Special Recommendations in High-Risk Prenatal Care

PART ONE

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2025 COLORADO PERINATAL SUBSTANCE USE DISORDER INTEGRATION CONFERENCE, AVON, CO







Disclosures

None



Disclaimers

Not ALL of the following recommendations have formal references or society recommendations.

There is still a significant amount of prenatal care that is based on historical delivery and/or local standard of care.

For High-Risk Pregnancy that standard is typically set by local Maternal Fetal Medicine (MFM) recommendations.

Objectives

- Briefly review evidence-based, routine prenatal care updates
- Work through recommended additions to care based on common pre-existing medical conditions such as:
 - > Diabetes
 - > Chronic HTN
 - Obesity and s/p Bariatric Surgery
 - > Thyroid Disease

Routine Prenatal Care

	PNC				<u>Fu Labs</u>	Antental Testing			<u>Delivery</u>	RISK Factors
	Labs	US Plan	Medication	Referral		Start	Frequ	Туре	<u>Plan</u>	
Routine	Prenatal Panal (CBC, Bld Type, Rh, Antibody screen, rubella, RPR, Hep B), Varicella, HIV, GC/CL, A1C, Hep C, Urine (cx, UDM). Pap if indicated.	Dating Anatomic (18-22 wks)	PNV	N/A	1 hr GTT at 24-28 wks	41 wk (39 wk if first US ≥ 22 wks)	twice weekly	mBPP (NST+AFI)	Recommend IOL at 41 wks, test if 41-42 weeks, refer to MFM by 42 wks if pt refuses IOL	for late term: stillbirth, macrosomia, CPD

Routine Prenatal Care

Labs

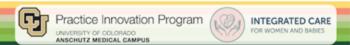
- CBC
- Blood Type, Rh, & Antibody
- Rubella
- RPR vs TPA
- Hep B
- HIV
- Urine Cx
- Gc/Ct

Often Missed "NEW" Additions

- Hep C
- Varicella
- A1C

What is not here?

- Urine Toxicology
- TSH
- Pap



Routine Prenatal Care

Ultrasounds

- Dating
- Screening Anatomic

Screenings

- Mental Health
 - PHQ9, GAD7, CIDI
- Substance Use
 - o 5Ps
- IPV USPSTF B
 - E-HITS

Other Labs

GTT

RPP at 28 wks & Delivery* ACOG PA 2024

Antenatal Testing if PostDates

2x/wk mBPP at 41 wk (*39 wks if first US >22 wks)

Referral

MFM by 42 wks if IOL refused



A little more information . . .

Rh status is NOT antibody status!

No longer indicated for 1st trimester bleeding Dec 2024 ACOG CPU

TPA is the preferred initial screen for Syphilis. If +, an RPR w/ quant is performed and TPPA

Hepatitis - Sept 2023 CPG

- HBsAg universal (SR, MQ)
 Triple Panel screening (HBsAg, anti-HBs, and total anti-HBc) (SR, LQ)
 If > 18 y/o and don't have the above documented as Anyone who hasn't completed the vaccine series
 Those w/ ongoing risks of infection regardless of testing hx or vaccination status
 HepC virus antibodies (SR, LQ)

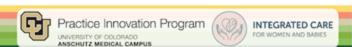
Varicella

A1C - Risk-based (*USPSTF*)

Obesityhx of gDM or DM

family hx of gDM or DM

Ethnicity (Hispanic, Native American, South or East Asian, African American, Pacific Islander)



