ethinyl estradiol (EE), or use alternative method •No data on OCs with <25 µg EE or progestins other than norethindrone or norgestimate	No additional contraceptive needed	No additional contraceptive needed
FPV	Use alternative contraception (use with estradiol/norethindrone may cause virologic failure)	No additional contraceptive needed	Use alternative or additional contraception





Drug Interactions between ARVs and Hormonal Contraceptives (CIII) (6)

Integrase Inhibitors: See Guidelines Table 3

ARV Drug	Recommendation for Combined Hormonal Methods and Progestin- Only Pills	Recommendation for DMPA	Recommendation for Etonogestrel Implants
RAL	No additional contraceptive needed	No additional contraceptive needed	No additional contraceptive needed
DTG			
EVG/COBI			

CCR5 Antagonist: See Guidelines Table 3

ARV Drug	Recommendation for Combined Hormonal Methods and Progestin-Only Pills	Recommendation for DMPA	Recommendation for Etonogestrel Implants
MVC	No additional contraceptive needed		





Other hormonal options

- □ Patch (Ortho Evra), vaginal ring (Nuva Ring), and transdermal implant (Implanon)
 - Warnings are similar to OCPs regarding drug-drug interactions
 - ■However, in theory, they avoid the "first pass" effect of liver metabolism that may occur with oral agents and should not be subject to the same limitations as OCPs





Intrauterine devices (IUDs)

- No known drug interactions
- No increase in shedding of HIV
- 2 types
 - Copper (Paragard) works for 10 years, may be associated with heavier menses, periods regular)
 - Levonorgestrel IUD (Mirena) works for 5 years, reduces menstrual blood loss (is FDA-approved as a treatment for menorrhagia), periods scant and not regular





Effectiveness of Contraception for HIV-Infected Women using Antiretroviral Therapy

Maria Pyra^{a,b}, Renee Heffron^{a,b}, Nelly R. Mugo^{b,d,e}, Kavita Nanda^f, Katherine K. Thomas^a, Connie Celum^{a,b,c}, Athena P. Kourtis^g, Jared M. Baeten^{a,b,c} for the Partners in Prevention HSV/HIV Transmission Study and Partners PrEP Study Teams

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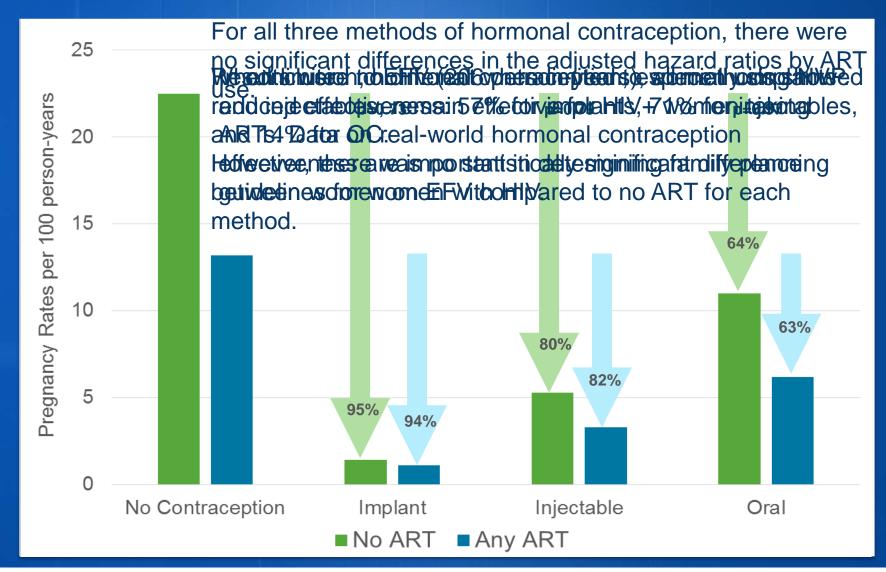
Question: Does effectiveness of hormonal contraception (implant, injectable, oral pill) differ by ART use?

