



CLINICAL GUIDELINES

Obstetrical Ultrasound Imaging Guidelines

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EviCore healthcare Clinical Decision Support Tool Diagnostic Strategies: This tool addresses common symptoms and symptom complexes. Imaging requests for individuals with atypical symptoms or clinical presentations that are not specifically addressed will require physician review. Consultation with the referring physician, specialist, and/or individual's Primary Care Physician (PCP) may provide additional insight.

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Table of Contents

Guideline	Page
Obstetrical Ultrasound Imaging General Guidelines (OB-1)	6
Abbreviations and Glossary for OB Ultrasound Imaging Guidelines.....	7
General Guidelines (OB-1.0).....	10
Inappropriate Use of OB Ultrasound (OB-1.2).....	12
Ultrasound Code Selection (OB-1.3).....	13
References (OB-1).....	17
Uncertain Dates (OB-2)	18
Uncertain Dates/Unknown Last Menstrual Period (LMP) (OB-2.1).....	19
References (OB-2).....	20
Intrauterine Device (IUD) (OB-3)	21
Locate an Intrauterine Device (IUD) (OB-3.1).....	22
References (OB-3).....	23
Infertility (OB-4)	24
History of Infertility (OB-4.1).....	25
Present Pregnancy with ART Treatment (IVF) (OB-4.2).....	26
Recurrent Pregnancy Loss (OB-4.3).....	27
References (OB-4).....	28
Vaginal Bleeding and/or Abdominal/Pelvic Pain/Cramping (OB-5)	29
Acute Abdominal/Pelvic Pain (OB-5.1).....	30
Vaginal Bleeding (OB-5.2).....	31
Ectopic Pregnancy (OB-5.3).....	32
Spontaneous Abortion/Threatened/Missed Abortion (OB-5.4).....	34
Hydatidiform Mole (OB-5.5).....	35
References (OB-5).....	36
Fetal Aneuploidy and Anomaly Screening (OB-6)	37
First Trimester Screening (OB-6.1).....	38
Second Trimester Screening (OB-6.2).....	40
References (OB-6).....	41
Fetal Anatomic Scan/Cervical Length Screening (OB-7)	42
Fetal Anatomic Scan (OB-7.1).....	43
Fetal Anatomic Scan – Follow-up (OB-7.2).....	44
Cervical Length Screening (OB-7.3).....	45
References (OB-7).....	46
Third Trimester Imaging (OB-8)	47
Third Trimester Imaging – Ultrasound (OB-8.1).....	48
Reference (OB-8).....	49

High Risk Pregnancy (OB-9)	50
High Risk General Information (OB-9.0).....	51
High Risk Group One – Risk Factors (OB-9.1).....	53
High Risk Group Two - Ultrasound Findings (OB-9.2).....	59
High Risk Group Three – Pre-pregnancy BMI ≥ 30 kg/m ² (OB-9.3).....	61
High Risk Group Four – Macrosomia (OB-9.4).....	64
High Risk Group Five – Zika and COVID-19 Virus (OB-9.5).....	65
High Risk Group Six - Pre-Gestational or Early Diagnosed (≤ 20 weeks) Diabetes (OB-9.6).....	68
High Risk Group Seven Gestational Diabetes (OB-9.7).....	70
Hypertensive Disorders in Pregnancy (OB-9.8).....	72
History of Spontaneous Pre-Term Delivery/History of PPRM (OB-9.9).....	76
History of Stillbirth (OB-9.10).....	77
Short Interval Pregnancy (≤ 18 months between last delivery and conception of current pregnancy) (OB-9.11).....	78
Detailed First Trimester Fetal Anatomic Scan (OB-9.12).....	79
References (OB-9).....	80
High Risk Medications and Substances (OB-10)	83
Potentially Teratogenic Medications/Substances (OB-10.1).....	84
Medications/ Exposures Associated with Poor Pregnancy Outcome (OB-10.2).....	87
References (OB-10).....	90
Multiple Gestations (OB-11)	91
Suspected Multiple Gestations (OB-11.1).....	92
Known Dichorionic Multiple Gestations (OB-11.2).....	93
Known Monochorionic-Diamniotic or Monochorionic-Monoamniotic Multiple Gestations (OB-11.3).....	96
References (OB-11).....	99
Fetal Echocardiography (ECHO) (OB-12)	101
Fetal Echocardiography – Coding (OB-12.1).....	102
Fetal Echocardiography - Indications for Fetal Conditions (OB-12.2).....	103
Fetal Echocardiography - Indications for Maternal Conditions (OB-12.3).....	105
Fetal Echocardiography - Indications for Medication or Drug Exposure (OB-12.4).....	107
References (OB-12).....	108
Fetal MRI (OB-13)	109
Indications for Fetal MRI (OB-13.1).....	110
References (OB-13).....	113
Abnormal Fetal Position/ Presentation and Pelvimetry (OB-14)	114
Abnormal Fetal Position or Presentation (OB-14.1).....	115
Pelvimetry (OB-14.2).....	116
References (OB-14).....	117
Adnexal Mass/Uterine Fibroids and Uterine Anomalies (OB-15)	118
Adnexal Mass (OB-15.1).....	119