And now . . . the fun stuff





DIABETES

		PNC			Fu Labs	Ar	ntental Tes	ting	Delivery	RISK Factors
	Labs	US Plan	Medication	Referral		Start	Frequ	Туре	Plan	ALCOHOLOGICA CONTROL C
hx gestDM	A1C on intake screens for pregestational diabetes	routine	routine	N/A	Routine 1 hr	Routine	Routine	Routine	Routine	70% w/ gDM have DMII w/i 20 years, latino> 60% develop w/i 5 yr
A1 gDM (diet controlled)	n/a	n/a	n/a	N/A	check 2hr	Routine (ur	nless other ris treat like A2	k factor then	Routine	
A2 gDM (well controlled)	n/a	n/a	metformin vs insulin	N/A	GTT at	32 wk 32 wk	q 4 wk weekly	growth US NST	39w0d - 39w6d	
A2 gDM (poorly controlled)	n/a	n/a	insulin vs metformin	MFM ENDO	PP	32 wk 32 wk	q 4 wk 2x/wk	growth US NST	37w0d - 38w6d (per MFM)	
preexist PRE-DM		n/a	metformin?	nutrition	early 1 hr	-	-	-	routine	Increased congenital malformations
preexist Type 2 - controlled (A1C ≤ 7)	CMP, 24 hr urine	Dating . Anatomic (per MFM)	ASA* Cont metformin if already on & controlled	MFM Endo Optho q trm	repeat	28 wk 32 wk	q4 wk 2x/wk	growth US NST	39w0d - 39w6d	(2-6x), cns malform 10x inc., 5x inc in heart malf; fetal macrosomia, fetal
preexist Type 2 - Uncontrolled (A1C > 7)	CMP, 24 hr urine, prot:cr	Dating Anatomic @ MFM (+ fetal echo)	ASA* . intensify insulin therapy	MFM Endo Optho q trm TRANFER CARE	cMP, 24 hr urine and prot/cr	28 wk 28 wk	q 4 week 2x/wk	growth US @ MFM NST	per MFM	hypoglycemia, fetal RDS
Type 1	CMP, 24 hr urine, prot/cr ratio, TSH	Dating . Anatomic @ MFM (+ fetal echo)	ASA*	MFM Endo Optho q trm <u>Transfer based</u> on control	ratio q 1-3 months	28 wk 28 wk	q 4 week 2x/wk	growth US @ MFM NST	per MFM	

A little more information . . .

- May 2024 American Journal of Perinatology

 "Even in patients with multiple risk factors, early screening for GDM does not improve outcomes."

Early Glucose Tolerance Testing Reminders 50g (1 hr) at the time of diagnosis If NORMAL - repeat at 24-28 weeks

- If Elevated → 100g (3 hr)

 If Normal → Repeat ONLY the 3 HR at 24-28 wk

Post Partum

75g (2hr) GTT at 4-12 wks due to high incidence of Type 2 DM

Hyperglycemia

- Hyperglycemia can be teratogenic, especially prior to 10 wks gestation
 Hospitalization is indicated for IMMEDIATE glycemic control w/ insulin drip if A1C > 9

 Detailed anatomic US by MFM w/ fetal echo

"Control"

- A1C goal < 7
- Fasting (<95), 1 hr PP (<140) OR 2 hr PP (120), HS (<110)

 Looking for 90+% to be in range each wk

 Rapid advancement of medical management due to potential impact on fetal growth daily

Aspirin

ACOG CO update 2023

Recommended if high risk for preeclampsia (or multiple moderate risk factors)

HIGH

★ Hx of preE

★ multigestation

★ cHTN

★ DM

★ CKD

★ Al

MODERATE

★ Nulliparity

★ Obesity

★ Fam Hx preE

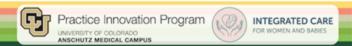
★ AMA

★ hx of LBW/SGA, adverse preg, >10 year interval

★ Sociodemographic characteristics

81mg PO daily starting at 12-28 wks (< 16 wk optimal)

NOT indicated for solely: stillbirth, FGR, PTB, EPL



keeping it going





HYPERTENSION

ACOG PBs 2019/2020

		PNC					Antental Testing			RISK Factors
	Labs	US Plan	Medication	Referral		Start	Frequ	Туре	<u>Plan</u>	
HTN				,			,	continue A	ASA until delivery	
Chronic	CMP, prot:cr, 24 hr urine protein	Dating Anatomic by MFM	ASA*	MFM	Check BP 1-2 wks PP	32 wk 28-32 wk	twice weekly monthly	mBPP growth US	38w0d	superimp preE (14-28%)>plac abruption, FGR, PTD.
hx preE or HELLP	CMP, prot:cr, 24 urine protein	Routine	ASA*	MFM if + hx in ≥ 2 pregs	yearly post partum BP, lipids,				Routine	21% chance of having in sub preg. -Increased risk for: PTL, FGR, abruption, fetal demise
pre E w/o SF & gHTN	N/A	N/A			FBG, & BMI	32 wk @ dx @ dx	2x/wk weekly 2x/wk	mBPP Plt/LFT/ur pro office BP's	37w0d	OP management ok b/w 32 and 37 wks if BP < 155/105

A little more information . . .