- Sample: 5,153 women

 (1,376 pregnancies) in
 serodiscordant couples from 3
 prospective studies in Africa
 - Partners in Prevention HSV/HIV Transmission Study
 - Couples Observation Study
 - Partners PrEP Study
- Characteristics:
 - Young (median age 29), healthy (51% CD4≥500), and ART naïve at enrollment
 - Median follow-up 1.8 years
 - 24% became pregnant and 31% ever took ART

- Analysis: Cox proportional hazard models with repeated outcomes (pregnancy)
 - Tested interactions between ART use and contraceptive method

Results



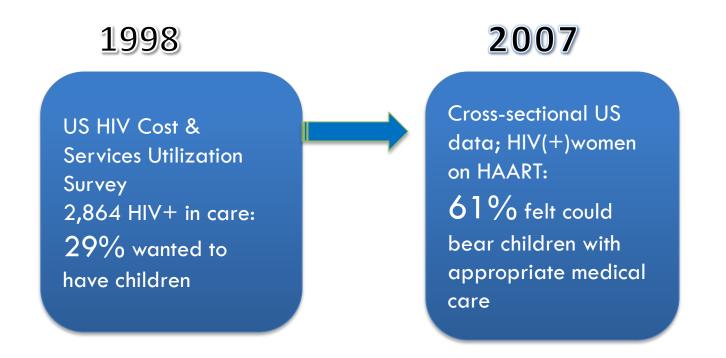


CONCEPTION





Fertility desires among HIV+ adults







What if both partners are HIV-positive?

 When a couple is not attempting conception, we recommend condoms to avoid superinfection and sharing of antiretroviral resistant virus

 If pregnancy desired: Ovulation predictor kit, maintaining an undetectable viral load, and once monthly unprotected sex is a reasonable approach





Reproductive Options for HIV-Concordant (both positive) and Serodiscordant Couples

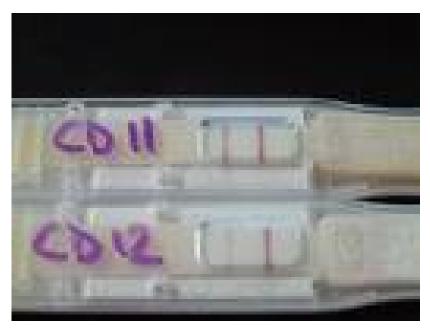
- Expert consultation is recommended so that approaches can be tailored to specific needs (AIII).
- Partners should be screened and treated for genital tract infections before attempting to conceive (AII).
- The HIV-infected partner should attain maximum viral suppression before attempting conception (AIII).