Uterine Fibroids in Pregnancy (OB-15.2).....	121
Uterine Anomalies in Pregnancy (OB-15.3).....	123
References (OB-15).....	124
Alloimmunization/Rh Isoimmunization/Other Causes of Fetal Anemia/Parvo/	
Hydrops (OB-16).....	125
Alloimmunization/Rh Isoimmunization (OB-16.1).....	126
Exposure to Parvovirus B-19 (OB-16.2).....	128
Twin Anemia Polycythemia Sequence (OB-16.3).....	129
Other Fetal Hydrops/Nonimmune Hydrops (OB-16.4).....	130
Other Causes of Fetal Anemia (OB-16.5).....	131
References (OB-16).....	132
Amniotic Fluid Abnormalities/ Oligohydramnios/Polyhydramnios (OB-17).....	133
Amniotic Fluid Abnormalities (OB-17.1).....	134
References (OB-17).....	136
Cervical Insufficiency/Current Preterm Labor (OB-18).....	137
Cervical Insufficiency (OB-18.1).....	138
Cerclage in Place in Current Pregnancy (OB-18.2).....	139
Current Preterm Labor (OB-18.3).....	140
References (OB-18).....	141
No Fetal Heart Tones/Decreased Fetal Movement (OB-19).....	142
No Fetal Heart Tones (OB-19.1).....	143
Decreased Fetal Movement (OB-19.2).....	144
References (OB-19).....	145
Fetal Growth Problems (FGR and Macrosomia) (OB-20).....	146
Fetal Growth Restriction Current Pregnancy (OB-20.1).....	147
Macrosomia – Large for Dates Current Pregnancy (OB-20.2).....	149
References (OB-20).....	150
Placental and Cord Abnormalities (OB-21).....	151
Single Umbilical Artery (Two Vessel Cord) (OB-21.1).....	152
Persistent Right Umbilical Vein (PRUV) (OB-21.2).....	153
Placental/Cord Abnormalities (OB-21.3).....	154
Subchorionic Hematoma/Hemorrhage (Placental Hematoma) (OB-21.4).....	157
Suspected Abruptio Placentae (OB-21.5).....	158
Previa (Placenta Previa and Vasa Previa) (OB-21.6).....	159
Placenta Accreta Spectrum (PAS): Accreta, Increta, Percreta (OB-21.7).....	161
References (OB-21).....	163
Late-term/Post-term Pregnancy (OB-22).....	165
Late-term/Post-term Pregnancy (OB-22.1).....	166
References (OB-22).....	167
Preterm/Prelabor Rupture of Membranes (OB-23).....	168
Current Preterm/Prelabor Rupture of Membranes (PPROM) (OB-23.1).....	169
Current Prelabor Rupture of Membranes (PROM) (OB-23.2).....	170

