Boston Medical Center Maternity Care Guideline

Guideline: Routine Prenatal Screening

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Introduction

Definition: Routine screening tests are recommended during prenatal care to promote optimal maternal-fetal outcomes. There are also vaccinations recommended during routine pregnancy care for disease prevention (see separate guideline).

Prevalence: All pregnant patients should be offered routine screening.

Risk factors: Some labs are recommended for high risk populations only. These populations include, but are not limited to the following:

- Hemoglobin electrophoresis: African American, African, Southeast Asian, and Mediterranean descent
- Early GDM screening: High risk for GDM
- TB screening: High risk for TB
- Hepatitis C: History of IVDU
- Hepatitis B: Southeast Asian, History of IVDU

Diagnosis

ROUTINE PRENATAL LABS FIRST TRIMESTER

- Blood type and Antibody Screen
- CBC (HYPERLINK ANEMIA GUIDELINE)
- Syphilis IgG/IgM UPDATE [see Appendix I]
- HIV (document verbal consent obtained)
- Hepatitis B Surface Antigen (HBSaG)
- Gonorrhea/Chlamydia (urine, cervical, or vaginal specimen)
- Urine culture
- PAP if indicated (hyperlink BMC PAP testing guideline)
- Varicella antibody (if unknown or prior non-immune status)
- Rubella antibody (if unknown or prior non-immune status)
- Early GLT if at risk (see Appendix II)
- TB screen if at risk (see Appendix III)
- Hepatitis C if at risk (See Appendix IV)
- Hepatitis B antibody if at risk (See appendix V)
- Genetic Testing: First Trimester Screen (FTS) 11w3d-13w4d

ONCE IN A LIFETIME

- Varicella Antibody [once in lifetime if found to be immune]
- Rubella Antibody [once in lifetime if found to be immune]
- Hepatitis B core antibody if high risk (see appendix VI)

Carrier Screening for Genetic Condition

- Hemoglobin Electrophoresis
 - o African American, African, Southeast Asian, and Mediterranean are high risk populations, but reasonable to offer in all populations
- Cystic Fibrosis
 - o Recent updates recommend CF carrier screening in all pregnant women
- SMA: spinal muscular atrophy
 - o Is recommended for all pregnant women for carrier screening
- Fragile-X
 - o Is recommended for women with a family history of fragile X-related disorders or intellectual disability suggestive of fragile X
- Tay-Sachs
 - Screening for Tay–Sachs disease is recommended for women of Ashkenazi Jewish, French–Canadian, Cajun descent, or with a family history consistent with Tay–Sachs disease

15-20 WEEKS

- QUAD screen if 1st trimester screen/early risk assessment or MaterniT21 was not done
- Offer isolated AFP if 1st trimester screen/early risk assessment was done to evaluate for risk of neural tube defects

24-28 WEEKS

- CBC
- 1hr Glucola (GDM GUIDELINE)
- Repeat Syphilis IgG/IgM if at high risk
 - o multiple sexual partners, sex worker, IVDU, prior positive

28 WEEKS

- RH NEG: Anti-body screen and RHOGAM
 - Antibody screen should be drawn prior to RHOGAM administration, but you DO NOT need to wait until anti-body screen is resulted

35-37 WEEKS

- Vaginorectal GBS culture.
 - Swab the lower vagina (vaginal introitus), followed by the rectum (insert swab through the anal sphincter)
 - Exceptions: If history of positive GBS UTI in current pregnancy or prior infant with invasive GBS disease—no testing indicated, plan on treatment in labor.
 - o Order sensitivities if patient has a PCN allergy
 - o If positive GBS vaginal culture prior to 34 weeks (eg: during a PTL work up) GBS needs to be re-tested.

- o GBS testing is considered valid for 6 weeks. Consider re-swab if greater than 6 weeks since GBS testing
- HIV: Third trimester HIV testing should be offered to all pregnant patients.
 - o If patients decline testing, please document this in problem list under supervision of pregnancy. If patients have a documented negative first trimester HIV test and decline 3rd trimester screening, they do not need to be offered testing on L and D
- GC/Chlamydia: Re-screen if at risk
 - o teen, prior positive in this pregnancy, high risk sexual activity, or patient desires

POSTPARTUM

Screening for diabetes if GDM at 6 weeks postpartum (fasting glucose or 2 hr 75 gm GT)*
GDM GUIDELINE)

GENETIC TESTING

Genetic testing should be offered to all pregnant patients. Some patients may opt not to have testing. Genetic counseling and testing is recommended in high risk populations who are AMA, or have a personal/family history of a child with a genetic disorder or anomaly. (Hyperlink to GENETICS GUIDELINE)

- First Trimester Screen (FTS) 11w3d-13w4d
- NIPT (MT21) offered as early as 10wks for high risk populations. There may be insurance coverage issues. This test should only be ordered by the genetic counselor in the ATU.
- Quad screen (if FTS not done) 15-22w6d weeks (most accurate between 16-18 weeks)
- MSAFP only (if FTS is done) 15-22w6d weeks (most accurate between 16-18 weeks)

Treatment/Management

- CDC 2015 STD Screening Recommendations
 - o http://www.cdc.gov/std/tg2015/screening-recommendations.htm
- Recommendations based on evidence based protocols for positive testing
- For abnormal genetic screening follow ATU genetic screening workflow (HYPERLINK TO GENETICS GUIDELINE)

Logistical Practice for BMC and CHC providers

- Use the EPIC Smartphrase ".bmcobprenatallabs" to pull most recent labs into problem list under "supervision of pregnancy" at BMC
- Patients with positive carrier or genetic screening should be referred to Genetics for consultation. Call 617-414-7292 to schedule or contact Philip Conner, Genetic Counselor, via EPIC flag.
- For +TB screening: Put Screening for Tuberculosis on the problem list. Use the smart phrase .bmcobtbrisk. (HYPERLINK TO TB PROTOCOL)