Ovulation predictor kits



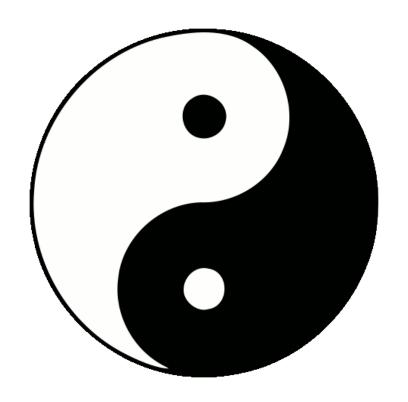


These replace the old basal body temperature charts





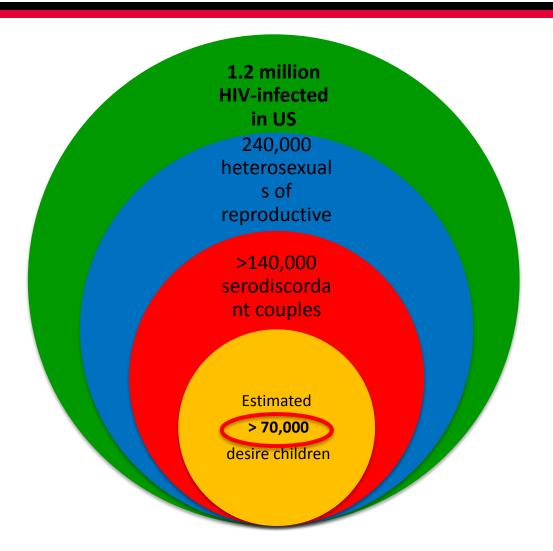
The Serodiscordant Couple







Serodiscordant Heterosexual Couples US 2006







Case 3-- Elizabeth

- ☐ Julia is 31, HIV+, diagnosed 1☐ Plan: after ending ago year relationship with an HIV-infected partner
- No history of HIV-related illness
- Not on HIV medications
- \square CD4 in the 600's
- □ VL is 8,000
- New partner is HIV-uninfected
- Seems anxious and upset

- - Discuss pros and cons of starting HIV treatment
 - Recommend HIV testing for partner
 - Reinforce the importance of using condoms.
 - Refer to a support group
 - Re-check her VL and CD4 in 3 months.
 - Continue to evaluate for and discuss HIV treatment





Case 3—Elizabeth...

- You ask Elizabeth if she wants to have another child.
 - She says, "Yes."
 - You ask, "When?"
 - She says, "Now."
- How do you counsel her?



Preconception counseling

- If a woman is not on ARVs, consider starting them prior to attempting conception
- If a woman is on ARVs and is considering pregnancy
 - Substitute other ARVs for efavirenz (Sustiva) because of possible risk of neural tube defects (NTDs)
 - Recommend folate or prenatal vitamins preconceptionally to reduce chance of NTDs





Discordant couples with HIV-infected women, HIV- man:

•The safest conception option is artificial insemination, including the option of self-insemination with a partner's sperm during the periovulatory period (AIII).





Serodiscordant couples

- If the woman is HIV+ and the man is HIV-, discuss the options of:
 - Ovulation predictor kits
 - Home insemination ("turkey baster method")





When the time is right, the choices are:

Home insemination with partner's semen

The "turkey baster" method

*A needle-less syringe works fine







HIV-infected Serodiscordant Couples are Willing to Have Unsafe Sex to Conceive

20% of couples seeking Assisted Reproduction Services reported engaging in unprotected intercourse to achieve pregnancy at some point in the past

Barreiro et al AIDS Rev 2006





Periconception Prep

- Very few data to date on periconception PrEP; studies under way.
- Infected partner should be on ART with fully suppressed HIV viral load.
- Once daily tenofovir/emtricitabine is currently FDA approved for PrEP; CDC recommends 1 month before and 1 month after conception attempted.
- Couples should use condoms at all times except during periovulatory intercourse.
- No reported increase in congenital anomalies for children whose mothers were exposed to tenofovir or emtricitabine during first trimester.





Case 4—Davis

- □32 year old HIV-positive male diagnosed with HIV 3 years ago,
- □On ARVs. CD4 882 and VL<20 (undetectable)
- ☐ Excited about plans to get married next month to a woman he's been dating for a year

- □Plan:
 - Refill medications
 - ■Counsel on use of condoms
 - ■Return in 6 months





Case 4—Davis

- ☐You ask Davis whether his fiancee has been tested for HIV
 - ■He says, "Yes, and she is HIV-negative."
- ☐You ask whether they are thinking about having children
 - He tells you, "Yes, sooner rather than later."
- ☐ How do you counsel him?





If the infected male has an undetectable plasma viral load, timed unprotected intercourse in order to conceive does not pose a risk to the negative female partner

- 1. TRUE
- 2. FALSE





Discordance of Genital and Plasma HVL

- Well-documented evidence that HIV RNA can be detected in genital secretions despite undetectable plasma HVL
- Prospective study, 25 men starting HAART, despite undetectable HVL plasma, 48% had intermittent shedding, no STIs
- 5% of 145 men on HAART seeking ART services found to have detectable genital HIV despite negative plasma HVL x 6 months, no documented STIs

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Discordant couples with HIV-infected men, HIV- woman:

- The use of donor sperm from an HIV-uninfected male with artificial insemination is the safest option (AIII).
- When the use of donor sperm is unacceptable, the use of sperm preparation techniques coupled with either intrauterine insemination or in vitro fertilization should be considered (AII).
- Semen analysis is recommended for HIV-infected males before conception is attempted to prevent unnecessary exposure to infectious genital fluid when the likelihood of getting pregnant is low because of semen abnormalities (AIII).