References (OB-23).....	171
Previous C-section or History of Uterine Scar (OB-24).....	172
Previous C-section or History of Uterine Scar (OB-24.1).....	173
References (OB-24).....	175
Termination of Pregnancy – Imaging (OB-25).....	176
Imaging for Planned Pregnancy Termination (OB-25.1).....	177
References (OB-25).....	178
Trauma (OB-26).....	179
Trauma – Imaging (OB-26.1).....	180
References (OB-26).....	182
Unequal Fundal Size and Dates (OB-27).....	183
Unequal Fundal Size and Dates (OB-27.1).....	184
References (OB-27).....	185
Procedure Coding Basics for Established Pregnancy (OB-28).....	186
Procedure Coding Basics for Established Pregnancy General Considerations (OB-28.1).....	187
Required Elements for Complete First Trimester Ultrasound (OB-28.2).....	188
Required Elements for Second or Third Trimester Fetal Anatomic Evaluation Ultrasound (OB-28.3).....	189
Required Elements for a Detailed Fetal Anatomic Evaluation Ultrasound (OB-28.4)...	191
Fetal Nuchal Translucency (OB-28.5).....	193
Limited and Follow-up Studies (OB-28.6).....	194
Obstetric Transvaginal Ultrasound (OB-28.7).....	195
Biophysical Profile (BPP) (OB-28.8).....	196
Fetal Doppler (OB-28.9).....	197
Duplex Scan (OB-28.10).....	198
Fetal Echocardiography (OB-28.11).....	199
3D and 4D Rendering (OB-28.12).....	200
Required Elements for a Detailed First Trimester Fetal Anatomic Evaluation Ultrasound (OB-28.13).....	201
References (OB-28).....	203

Obstetrical Ultrasound Imaging General Guidelines (OB-1)

Guideline	Page
Abbreviations and Glossary for OB Ultrasound Imaging Guidelines.....	7
General Guidelines (OB-1.0).....	10
Inappropriate Use of OB Ultrasound (OB-1.2).....	12
Ultrasound Code Selection (OB-1.3).....	13
References (OB-1).....	17

Abbreviations and Glossary for OB Ultrasound Imaging Guidelines

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Abbreviations and Glossary for OB Ultrasound Imaging Guidelines	
ACOG	American College of Obstetricians and Gynecologists
AFI	amniotic fluid index
AFP	alpha-fetoprotein
ART	Assisted Reproductive Technology
B-mode (brightness)	two dimensional imaging procedure, B-mode ultrasound is the basis for all static and real time B-scan images
BPP	Biophysical Profile includes the ultrasound variables: fetal breathing, muscle tone, and movement as well as amniotic fluid volume. BPP can be performed with or without a non-stress test (NST) which involves fetal heart rate (FHR) monitoring.
CST	contraction stress test
D & C/D & E	dilatation and curettage/ Dilation and Evacuation
Dichorionic twins	twins having distinct chorions (membrane that forms the fetal part of the placenta), including monozygotic twins (from one oocyte [egg]) separated within 72 hours of fertilization and all dizygotic twins (from two oocytes fertilized at the same time

Obstetrical Ultrasound Imaging Guidelines

Abbreviations and Glossary for OB Ultrasound Imaging Guidelines

Doppler	involves measuring a change in frequency when the motion of vascular flow is measured
EDC	Estimated Date of Confinement; determined from the first day of the last menstrual cycle
EDD	Estimated Date of Delivery
FGR	Fetal growth restriction; an estimated weight of the fetus at or below 10th percentile for gestational age; and/or abdominal circumference of the fetus at or below 10th percentile for gestational age
FHR	fetal heart rate
hCG	human chorionic gonadotropin
IDDM	insulin-dependent diabetes mellitus
M-mode	ultrasound imaging technique in which structure movement can be depicted in a wave-like manner; primarily used in cardiac and fetal cardiac imaging
Macrosomia	estimated fetal weight of greater than 4000 or 4500 grams
Monochorionic twins	twins developed from one oocyte (egg) developing with a single chorions (membrane that forms the fetal part of the placenta)
NICU	Neonatal Intensive Care Unit
NST	fetal non-stress test