Pre Pregnancy Eval

End organ evolvement (level C) Secondary (11-14%)

Optimize comorbidities

Optimize control

Medication Review

AVOID ACE inhibitors and ARBs

fetogenic (renal dysgenesis, calvarial hypoplasia, FGR)

Superimposed preE on cHTN
 Occurs in 20-50% of patients
 5x higher than baseline population

Difficult to diagnose

sudden increase in baseline or proteinuria

preE labs

Eval for Heart Disease

Poor control > 4 yrs, > 30 y/o EKG +/- Echo

"Control"

No benefit to tight control

Initiate at 160/110* (level B)
Labetalol and Nifedipine are 1st line (level B)

Delivery

cHTN 38 wks+ if not on medication (37 wks+ if on meds) *level A* at 37 wks if superimposed or preE/gHTN w/o severe features *level A* 34 wks + if superimposed preE w/ severe features *level B*

Box 1. Risks of Chronic Hypertension in Pregnancy

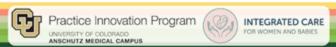
Maternal

- Death
- Stroke
- Pulmonary edema
- Myocardial infarction
- Preeclampsia
- Placental abruption
- Cesarean delivery
- · Postpartum hemorrhage
- Gestational diabetes

Fetal and Neonatal

- · Stillbirth or perinatal death
- · Growth restriction
- Preterm birth
- Renal insufficiency and failure • Congenital anomalies (eg, heart defects, hypospadias, esophageal atresia)





another doozy





Obesity

ACOG PB June 2021/2025

		PNC			Antental Testing Fu Labs			Delivery	RISK Factors	
	Labs	US Plan	Medication	Referral	1 4 5 4 5	Start	Frequ	Туре	Plan	NISK TUCKOTS
Obesity									umenable	
pregravid BMI 30-35		routine		routine		26 wk	q 6 wk	growth US	Routine	
pgBMI 35 - 40		Anatomic?		MFM for imaging based on shape		26 wk 36 wk	q 4 wk weekly	growth US BPP		DM, gDM, preE, medically indicated PT delivery, twin, post date gestation, OSA, labor dystocia, CSX, macrosomia, congen abnml, stillbirth, VTE, PP infection
pgBMI 40-45	CMP, prot:creat	Detailed Anatomic + fetal echo @MFM	if > 2 moderate risk factors for preE	MFM for imaging and		26 wk 34 wk	q 4 wk weekly	growth US BPP	Consider PP pharmacologic DVT prophylaxis based on risk;	
BMI > 45		Detailed Anatomic + fetal echo @MFM		TRANSFER CARE		26 wk 34 wk	q 4 wk weekly	growth US (@ MFM) . BPP	use Lovenox Post CSX	
s/p Bariatric Surgery	CBC, CMP, PTH, Iron, ferritin, folate, Vit A, Vit B12, Vit D	MFM (if surgery within the past 12-24 months)	PNV . Calcium citrate 1200 mg . Ferrous Sulfate 325 mg QOD	Nutrition referral Bariatric surgery if any concerns	CBC, BMP, iron, ferritin, folate Ca, & Vit D q trimester	28 wk	q 4 wk	growth US (if surg was w/i last 12-24m)	Routine	N/V, electrolyte derrangement, dumping syndrome FGR (if surgery within 12-24 months, cesarean

A little more information . . .

- Level A

 - PreGravid BMI should guide care
 SubQ drain should not be used routinely after csx
 Diet and Excercise better than exercise alone
- Level B

 - Encourage wt loss pre-pregnancy*
 Allowing longer 1st stage is reasonable to reduce csx
 Thromboprophylaxis
- Level C
 - US limitations
- early glucose screen
 Weekly antenatal testing
 by 37 wks if pgBMI 35.0-39.9
 at 34 wks if pgBMI > 40
- So what about growth Ultrasounds?

