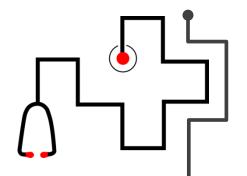


DM IN PREGNANCY

พญ.ธารทิพย์ อุทัยพัฒน์ สูติแพทย์ โรงพยาบาลกำแพงเพชร



Scope

- **01** Gestational Diabetes Mellitus
- **02** Pregestational Diabetes Mellitus
- 03 KPH Guideline



INTERIM UPDATE

ACOG PRACTICE BULLETIN

Clinical Management Guidelines for Obstetrician-Gynecologists

NUMBER 190, FEBRUARY 2018

(Replaces Practice Bulletin Number 180, July 2017)

Committee on Practice Bulletins—Obstetrics. This Practice Bulletin was developed by the American College of Obstetricians and Gynecologists Committee on Practice Bulletins—Obstetrics with the assistance of Aaron B. Caughey, MD, PhD, and Mark Turrentine, MD.

INTERIM UPDATE: This Practice Bulletin is updated as highlighted to reflect a limited, focused change to clarify and provide additional information on the pharmacologic treatment of gestational diabetes mellitus.

Gestational Diabetes Mellitus



ACOG PRACTICE BULLETIN

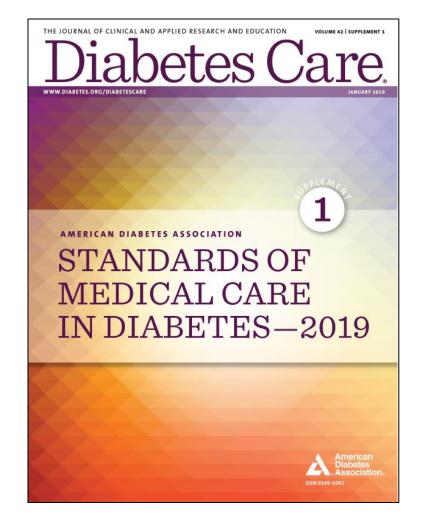
Clinical Management Guidelines for Obstetrician-Gynecologists

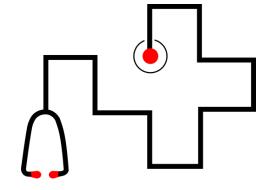
NUMBER 201

(Replaces Practice Bulletin Number 60, March 2005)

Committee on Practice Bulletins—Obstetrics. This Practice Bulletin was developed by the American College of Obstetricians and Gynecologists' Committee on Practice Bulletins—Obstetrics with the assistance of Aaron B. Caughey, MD, PhD; Anjali J. Kaimal, MD, MAS; and Steven G. Gabbe, MD.

Pregestational Diabetes Mellitus





Gestational Diabetes Mellitus

Maternal and Fetal Complications





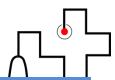
Maternal complication

- Preeclampsia
- Cesarean delivery
- Increased risk of developing diabetes

Fetal complication

- Macrosomia
- Neonatal hypoglycemia
- Hyperbilirubinemia
- Shoulder dystocia
- Birth trauma
- Stillbirth
- Childhood and adultonset obesity and diabetes in offspring

GDM: Screening



Universal screening¹

• GA 24-28 week

Preventive Services Task Force recommendation statement 2014

Risk based screening (KPH)

- Age \geq 30 years (GCT at GA 24-28 week)
- Family history of DM in first degree relatives
- History of GDM
- Obesity as defined as pre-pregnancy BMI \geq 25 kg/m²
- History of fetal macrosomia (BW \geq 4000 g)
- History of unknown congenital fetal anomaly
- History of unexplained stillbirth/fetal death
- Hypertension
- Glucosuria; dipstick $\geq 1+$
- Excessive weight gain; ≥ 2 kg in 1 week

Screening (NEW)

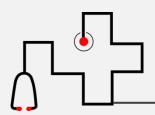


Early screening at 1st ANC

Pregestational DM or early GDM

GA 24-28 week(Universal)

• GDM



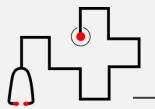
ADA 2019 ACOG 2018

Pregestational DM

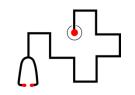
Early screening at 1st ANC

- BMI > 25 and one or more additional risk factors
 - Physical inactivity
 - First-degree relative with diabetes
 - High-risk race or ethnicity
 - Previous child \geq 4000 g
 - Previous GDM
 - Hypertension
 - HDL < 35 mg/dL, TG > 250 mg/dL
 - PCOS
 - HbA1C \geq 5.7%, IGT, IFG
 - Insulin resistance; severe obesity, acanthosis nigricans
 - Hx of cardiovascular disease





Pregestational DM



FPG ≥ 126 mg/Dl (no caloric intake ≥ 8 hr)