Abbreviations and Glossary for OB Ultrasound Imaging Guidelines

Oligohydramnios	diminished amniotic fluid volume (AFV) for gestational age; definitions include: maximum deepest pocket of ≤ 2 cm and/or AFI of ≤ 5 cm or $<$ the 5 th percentile for gestational age if <30 weeks.
PACS	Picture Archiving and Communications System
Polyhydramnios	AFI ≥ 24 cm or maximum vertical pocket of ≥ 8 cm
PROM	preterm rupture of membranes
Quad screen	alpha-fetoprotein (AFP), estriol, human chorionic gonadotropin (hCG), inhibin A
Real time scan	considered the most common type of ultrasound; a 2-dimensional scan that reflects structure and motion over time, scanning and display of images are run at a sufficiently rapid rate so that moving structures can be viewed moving at their natural rate; frame rates ≥ 15 frames per second are considered "real time"

General Guidelines (OB-1.0)

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- This document offers an in-depth, indication driven guide to obstetrical imaging
- The use of an obstetrical CPT code is only indicated with a positive pregnancy test or an otherwise confirmed pregnancy. It is not appropriate to report non-obstetrical, pelvic ultrasound procedure codes (CPT[®] 76830, CPT[®] 76856, and CPT[®] 76857) with a positive pregnancy test or a confirmed pregnancy
- An evaluation of pregnancy with a history and physical exam (an initial office visit) is necessary prior to obstetric ultrasound imaging requests
 - The following information must be submitted with each request:
 - Expected date of delivery
 - Gestational age at date of service
 - Results of prior ultrasound studies if available
- Ultrasound assessment is an accurate method of determining gestational age, fetal number, viability, and placental location, and it is recommended for all pregnant patients
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811 if high-risk) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, the optimal time for a single ultrasound examination is at 18 to 22 weeks of gestation. This timing allows for a survey of fetal anatomy in most females and an accurate estimation of gestational age.²
 - For a Normal (Low Risk) Pregnancy report a fetal anatomy ultrasound CPT[®] 76805 if ≥16 weeks
 - If pregnancy is **High Risk** can report:
 - A detailed first-trimester obstetric ultrasound^{3,15} [requested as CPT[®] 76801 plus CPT[®] 76813 (and CPT[®] 76802 plus CPT[®] 76814 for each additional fetus)] between 12 weeks 0 days and 13 weeks 6 days (if indicated), See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)**, and
 - A detailed fetal anatomy ultrasound (CPT[®] 76811) if ≥16 weeks
 - These high risk scans are indication driven and are generally performed by a Maternal Fetal Medicine (MFM) specialist/Perinatologist, or a Radiologist at an AIUM or ACR accredited facility. See **High Risk Pregnancy (OB-9)**
 - Current ACOG and SMFM guidelines state that the method of cervical length (CL) screening in singleton gestations without a prior spontaneous preterm birth (PTB) cannot yet be universally mandated. Initial cervical length screening by transabdominal or transperineal ultrasound may limit the need for transvaginal exams.
 - Transvaginal ultrasound (CPT[®] 76817) can be performed if:

- The screening transabdominal CL appears shortened (≤ 3.6 cm) and transvaginal exam would be needed to guide therapeutic recommendations
- If a documented screening transabdominal attempt fails to visualize the cervix
- SMFM currently recommends against routine use of progesterone, pessary, or cerclage for the treatment of cervical shortening in twin gestations outside of the context of a clinical trial
 - Therefore, serial measurements of cervical length in multiple gestations is not recommended
- Fetal Nuchal Translucency (CPT[®] 76813) can be performed if Cell-Free DNA (cfDNA) is not planned or has not already been performed, as they are both screening tools for fetal aneuploidy

Inappropriate Use of OB Ultrasound (OB-1.2)

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v1.0.2025

Obstetrical ultrasound studies cannot be authorized for payment for individuals who do not have a positive pregnancy test or clinical evidence of a pregnancy (fetal heart tones)

- Obstetrical ultrasound is **not** medically indicated for the following:
 - Sex determination only
 - To provide a keepsake or souvenir picture

Ultrasound Code Selection (OB-1.3)

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- See **Procedure Coding Basics for Established Pregnancy (OB-28)**
 - It is not appropriate to report non-obstetrical pelvic ultrasound procedure codes (CPT® 76830, CPT® 76856, and CPT® 76857) with a positive pregnancy test or confirmed pregnancy

CPT® Code Guidance

CPT® 76801 and CPT® 76802 (for each additional fetus) are reported for complete studies performed during the first trimester (<14 weeks). These codes should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication for ultrasound.