 2022 AJOG lower rates of FGR/SGA in obesity
 2024 AJP Consider delaying until 32 wks, limited utility
 - But . . .

last one (for this session)





Thyroid Disease

ACOG PB 2020

		PNC			Fu Labs	An	tental Tes	ting	Delivery	RISK Factors
	Labs	US Plan	Medication	Referral	1 G LGDS	Start	Frequ	Туре	Plan	NISK Tuctors
Нуро	TSH . (if decide to check TSH and NEW dx made, initiate treatment for TSH > 4)	Routine (unless TSH was > 10 in pregnancy then Anatomic by MFM)	Increase levothyrozine by 20-30% of home dose at dx of pregnancy.	n/a unless uncontrolled	TSH q 4 wk post change & atleast 1x per trim. (goal<2.5) TSH 6wk PP	Routine	Routine	Routine	Reduce to original dose after delivery Routine	Miscarriage, neonatal neurodevelopment issues if TSH >10, PPH, postpartum depression, lower APGARS, preeclampsia, preterm birth, abruption, FGR (if untreated
HYPER	TSH, FT4	antatomic @ MFM	PTU in first trimester, methimazole in second and third	Endo MFM <u>TRANSFER</u> <u>CARE</u>	FT4 q2 wks until controlled , then q2-4 wks. Goal FT4 is upper limit of nml	After 20 wk	q 4 wk	growth US	l	Miscarriage, hypertensive disorder, preterm birth, abruption, low birthweight, FGR, neonoatal hypo- and hyperthyroidism

A little more information . . .

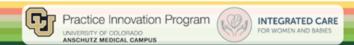
Level A

- Universal screening is NOT recommendedTSH is firstline

- Hypothyroidism
 Monitor q 4-6 wks
 treatment goal of TSH <2.5
 HYPERthyroidism
 Monitor free T4 (sometimes T3), goal upper limit of normal

Level B

- Use PTU or methimazole for treatment based on trimester, etc
- Level C
 - Continue to test as clnically indicated (symptoms, personal or family) hx, DM 1)
 - Thyroid tésting NOT indicated in hyperemesis



Part 1 Conclusion

- Not everyone needs a TSH or screening with urine toxicology
- Make sure to add in Hep C, Varicella, and triple Hep B
- Strongly evaluate need for A1C and ASA (indicated more often than not)
- Glycemic control is crucial to improved outcomes in diabetes
- If nothing else the presentation should have convinced you that preconception counseling and optimization of chronic health conditions is crucial

QUESTIONS?



Special Recommendations in High-Risk Prenatal Care

PART TWO

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Disclosures

None



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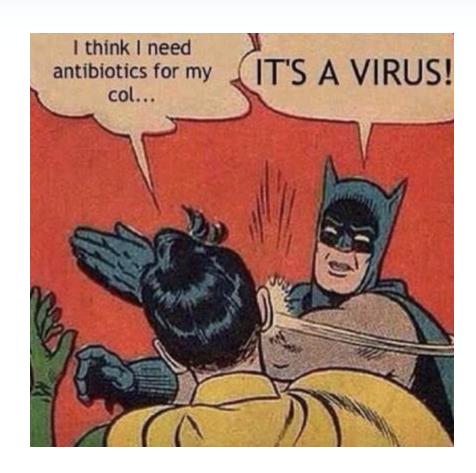
There is still a significant amount of prenatal care that is based on historical delivery and/or local standard of care.

For High-Risk Pregnancy that standard is typically set by local Maternal Fetal Medicine (MFM) recommendations.