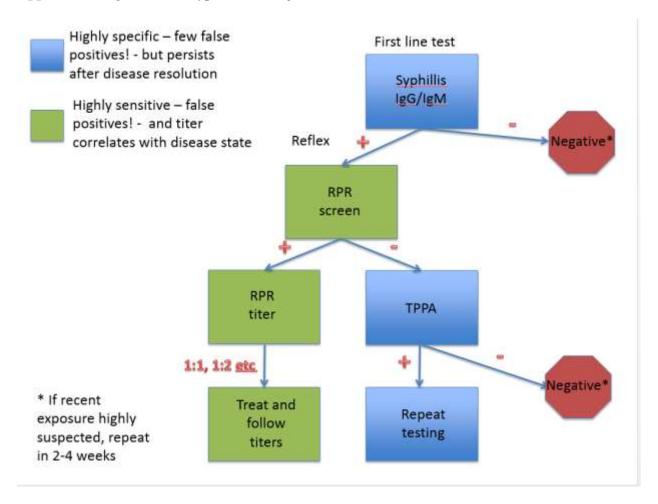
Patient Education/Patient Education Materials

• GBS patient materials in English and Spanish http://www.cdc.gov/groupbstrep/resources/print-materials.html

Appendix

SPECIAL POPULATIONS TESTING

Appendix I: Algorithm for Syphillis Testing



Appendix II: Early GLT Screening: (GDM guideline)

Pregnant women who meet the following criteria should be screened as early as possible, preferably at the first prenatal visit. If the initial screening result is normal, they should be re-screened at 24-28 weeks. Indications for early testing include, but are not limited to:

- Personal history of GDM
- Obesity (BMI≥30)
- PCOS
- Impaired glucose tolerance
- Glycosuria early in pregnancy
- Strong family history of diabetes (one first degree relative, or more than one second degree relative)
- Previous macrosomic infant.
- Previous unexplained third trimester loss or neonatal death.

Appendix III Tuberculosis screening: Pregnant women at high risk for tuberculosis should be screened in the first trimester of pregnancy. (TB PROTOCOL)

- Symptomatic for TB disease (fever, night sweats, cough, and weight loss)
- Close contact with someone with active TB
- Foreign-born or >1month travel to high prevalence country (>25/100,000) www.who.int/tb/data for country atlas
- Resident of large congregate shelter (NOT small family shelters) or recent incarceration (without test)
- HIV infection, organ transplant, other immunosuppression
- Active IV drug use
- Medical conditions associated with risk of progression to TB disease if exposed/infected: diabetes mellitus (preexisting), silicosis, cancer of head or neck, Hodgkin's disease, leukemia, end-stage renal disease, intestinal bypass or gastrectomy, chronic malabsorption syndrome, extremely low body weight
- CXR with fibrotic changes suggestive of old TB

At this time, **TST/PPD** is the preferred test for our patients. The BMC TB clinic strongly discourages use of Quant Gold because of known "problems with conversions, reversions, reliability, and inadequate data to interpret the findings" (A. Barry, October 2015, personal communication). Use of Quant Gold may be **considered** ONLY for women who are high risk AND absolutely will not return for PPD reading or are from countries where BCG vaccine **boosters** are given past infancy. **www.bcgatlas.org** Patient's from East Boston Neighborhood Health Center are generally screened with Quantiferon Gold testing at this time.

Get a CXR for women with a positive test and refer to TB clinic **during the pregnancy** and follow up postpartum. Treatment for LTBI is most commonly started three months postpartum if indicated.

Appendix IV

Hepatitis C antigen Screen:

- History of IV drug use
- Tattoos

Appendix V

Hepatitis B antibody screen

Anti-Hbc testing should be performed in the following high risk populations:

- Household, sex, and needle-sharing contacts of HBsAg-positive persons
- HIV-infected persons
- Injection drug-users
- Incarcerated persons

Foreign born persons from countries with Hepatitis B prevalence of >8% (listed below) Africa: all countries except Algeria, Djibouti, Egypt, Libya, Morocco, Tunisia

- Southeast Asia: All countries except Malaysia
- East Asia: China, Hong Kong, Mongolia, North Korea, South Korea, Taiwan
- Australia and South Pacific: all countries except Australia, Guam, and New Zealand
- Middle East: Jordan and Saudi Arabia
- Eastern Europe and Northern Asia: Albania, Armenia, Azerbaijan, Moldova, Tajikistan, Turkmenistan, Uzbekistan

Appendix VI

HIV High Risk

- CDC defines high HIV prevalence as an area in which 17/100,000 people are infected with HIV. Almost all neighborhoods in Boston fall into this category.
- CDC recommends a repeat screening in the third trimester in places with elevated rates of HIV infection in pregnant women, which includes Boston

References

ACOG. ACOG Committee Opinion No.691. Carrier Screening for Genetic Conditions. March 2017:1-15.

ACOG. ACOG Practice Bulletin No. 78. Hemoglobinopathies in pregnancy. Obstet Gynecol. 2007 Jan;109(1):229-37.

References for HIV screening recommendations:

Specific most recent numbers by neighborhood are in page 50 in this document:

http://www.bphc.org/healthdata/archive/Documents/Health%20of%20Boston%202008_Online.pdf General guidelines: 17/100,000 is considered high HIV prevalence by CDC; please see this document for prenatal HIV screening guidelines by CDC.

http://www.cdc.gov/mmwr/preview/mmwrhtml/rr5514a1.htm