CPT® 76813 and CPT® 76814 (each additional fetus) are used to report nuchal translucency screening: an ultrasound measurement of the clear (translucent) space at the back of the fetal neck to assess risk for Down Syndrome (Trisomy 21), Trisomy 18, and other genetic disorders.

CPT® 76801 plus CPT® 76813 (and CPT® 76802 **plus** CPT® 76814 for each additional fetus) when billed together, are used to report a detailed first-trimester obstetric ultrasound examination between 12 weeks 0 days and 13 weeks 6 days.^{3,15}

- This indication-driven detailed first trimester fetal anatomic evaluation is generally performed by those with special skills to perform this study, such as a Maternal Fetal Medicine specialist (Perinatologist), or a Radiologist with advanced training in fetal imaging.
- These codes should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication for ultrasound. See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)**

CPT® 76805 and CPT® 76810 (for each additional fetus) are used to report complete studies (anatomy scan) performed during the second and third trimester, in a normal (low risk) pregnancy. These studies should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication for ultrasound.

CPT® Code Guidance

CPT® 76811 and CPT® 76812 (for each additional fetus) describe a detailed fetal anatomic survey and are used only when the study includes this service. These studies should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication for ultrasound.

This detailed fetal anatomic evaluation is generally performed by those with special skills to perform this study, such as a Maternal Fetal Medicine specialist (Perinatologist), or a Radiologist with advanced training in fetal imaging.

In circumstances where a detailed fetal anatomy (CPT® 76811) is indicated but access is limited due to geographic or other constraints, a standard fetal anatomy survey (CPT® 76805) can be authorized instead at the appropriate gestational age.

CPT® 76817 is used to report a transvaginal ultrasound. The other OB ultrasound codes are used for transabdominal studies.

CPT® 76816 is used to report a follow up study, such as a growth scan or follow up on anatomy when more than one area requires reexamination.

- CPT® 76816 [should not be performed prior to a CPT® 76801 or an anatomy scan CPT® 76805 (normal pregnancy) or Detailed anatomy scan CPT® 76811 (high risk pregnancy)]
- CPT® 76816 should not be done on same date of service as CPT® 76815

CPT® 76815 describes a limited or 'quick look' study

- It can be used at any gestational age for various indications, including quick look for AFI assessment, fetal heart-beat, fetal position, placental location etc.
- It can be used specifically for 'dating' (when indicated) in those that don't meet gestational age criteria for dating with CPT® 76801 or are too early for anatomy scan (i.e. >14 weeks but <16 weeks)
- It is also used to report a modified BPP.
- Note: CPT® 76815 should never be reported with complete studies CPT® 76801/ CPT® 76802, CPT® 76805/CPT® 76810, CPT® 76811/CPT® 76812, or with CPT® 76816 or BPP (CPT® 76818 and CPT® 76819).

CPT® 76818 (includes non-stress test) and CPT® 76819: are used to report a Biophysical profile (BPP), a test for antepartum fetal surveillance (A BPP is not typically performed before 26 weeks due to lack of fetal brain stem maturity prior to this gestational age).

CPT® Code Guidance

CPT® 76820 describes Doppler velocimetry of the umbilical artery.

CPT® 76821 describes Doppler velocimetry of the middle cerebral artery.

CPT® 76825 describes fetal echocardiography and CPT® 76827 describes the Doppler portion of the echocardiogram. These codes should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office or there is a new medical indication for ultrasound.

CPT® 76826 describes a follow up fetal echocardiography and CPT® 76828 describes a follow up Doppler portion of the echocardiogram.

CPT® 93325 can be added for color mapping in conjunction with fetal echocardiography procedures.

CPT® 93976 describes a limited duplex scan and is used during pregnancy for characterizing the pattern and direction of blood flow in arteries and veins. It can be used to report fetal umbilical-placental flow evaluation (accreta or other placental or cord abnormalities).