OR

2hr PG ≥ 200 mg/dL (75-g OGTT)

OR

A1C ≥ 6.5%

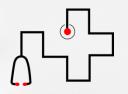
OR

Random plasma glucose
≥ 200 mg/dL
with classic symptoms of
hyperglycemia or
hyperglycemic crisis

Pregestational DM

ADA 2019

GDM: Screening and diagnosis



Two-step approach





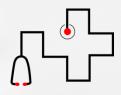


100-g OGTT

	Carpenter and Coustan	National Diabetes Data Group
Fasting	95 mg/dL	105 mg/dL
1 hour	180 mg/dL	190 mg/dL
2 hours	155 mg/dL	165 mg/dL
3 hours	140 mg/dL	145 mg/dL

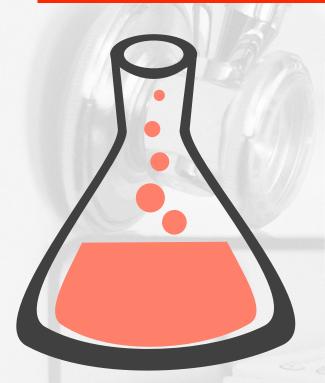
Abnormal ≥ 2

GDM: Screening and diagnosis



One-step approach

75-g 2-hour OGTT



	International Association of Diabetes and Pregnancy Study Group (IADPSG)
Fasting	92 mg/dL
1 hour	180 mg/dL
2 hours	153 mg/dL

Abnormal ≥ 1

Blood glucose monitoring





Fasting or preprandial;

GOAL < 95 mg/dL

Postprandial (1 or 2 hour); GOAL

1 hour < 140 mg/dL

2 hour < 120 mg/dL

Fasting: Neonatal fat mass

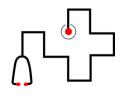
childhood obesity and diabetes

better glycemic control

lower incidence of LGA

lower rates of cesarean delivery

Nonpharmacologic treatments





Diet

Carbohydrate: Protein: Fat

33-40: 20: 40

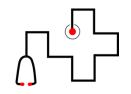
3 meals and 2 snacks

Exercise

- 30 minutes of moderate-intensity aerobic exercise at least 5 days a week
- or a minimum of 150 min/wk
- walking for 10-15 min after each meal



Pharmacologic treatments



First line therapy: Insulin



Does not cross the placenta



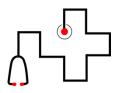
Starting dose is 0.7-1.0 units/kg/day



Long-acting or intermediate acting insulin in combination with short-acting insulin



Pharmacologic treatments





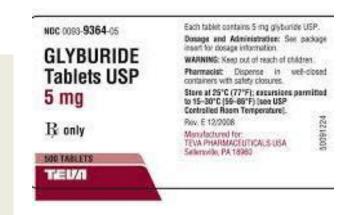
Metformin

- Equivalent to insulin
- Cross placenta
- Limit long term data
- may be a reasonable alternative approach

Oral Medications

Glyburide

- Cross placenta
- Increased risks of macrosomia and Hypoglycemia
- Should not be recommended as a first-choice pharmacologic treatment