Accessibility

Prohibitive Advisories/Regulations

Cost Limitations





Nearly each state in the US has a fertility clinic that offers assisted reproduction services to HIV serodiscordant couples

- 1. TRUE
- 2. FALSE





- Accessibility
- 80% US Fertility Clinics NOT supporting services for HIVaffected couples (estimated 70,000 couples in US)
- Per Perinatal HIV Hotline only 6 fertility clinics in US offering sperm wash-IUI





Accessibility

Prohibitive Advisories/Regulations

- In 1990, CDC advised against sperm wash due to one isolated case of transmission when current standard sperm wash protocol not employed
- As of 2006 ASRM "endorses" sperm wash; reversing 1994 advisory against sperm wash
- As of 2014, DHHS Guidelines support use of sperm wash
- Some states still have prohibitive legislation; criminal penalties for using HIV-infected semen

Cost Limitations





Accessibility

Prohibitive Advisories/Regulations

Cost Limitations





Reproductive Options Comparison

Cost and HIV transmission risk by reproductive options					
Variable	IUI with SW	IVF with SW	ICSI with SW	Self-insemination	Intercourse
Average number of cycles ^{48,49}	2.8	1.4	0.6	5.5	5.5
Average cost/cycle					
Without PrEP ⁵⁰	\$1,265	\$ 12,513	\$15,128	\$30 (kit)	\$0
With PrEP ^{ab}	\$2,195	\$13,443	\$16,058	\$960	\$930
Average cost/live birth					
Without PrEP ⁵⁰	\$12,635	\$41,132	\$46,256	\$30	\$0
With PrEPab	\$16,835	\$42,062	\$47,156	\$5,145	\$5,115
Pregnancy rate/procedure 48,49c	19% ²⁹	38.1% ⁵¹	23% ²⁹	20%	20%51
Risk of HIV transmission ^d	0.1-0.5%52	0-0.4%	0-0.09%	0.03-0.14% ^{50,51} or 0.1-0.5% ⁵³	0.03-0.14% ^{50,8} or 0.1-0.5% ⁵³





Preexposure prophylaxis and timed intercourse for HIV-discordant couples willing to conceive a child

Pietro L. Vernazza^a, Irma Graf^b, Ulrike Sonnenberg-Schwan^c, Maria Geit^d and Anja Meurer^c

Many HIV-discordant couples express a strong wish to conceive a child. Insemination with processed semen is offered to these couples in many countries. Given the very low level of transmission risk during fully suppressive antiretroviral therapy, we offered timed intercourse combined with preexposure prophylaxis to further reduce the transmission risk. In 53 cases, natural conception was attempted using the proposed method. Pregnancy rates were high and reached a plateau of 75% after six cycles. Advanced age in the female partner was a predictor for infertility in these couples.

© 2011 Wolters Kluwer Health | Lippincott Williams & Wilkins

AIDS 2011, **25**:2005–2008





Prep and Timed Intercourse for Conception

- Male partner on HAART with undetetectable HIV-RNA in plasma (<50 copies/ml)
- No current symptoms of genital infections
- Urine LH-test to determine the optimal time of conception (36 h after LH-peak)
- Administration of PrEP (tenofovir po), First does at LH-peak-Second 24 hours later
- After 6 unsuccessful attempts, fertility evaluation recommended

(Vernazza et al, AIDS 2011)





PrEP for Conception

- Outcomes: March 2004 2007
 - 46 serodiscordant couples
 - 75% became pregnant: 50% after 3 or fewer attempts
 - 0 seroconversions or adverse events

Vernazza et al AIDS 2011;25:2005-2008





Periconception Prep

- Very few data to date on periconception PrEP; studies under way.
- Infected partner should be on ART with fully suppressed HIV viral load.
- Once daily tenofovir/emtricitabine is currently FDA approved for PrEP; CDC recommends 1 month before and 1 month after conception attempted.
- Couples should use condoms at all times except during periovulatory intercourse.
- No reported increase in congenital anomalies for children whose mothers were exposed to tenofovir or emtricitabine during first trimester.