CPT® 74712 and CPT® 74713 (for each additional fetus) are used to report a fetal MRI (indicated for more in depth imaging of certain fetal abnormalities).

Background and Supporting Information

- ACOG recommendations for imaging during pregnancy and lactation:
 - Ultrasonography and magnetic resonance imaging (MRI) are not associated with risk and are the imaging techniques of choice for the pregnant patient, but they should be used prudently and only when use is expected to answer a relevant clinical question or otherwise provide medical benefit to the patient.
 - With few exceptions, radiation exposure through radiography (Xrays), computed tomography (CT) scan, or nuclear medicine imaging techniques is at a dose much lower than the exposure associated with fetal harm. If these techniques are necessary in addition to ultrasound or MRI or are more readily available for the diagnosis in question, they should not be withheld from a pregnant patient.
 - The use of gadolinium contrast with MRI should be limited; it should be used as a contrast agent in a pregnant female only if it significantly improves diagnostic performance and is expected to improve fetal or maternal outcome.

- With regards to iodinated IV contrast media, “it is generally recommended that contrast only be used if absolutely required to obtain additional diagnostic information that will affect the care of the fetus or female during pregnancy.”
- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a fetal anatomic ultrasound examination (CPT[®] 76811) **due to geographic or other constraints**, a standard (after first trimester) fetal anatomic ultrasound examination (CPT[®] 76805) can be authorized instead.
- CPT[®] 76816 [should not be performed prior to a CPT[®] 76801 or an anatomy scan CPT[®] 76805 (normal pregnancy) or Detailed anatomy scan CPT[®] 76811 (high risk pregnancy)], and is typically not performed prior to 14 weeks gestation.
- Typically all components of the BPP (CPT[®] 76818 and CPT[®] 76819), such as breathing, are not present until ≥ 26 weeks gestation. However, a modified BPP (CPT[®] 76815) can be utilized sooner in certain high risk cases but should not be done prior to 22 weeks.
- SMFM suggest that ductus venosus, middle cerebral artery, or uterine artery Doppler use for routine clinical management of early- or late-onset FGR *is not recommended*
- The minimal use of color Doppler alone (CPT[®] 93976), when performed for anatomical structure identification, during a standard ultrasound procedure, is not separately reimbursable.

References (OB-1)

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Uncertain Dates (OB-2)

Guideline	Page
Uncertain Dates/Unknown Last Menstrual Period (LMP) (OB-2.1).....	19
References (OB-2).....	20

Uncertain Dates/Unknown Last Menstrual Period (LMP) (OB-2.1)

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- If there is a difference in the clinical size of the uterus on pelvic exam and the estimated gestational age calculated by the LMP OR there is an uncertain/unknown LMP OR there have been irregular periods in the last year, one of the following can be performed:
 - If <14 weeks by pelvic exam CPT[®] 76801 one time (plus CPT[®] 76802 for each additional fetus) and/or CPT[®] 76817 one time if a complete ultrasound has not yet been performed
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 **and/or** CPT[®] 76817 for a transvaginal ultrasound is indicated
 - If ≥14 weeks by abdominal exam CPT[®] 76815 or CPT[®] 76805 (CPT[®] 76811 if high risk) if complete fetal anatomic scan has not yet been performed

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though a fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.¹

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v1.0.2025

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Intrauterine Device (IUD) (OB-3)

Guideline	Page
Locate an Intrauterine Device (IUD) (OB-3.1).....	22
References (OB-3).....	23

Locate an Intrauterine Device (IUD) (OB-3.1)

OB.ID.0003.1.A

v1.0.2025

- CPT[®] 76801 and/or CPT[®] 76817 if <14 weeks and a complete ultrasound has not yet been performed or
- CPT[®] 76815 and/or CPT[®] 76817 if complete ultrasound has already been performed or if ≥14 weeks.
- CPT[®] 76805 (CPT[®] 76811 if otherwise indicated), if ≥16 weeks when an anatomy ultrasound (CPT[®] 76805/CPT[®] 76811) has not yet been performed, and
- 3-D Rendering (CPT[®] 76376/CPT[®] 76377) can be added for suspected retained IUD
- For continued pregnancy with retained IUD⁴ image as per **High Risk Group One – Risk Factors (OB 9.1)**