Antepartum Care



Fetal testing

GA 32 wk

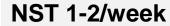
GA 40+6 wk

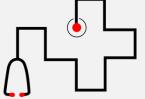


- Poor-controlled GDM
- Medication use

Well-controlled GDMA1

NST 1/week





Antepartum Care

Fetal Growth



GA 32 wk GA 36-39 wk

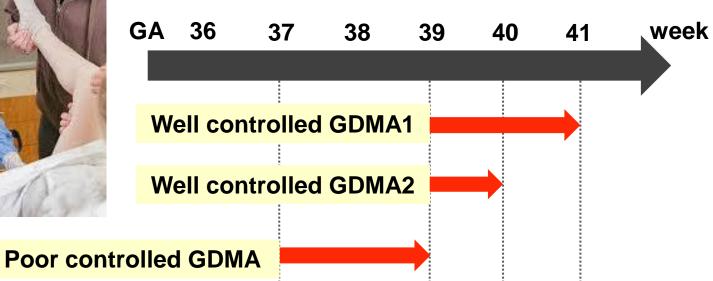
Ultrasound

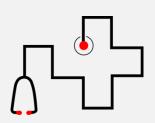
Fetal weight AFI

Delivery



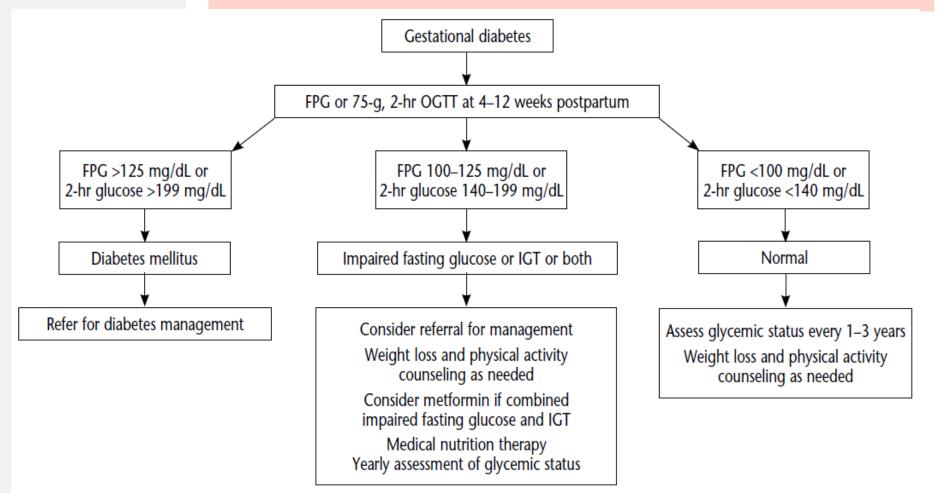
Timing

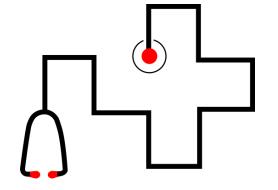




Reasonable to scheduled C/S when EFW ≥ 4,500 g

Postpartum Care

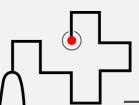




Pregestational Diabetes Mellitus

Maternal and Fetal Complications





Maternal complication

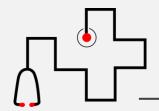
- Diabetic retinopathy
- Diabetic nephropathy
- Hypertension
- Acute myocardial infarction
- Diabetic neuropathy
- Diabetic ketoacidosis
- Preterm labor
- Polyhydramnios

Fetal complication

- Congenital anomalies; Complex cardiac defects; CNS, anencephaly and spina bifida; and skeletal malformations, sacral agenesis
- Spontaneous abortion
- Macrosomia
- Neonatal hypoglycemia
- Hyperbilirubinemia, Polycythemia
- Shoulder dystocia
- Stillbirth
- Obesity and diabetes

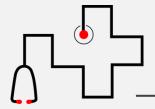
Management: Prepregnancy visit

- 1. Counsel about potential complications
- 2.Evaluate for baseline complications;
 hypertension, nephropathy, retinopathy, and
 cardiovascular disease
- 3. Adequate contraception
- 4. Optimize HbA1C (< 6.0%)
- 5.Start increased folic acid when attempting to get pregnant (400 micrograms of folic acid)



Management: First trimester

- 1.Prenatal labs/tests
 - HbA1C
 - TSH
 - 24-hour urine
 - electrocardiogram
- 2. Evaluation by ophthalmologist, dietitian, possibly endocrinologist, cardiologist, nephrologist
- 3.Regular ongoing assessment of blood glucose



Management: Second trimester

1.Start low-dose aspirin 12-28 weeks of gestation (optimally before 16 week)

2. Ultrasonography; detailed anatomical survey

3. Consider fetal echocardiography

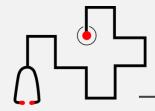


Table 1. Clinical Risk Factors and Aspirin Use*

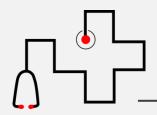
Level of Risk	Risk Factors	Recommendation	
High [†]	 History of preeclampsia, especially when accompanied by an adverse outcome 	Recommend low-dose aspirin if the patient has one or more of these high-risk factors	
	 Multifetal gestation 		
	Chronic hypertension		
	• Type 1 or 2 diabetes		
	Renal disease		
	 Autoimmune disease (ie, systemic lupus erythematosus, the antiphospholipid syndrome) 		
Moderate [‡]	 Nulliparity 	Consider low-dose aspirin if the patient has	
	 Obesity (body mass index greater than 30) 	more than one of these moderate-risk factors [§]	
	 Family history of preeclampsia (mother or sister) 		
	 Sociodemographic characteristics (African American race, low socioeconomic status) 		
	Age 35 years or older		
	 Personal history factors (eg, low birth weight or small for gestational age, previous adverse pregnancy outcome, more than 10-year pregnancy interval) 		
Low	 Previous uncomplicated full-term delivery 	Do not recommend low-dose aspirin	

Management: Third trimester

1. Evaluate fetal growth

2.Start low-dose aspirin by 28 weeks of gestation if not started in the second trimester

3.Fetal monitoring (nonstress test or amniotic fluid index, biophysical profile)



Blood glucose monitoring



Four times a day

Fasting or preprandial;