Objectives

- Work through recommended additions to routine prenatal care based on:
 - > Infections
 - Maternal Age at Delivery
 - > Incarceration
 - Substance Use Disorder
 - > Tobacco Use Disorder
- A look at the whole High-Risk Obstetrics Grid

starting big





Infections - pt 1

		PNC			Fu Labs	An	tental Tes	ting	Delivery	RISK Factors	
	Labs	US Plan	Medication	_	T G EGDS	Start	Frequ	Туре	<u>Plan</u>	NISK Tuctors	
Infections				and/or ivirivi							
HIV	CMP, TB, toxo, Trich, hep A, viral load (plasma HIV RNA) Hep A, B, & pneumocococal vaccines	Date ASAP . Detailed anatomic @ MFM	initiate or cont ART (earlier better, triple therapy, AVOID stab/didan/riton/ covicistat) Call 1-888-448-8765	TRANSFER CARE to OB and/or MFM + ID consult	HIV RNA atleast monthly, CBC/CMP q 3m, CD4 counts prn	28 wk	q4 wk	growth US	routine if supressed 38 week CSX if viral load > 1000	fetal transmission, FGR; co-infection> FGR, PTL BFing NOT rec'd in US	
Syphillis	TPA>RPR quant if + (TPPA if RPR -)	Dating . Anatomic (@ MFM if high titers, late tc, untrt)	PCN (1 vs 3 doses)	MFM?, HD	f/u titers at 30 wks, delivery, pp (3,6,12)	28 wk 36 wk	q 4 wk weekly	growth US mBPP * (for uncont)	routine (IOL for perinatology based on fetal US)	FGR, stillbirth, congenital syphillis	acog pa april 2024
Нер В	DNA quant, CMP	routine	Tenofovir df if load > 200,000	Consider GI/Hepatology	viral load prior to treatment	28 wk	q 4 wk	growth US	routine . baby (HBV & HBIG w/i 12 hrs), BFing encouraged	pp viral flare, 90% vert transmit if not treated	acog cpg 6 sept 2023
Нер С	HCV RNA quant, CMP	routine	Defer to after pregnancy	Consider GI/Hepatology	viral load prior to treatment	28 wk	q 4 wk	growth US	routine (VD and internals not CI even w/ high load), BFing not discouraged	cholestasis, 3-10%	acog cpg 6 sept 2023

A little more information . . .

- Early Universal screening for HBsAg (SR,MQ)
- Triple Panel (HBsAg, anti-HBs, total anti HBc) if never documented after age 18, ongoing risks, or not fully vaccinated (SR, LQ)
- Hep C screen w/ each pregnancy (SR, LQ)
- Screen and treat HepC pre pregnancy when able (SR, MQ)
- Use HepB DNA quant to guide therapy (SR, LQ) and treat > 200,000 viral load to decrease vertical transmission (SR, MQ)
- CSX based on obstetric indication not dx of HepB (LQ)
- HBIG and HepB Vaccine w/i 12 hrs when indicated (SR, MQ)
- Manage healthcare exposures to Hep B/C similar to if not pregnant (SR, LQ)
- Hep A and Hep B vaccination safe in pregnancy use as indicated (SR, MQ)

Infections - pt 2

		PNC			Fu Labs	An	tental Tes	ting	Delivery	RISK Factors	
	Labs	US Plan	Medication	Referral	T G EGDS	Start	Frequ	Туре	<u>Plan</u>	NISK Tuctors	
Pyelo	routine	routine	IV then PO to total 14d . consider proph abx	Consider Uroloy	insuf evidence	routine	routine	routine	routine unless active infx	PTB, recurrent infx (septic shock, ARDs, stillbirth)	acog cc #4 aug 2023
Gc/Ct	consider Trich	routine	Abx as indicated	n/a	repeat per trimester	routine	routine	routine	routine		
Varicella	routine	Anatomic by MFM for primary infx in early pregnancy	Exp: VZIG w/i 96 hrs if VzNI Infx: Acyclovir 800mg 5x/d	MFM	routine	28 wk	q 4 wk	growth US	routine unless otherwise indicated	FGR, fetal hydrops, heart malform	acog PB 151, june 2015
HSV	routine	routine	Acyclovir 400mg po TID at 36 wk	n/a	none	routine	routine	routine	routine (csx w/ active GENITAL lesions)	vertical transmission	acog PB 220, may 2020