ART DURING PREGNANCY





Principles of ARV Use during Pregnancy

Coordination of services among prenatal care providers, primary care and HIV specialty care providers, and when appropriate, mental health and drug abuse treatment services, and public assistance programs, is essential to ensure that infected women adhere to their ARV drug regimens (AIII).





General Principles of Drug Selection

- Guidelines for use of cART for maternal health during pregnancy generally are the same as for women who are not pregnant.
 - Some modifications based on concerns about specific ARVs during pregnancy and limited experience during pregnancy with newer ARVs.
- Ensure that at least 1 NRTI with high placental transfer is included in cART regimen for sufficient infant preexposure prophylaxis.
- Counsel women on the importance of close adherence to ARV regimen.
 - Offer support services, mental health services, smoking cessation, and drug abuse treatment plans as indicated.
- Coordinate between HIV and OB specialists.





Initial ART for ARV-Naive Pregnant Women (1) Preferred 2-NRTI Backbone Regimens

	Comments
ABC/3TC	 Available as FDC, can be given once daily Potential HSR: ABC should not be used in patients who test positive for HLA-B*5701 because of risk of hypersensitivity reaction Not recommended with ATV/r or with EFV if pretreatment HIV RNA >100,000 copies/mL
TDF/FTC or TDF + 3TC	 Available as FDC, can be given once daily TDF has potential renal toxicity, use with caution in patients with renal insufficiency
ZDV/3TC	 Most experience for use during pregnancy Available as FDC. Twice-daily administration Higher risk of hematologic toxicity

Initial ART for ARV-Naive Pregnant Women (2)

Preferred PI Regimens

	Comments
ATV/r + preferred 2-NRTI backbone	 Once daily administration Extensive experience in pregnancy Maternal hyperbilirubinemia
DRV/r + preferred 2-NRTI backbone	 Better tolerated than LPV/r. PK data available. Increasing experience in pregnancy Must be used twice-daily in pregnancy.





Initial ART for ARV-Naive Pregnant Women (3)

Preferred NNRTI Regimen

	Comments
EFV + preferred 2-NRTI backbone Note: May be initiated after the first 8 weeks of pregnancy	 Birth defects in primates; risk in humans is unclear. Postpartum contraception must be ensured. Preferred regimen in women requiring coadministration of drugs with significant interactions with PIs or the convenience of coformulated, single-tablet, oncedaily regimen.





Initial ART for ARV-Naive Pregnant Women (3)

Preferred Integrase Inhibitor Regimen

	Comments
RAL + preferred 2-NRTI backbone	 PK data available and increasing experience in pregnancy. Rapid viral load reduction. Useful when drug interactions with PI regimens are a concern. Twice-daily dosing required.





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Initial ART for ARV-Naive Pregnant Women (5)

Alternative PI Regimens

	Comments
LPV/r + preferred 2-NRTI backbone	 Abundant experience and established PK in pregnancy. More nausea than preferred agents. Twice-daily administration. Once-daily LPV/r is not recommended for use in pregnant women.





Initial ART for ARV-Naive Pregnant Women (6) Alternative NNRTI Regimen

	Comments
RPV + preferred 2- NRTI backbone	 RPV not recommended with pretreat-ment HIV RNA >100,000 copies/mL or CD4 cell count <200 cells/mm³. Do not use with PPIs. PK data available in pregnancy but relatively little experience with use in pregnancy. Available in co-formulated single-pill once daily regimen.