GOAL < 95 mg/dL

Postprandial (1 or 2 hour); GOAL

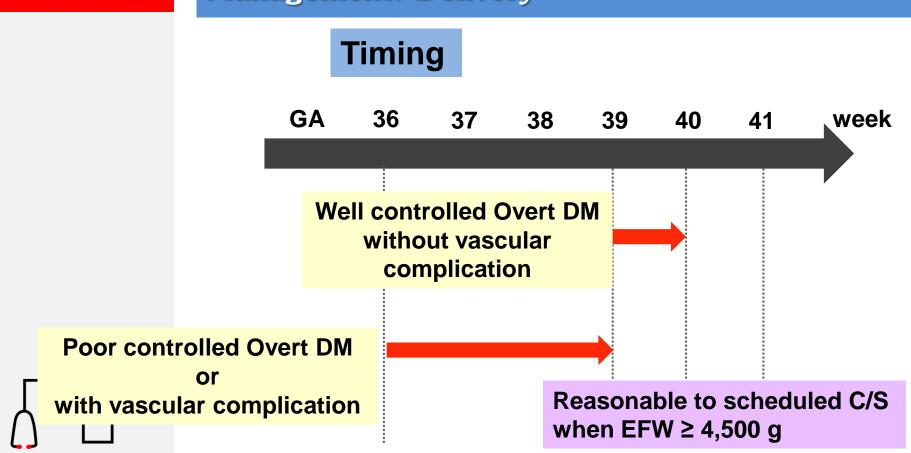
1 hour < 140 mg/dL

2 hour < 120 mg/dL

Mean capillary glucose ~100 mg/dL

HbA1C < 6%

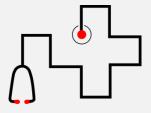
Management: Delivery



Management: Intrapartum

Box 3. Insulin Management During Labor and Delivery

- Usual dose of intermediate-acting or long-acting insulin is given at bedtime.
- Morning dose of insulin is withheld or reduced based upon the timing of admission or delivery.
- Intravenous infusion of normal saline is begun.
- Once active labor begins or glucose levels decrease to less than 70 mg/dL, the infusion is changed from saline to 5% dextrose and delivered at a rate of 100–150 cc/h (2.5 mg/kg/min) to achieve a glucose level of approximately 100 mg/dL.
- Glucose levels are checked hourly using a bedside meter allowing for adjustment in the insulin or glucose infusion rate.
- Regular (short-acting) insulin is administered by intravenous infusion at a rate of 1.25 units/h if glucose levels exceed 100 mg/dL.



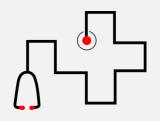
Management: Postpartum

Breastfeeding +500 kcal/d

Contraception

- Permanent contraception with tubal ligation
- Long-acting reversible contraception;

IUD or implantable progestin





KAMPHAENGPHET GUIDELINE

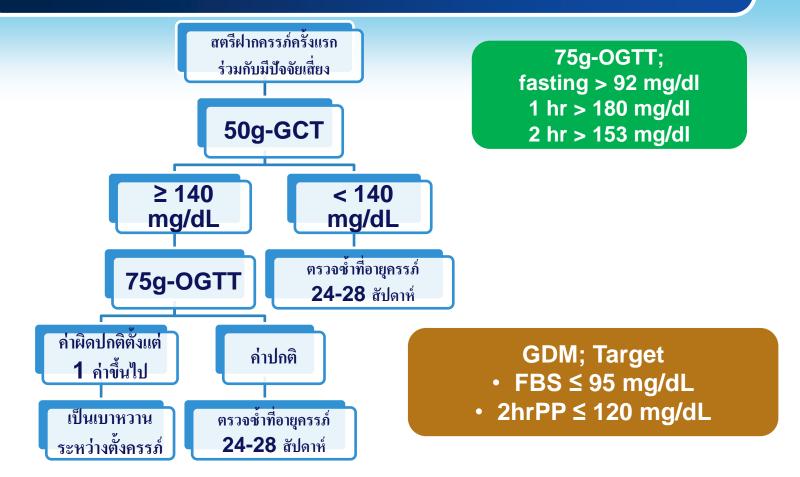


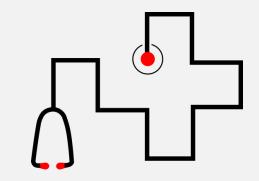
KPH guideline

ปัจจัยเสี่ยงต่อการเกิดเบาหวานระหว่างตั้งครร<u>ภ</u>์

- Age ≥ 30 years (GCT at GA 24-28 week)
- Family history of DM in first degree relatives
- History of GDM
- Obesity as defined as pre-pregnancy BMI ≥ 25 kg/m²
- History of fetal macrosomia (BW ≥ 4000 g)
- History of unknown congenital fetal anomaly
- History of unexplained stillbirth/fetal death
- Hypertension
- Glucosuria; dipstick ≥ 1+
- Excessive weight gain; ≥ 2 kg in 1 week







Thank you