always fun to address in "nice" terms





Age at Delivery

ACOG OCC Aug 2022

		PNC			Fu Labs	An	Antental Testing			RISK Factors
	Labs	US Plan	Medication	Referral	T G EGOS	Start	Frequ	Туре	<u>Plan</u>	NION Factors
Teen	Routine	Routine	Routine	N/A	N/A	30 wk	once then prn	growth US	Routine	PTL, (LBW infants, malnutrition, PP depression, preE, IUGR).
AMA										
35-39 y/o (at delivery)		Dating . Consider Anatomic @ MFM	consider ASA if > 2 moderate risk	prn NT US	N/A	28-32 wk 38 wk	once twice weekly	growth US mBPP	by 40w0d	Ectopic, spon Abortion, stillbirth, chromosomal abnml, some congen
≥ 40 y/o	(cell free DNA is first line)	Dating Anatomic @ MFM	factors for preE	MFM by 20 wk (sooner prn NT US)	N/A	28-32 wk 34 wk	once twice weekly	growth US mBPP	at 39w0d	abnml, placenta previa, gDM, preE, CSX, (PT birth)

A little more information . . .

Level A

Prenatal genetic screening and diagnostic testing options should be discussed and offered to all individuals regardless of age or risk

Level B

- Daily low dose ASA if 2 mod risks (ama already one)
 antenatal fetal surveillance if delivery > 40 y/o 2/2 stillbirth risk
 Deliver 39 39.6 for > 40 y/o
 not an indication for csx

- be aware of disproportionate rate of adverse outcomes in Black,
 American Indian, and Alaska Native pregnant individuals ages 35 and older

Level C

- Delivery at > 35 y/o is high risk and should be discussed vague
 1st trimester US 2/2 high risk of multiples
 Detailed anatomic US if >35 yo
 Growth US in 3rd trimester if delivery at > 40yo

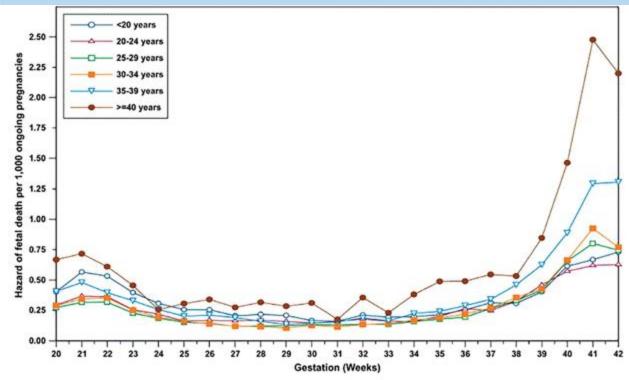


Figure 1. Risk of Fetal Death per 1,000 Ongoing Pregnancies by Week of Gestation. Reprinted from Reddy UM, Ko CW, Willinger M. Maternal age and the risk of stillbirth throughout pregnancy in the United States. Am J Obstet Gynecol 2006;195:764–70. doi: 10.1016/j.ajog.2006.06.019. Copyright 2006, with permission from Elsevier.

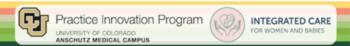
Term*						
	Trisomy 21	Trisomy 18	Trisomy 13	Sex Chromosome Aneuploidy (XXX, XY, XYY, 45, X)	Microarray or Rare Chromosomal Abnormality	All Chromosomal Abnormalities
Age 20	8 per 10,000 1 in 1,250	2 per 10,000 1 in 5,000	1 per 10,000 1 in 10,000	34 per 10,000 1 in 294	37 per 10,000 1 in 270	82 per 10,000 1 in 122
Age 25	10 per 10,000	2 per 10,000	1 per 10,000	34 per 10,000	37 per 10,000	84 per 10,000

Table 1. Chromosomal Abnormalities in Second-Trimester Pregnancies Based on Maternal Age at

1 in 1,000 1 in 5,000 1 in 10,000 1 in 270 1 in 294 1 in 119 Age 30 | 14 per 10,000 4 per 10,000 2 per 10,000 34 per 10,000 37 per 10,000 91 per 10,000 1 in 2,500 1 in 5,000 1 in 270 1 in 110 1 in 714 1 in 294 Age 35 34 per 10,000 9 per 10,000 4 per 10,000 35 per 10,000 37 per 10,000 119 per 10,000 1 in 1,111 1 in 2,500 1 in 270 1 in 84 1 in 294 1 in 285 Age 40 | 116 per 10,000 30 per 10,000 14 per 10,000 | 51 per 10,000 37 per 10,000 248 per 10,000 1 in 86 1 in 333 1 in 714 1 in 196 1 in 270 1 in 40

*Not all chromosomal abnormalities increase as maternal age increases.