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Initial ART for ARV-Naive Pregnant Women (8)

Insufficient Data to Recommend Routine Use

DTG

EVG/COBI/TDF/FTC FDC

FPV

MVC

RPV

Not Recommended		
ABC/3TC/ZDV	SQV	
d4T	ETR	
ddI	NVP	
IDV/r	T-20	
NFV	TPV/r	
RTV as single PI		





HIV-Infected Pregnant Women Who Are Currently Receiving Antiretroviral Therapy (1)

- In general, HIV-infected pregnant women receiving cART who present for care in the 1st trimester should continue treatment during pregnancy, assuming the regimen is tolerated and effective in suppressing viral replication (HIV-1 viral load less than lower limits of detection of the assay) (AII).
- It is recommended that efavirenz be continued in pregnant women receiving efavirenz-based cART who present for antenatal care in the first trimester provided the regimen is achieving virologic suppression (CIII).





Maternal and Fetal Monitoring during Pregnancy (2)

Monitor HIV RNA:

- At the initial visit (AI)
- 2-4 weeks after initiating or changing ARV drug regimens (BI)
- Monthly until HIV RNA is undetectable (BIII)
- At least every 3 months during pregnancy (BIII)
- HIV RNA should also be assessed at approximately 34-36 weeks' gestation to inform decisions about mode of delivery and about infant ARV prophylaxis (AIII).









POSTPARTUM CARE





Postpartum Care

- Because the immediate postpartum period poses unique challenges to antiretroviral adherence, arrangements for new or continued supportive services should be made before hospital discharge for women continuing cART (AII).
 - Counsel women about the challenge of adherence in the postpartum period.
 - Remain vigilant for signs of depression, intimate partner violence, and drug or alcohol use.
 - Consider simplifying cART regimens to improve adherence.







Background

- Postpartum HIV-infected women face challenges with treatment adherence.
- Women in the deep South may experience greater difficulties with care engagement due to poor access to care, stigma, lack of social support, and mistrust in the health care system.

- 1.Bardeguez et al. Adherence to antiretrovirals among US women during and after pregnancy. JAIDS 2008; 48:408.
- 2. Watts DH et al. Progression of HIV disease among women following delivery. JAIDS 2003; 33:585–93
- 3. Postpartum viral load rebound in HIV-1 infected women treated with HAART. HIV Clin Trials 2011; 12:9-23
- 4. Sex, race, and geographic region influence clinical outcomes following primary HIV-1 infection. JID Feb 15 2011;203(4):442-451
- 5.Reif S. et al, HIV Diagnoses, Prevalence and Outcomes in Nine Southern States, 39(6) J. Comm. Health (2015) 40:642–651







Background

 A growing number of HIV-infected women are giving birth every year.

 Pregnancy provides a unique opportunity to impact the HIV Treatment Adherence Cascade

Whitmore SK, et al. Estimated Number of Infants Born to HIV-Infected Women in the United States and Five Dependent Areas, 2006. *JAIDS* 2011.

Lando HA et al. Promoting smoking abstinence in pregnant and postpartum patients: a comparison of 2 approaches. Am J Manag Care. Jul 2001;7(7):685-693.

Matthey S, et al. Prevention of postnatal distress or depression: an evaluation of an intervention at preparation for parenthood classes. J Affect Disord. Apr 2004;79):113- 126. Wodak A et al. Evaluation of a cognitive-behavioural intervention for pregnant injecting drug users at risk of HIV infection. Addiction. Aug 1996;91(8):1115-1125.



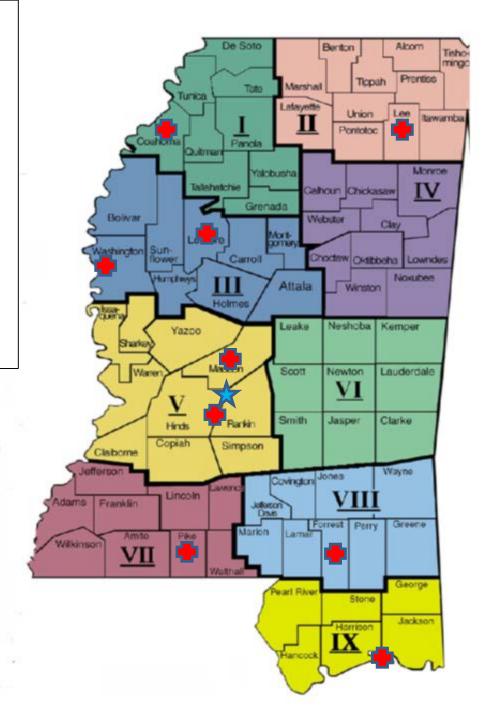


HIV in Mississippi, 2013

- PLWH: 10,473
 - 68% men, 32% women
- HIV Incidence: 536
 - 24.8% female (82% HS, 16% IDU)
 - AA women case rate 9x
 White women
- 9 Ryan White Funded Clinics

Mississippi Public Health Districts I-IX







Objectives

- Retrospective analysis of all HIV-infected women ≥16 years who delivered in Mississippi from January 1, 2002 to Dec 31, 2014.
- Focus on health care utilization and outcomes:
 - Death/Progression to AIDS
 - Engaged in care in 2015 (one medical visit or CD4/PVL in 2015)
 - HIV-1 Plasma Viral Load <200 copies/mL in 2015</p>







Methods

- Clinical data from all 9 federally funded Ryan White clinics in Mississippi (Careware)
 - Statewide implementation in 2005-2006
- Mississippi Department of Health (MSDH) Enhanced HIV/AIDS Reporting System (eHARS)
 - Mandatory CD4/HIV Viral Load reporting to MSDH started Jan 2013





Demographics

•	
<u>Total Women</u>	548
Total number of deliveries	685
Median Age at First Delivery (IQR)	26 (23,31)
Race Black White Multiple AI/AN Not reported	474 (86.5%) 57 (10.4%) 4 (0.7%) 3(0.5%) 10 (1.8%)
<u>Hispanic</u>	15 (2.7%)
Median Annual Income (IQR) (n=208)	\$9780 (\$4116, \$15570)
Insurance Medicaid* Uninsured Private Medicare Unknown	123 (22.4%) 65 (11.8%) 20 (3.6%) 9 (1.6%) 217 (39.6%)
Housing Status Stable/Permanent Temporary/Unstable	192 (35%) 23 (4.2%)

BOST Unknown

N chool

215 (39.2%)



Geography

Health District (N=300)	
I	10 (3.3%)
II	6 (2%)
III	36 (12%)
IV*	12 (4%)
V	134 (44.7%)
VI*	27(9%)
VII	23 (7.7%)
VIII	31 (10.3%)
IX	21 (7%)
Current State of Residence eHARS (N=415)	
Mississippi	383 (92.3%)