2025 COLORADO PERINATAL SUBSTANCE USE DISORDER INTEGRATION CONFERENCE, AVON, CO



scraping the surface of why we are all here this weekend





Additional Risk Factors

		PNC			Fu Labs	An	tental Tes	ting	Delivery	RISK Factors
	Labs	US Plan	Medication	Referral	Tu Labs	Start	Frequ	Туре	<u>Plan</u>	KISK Factors
Incarceration	Trichamonas, TB? (Pap/HPV, follow nml guidelines) Hep A Vac (Hep B if not prior)	routine	routine	N/A	Repeat Syphillis, HepB, HepC, HIV, Gc/Ct, & trich screening in 3rd trimester	Routine	Routine	Routine	after 39w0d, consider social IOL	PTL, LBW. (poor PNC, SA, low SE status, current or prior trauma, MH issues, STIs, Resp illness(flu and tdap extra important), hyperemesis increased due to poor diet)
Tobacco Use	Routine	routine	cessation aids prn	N/A	Routine	30 wk	once then prn	growth US	Routine	Placenta previa, abruption, PROM, PTL, FGR, SIDS, birth defects, childhood obesity (3)
Substance Use	CMP Hep A Vac (Hep B if not prior)	routine	ASA MOUD	ICWB, Peer Support, BH, OTP prn	UDM?	30 wk	once then prn	growth US	after 39w0d, consider social IOL	PTL, FGR, inadequate prenatal care

A lot of OLD information

- Reproductive Health Care for Incarcerated Pregnant, Postpartum, and Nonpregnant Individuals - CO 2021 (2024)
- Alcohol Abuse and Other Substance Use Disorders: Ethical Issues
 CO 2015 (2021)
- Marijuana Use During Pregnancy and Lactation CO 2017 (2021)
- Tobacco and Nicotine Cessation During Pregnancy CO 2020 (2023)
- Substance Abuse Reporting and Pregnancy CO 2011 (2022)

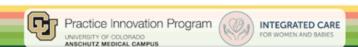
A lot of OLD information

- Opioid Use and OUD in Pregnancy CO 2017 (2021)
 - Screening and intervention improve outcomes
 - UNIVERŠAL screening should be part of comprehensive obstetric care
 - pharmacotherapy is preferable to withdrawal
 - infants should be monitored
 - consider modifying some elements of prenatal care
 - prescribe opioids appropriately
 - Breastfeeding encouraged if stable on opioid agonists
 - postpartum support important
 - contraceptive counseling and access should be a routine part o SUD treatment

Part 2 Conclusion

- HIV, Syphilis, and Hep B should be screened for and can safely be treated in pregnancy to improve outcomes
- Hep C is less easily treated in pregnancy (though possible) and we should improve preconception detection and treatment when possible
- Expand our use of HepA and HepB vaccines
- There is no current consensus on additional antenatal testing for most pre-existing or primary viral infections

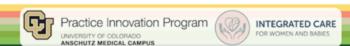
 Outstand additional antenatal testing.
 - Syphilis panel later today!
- Genetic screening should be offered universally but increasingly important in individuals delivering over age of 35
- Nationally, we need more study and evidence based care in substance use disorder in pregnancy



My Soapbox



SCFM's HROB Grid



QUESTIONS?



References



- American College of Obstetrics and Gynecologist - <u>acog.org/clinical</u>
 - Special THANK YOU to Dr Michael Growney, OBGYN
- Obstetrics Normal and Problem Pregnancies. Gabbe et al. 8th ed. 2020
- Obstetrics & Gynecology Handbook.
 Zheng. 3rd ed. 2020
- Southern Colorado Maternal Fetal Medicine Group