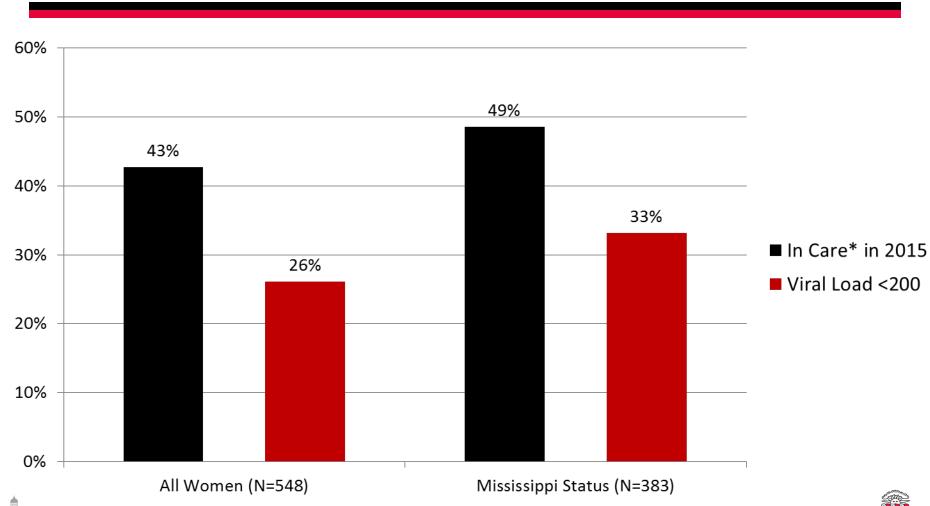


HIV

Median Age at HIV Dx (IQR) (n=548)	22 (19, 27)
HIV Risk	
Heterosexual	395 (72.1%)
Perinatal	15 (2.7%)
IDU	13 (2.4%)
Unknown	125 (22.8%)
HIV dx around pregnancy	206 (37%)
AIDS Diagnosis	268 (48.9%)
Median Age at AIDS (IQR) (n=268)	28 (23, 32)
AIDS within 1 year of HIV dx	68 (13%)
Median Time HIV to AIDS, years (n=268)	4.67 (.91, 8.3)
Median Last available CD4 cells/μL (IQR)	494 (305, 695)
Most recent HIV-1 PVL	
<200 copies/mL	146 (26.7%)
>200 copies/mL	228 (41.6%)
Missing	174 (31.8%)
Perinatal Transmission	9 (1.3%)

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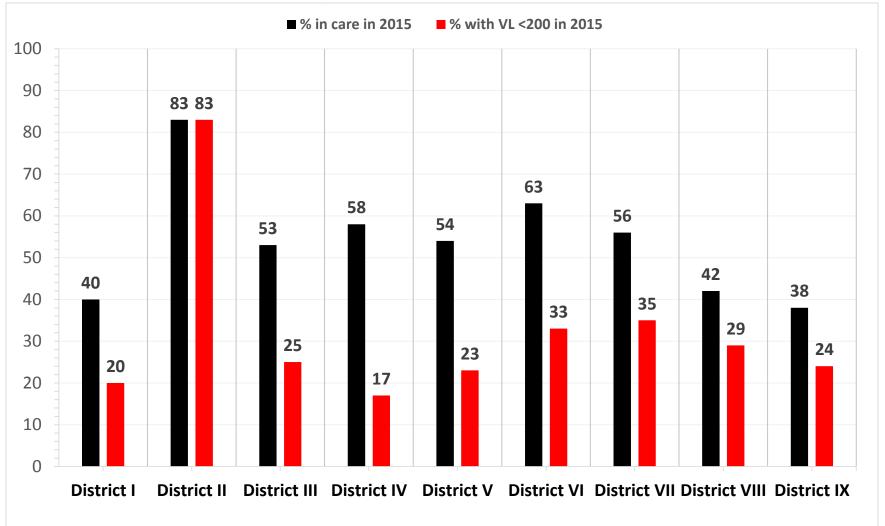
Engagement in 2015







Outcomes by Health District in 2015









Mortality

Number of Deaths	67 (12.2%)
Median Age at death (IQR)	32.4 (28.1, 36.7)
Median Time from HIV dx to death, years (IQR)	9.3 (6, 14.7)
Median Time from AIDs dx to death, years	4.5 (2.1-7.9)
Median time from last delivery to death, years (IQR)	5.35 (3.0, 7.0)
Median last available CD4 cells/μL (IQR) n=60	38 (9, 133)
Median last available HIV PVL copies/mL (IQR) n=58	59220 (7713, 195137)







Conclusions

- Young, HIV infected women in Mississippi experience low rates of retention and viral suppression, and significant morbidity and mortality following delivery.
- Systems based and innovative interventions initiated during pregnancy and continued through postpartum phase to support engagement with care may improve longitudinal treatment adherence and health outcomes.
- Interventions should be developed in collaboration with target health districts with lowest rates of care engagement.





Next Steps

- Analysis of predictors of retention, viral suppression and AIDS/mortality
- Cause of death
- GIS mapping (census tract data, health districts)
- Prospective study of HIV-infected pregnant and postpartum women
 - Followed longitudinally over 2 year period
 - Assessments of structural and behavioral barriers to care





Websites to Access the Guidelines

- http://www.aidsetc.org
 - http://aidsinfo.nih.gov











QUESTIONS??