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

References (OB-3)

v1.0.2025

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Infertility (OB-4)

Guideline	Page
History of Infertility (OB-4.1).....	25
Present Pregnancy with ART Treatment (IVF) (OB-4.2).....	26
Recurrent Pregnancy Loss (OB-4.3).....	27
References (OB-4).....	28

History of Infertility (OB-4.1)

OB.IF.0004.1.A

v1.0.2025

- If the current or a prior pregnancy was conceived using an ovulation induction agent (for example Clomid) and/or by intrauterine insemination (IUI), **or**
- If there is a history of infertility or history of IVF in a **prior** pregnancy
- Report:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 **and/or** CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76805 if ≥16 weeks, when complete fetal anatomic scan has not yet been performed
 - Then, follow normal/low risk imaging See **Fetal Anatomic Scan (OB-7.1)**
- Repeat ultrasound is not usually necessary unless there are new clinical indications

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

Present Pregnancy with ART Treatment (IVF) (OB-4.2)

OB.IF.0004.2.A

v1.0.2025

- If the **current** pregnancy was conceived by IVF, can perform
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 **and/or** CPT[®] 76817 for a transvaginal ultrasound is indicated
 - A detailed first-trimester obstetric ultrasound [requested as CPT[®] 76801 plus CPT[®] 76813 (and CPT[®] 76802 plus CPT[®] 76814 for each additional fetus)] between 12 weeks 0 days and 13 weeks 6 days⁶
 - Detailed Fetal Anatomic Scan CPT[®] 76811 ≥16 weeks
 - Initial Fetal echo CPT[®] 76825 **and/or** CPT[®] 76827 +/- CPT[®] 93325 at ≥16 weeks
 - A growth scan (CPT[®] 76816) is indicated in the third trimester (≥28 weeks)
 - If additional high risk factors are noted, imaging as per high risk factor (OB 9.0)
 - Starting at 36 weeks, weekly BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815)

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

Recurrent Pregnancy Loss (OB-4.3)

OB.IF.0004.3.A

v1.0.2025

- If there is a history of at least 2 consecutive or 3 non-consecutive clinical miscarriages/losses at <20 weeks gestation
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, and/or CPT[®] 76817
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 **and/or** CPT[®] 76817 for a transvaginal ultrasound is indicated
 - Detailed Fetal Anatomic Scan CPT[®] 76811 ≥16 weeks
 - Starting at 22 weeks follow-up growth scans (CPT[®] 76816) every 3 to 6 weeks
 - Starting at 36 weeks, weekly BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815)
- If history of one or more 2nd trimester loss (14 to 24 weeks gestation)
 - CPT[®] 76815 **and/or** CPT[®] 76817 every 2 weeks from 16 to 24 weeks.⁵ See **Cervical Insufficiency (OB 18.1)**

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

References (OB-4)

v1.0.2025

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Vaginal Bleeding and/ or Abdominal/Pelvic Pain/Cramping (OB-5)

Guideline	Page
Acute Abdominal/Pelvic Pain (OB-5.1).....	30
Vaginal Bleeding (OB-5.2).....	31
Ectopic Pregnancy (OB-5.3).....	32
Spontaneous Abortion/Threatened/Missed Abortion (OB-5.4).....	34
Hydatidiform Mole (OB-5.5).....	35
References (OB-5).....	36

Acute Abdominal/Pelvic Pain (OB-5.1)

OB.AP.0005.1.A

v1.0.2025

For acute abdominal/pelvic pain:

At the time of complaint, can perform:

- Complete first trimester ultrasound CPT[®] 76801 and/or CPT[®] 76817 if complete ultrasound has not yet been performed, and <14 weeks **or**
- CPT[®] 76815 and/or CPT[®] 76817 **or**
- CPT[®] 76805 (CPT[®] 76811 if otherwise indicated), if ≥14 weeks when an anatomy ultrasound (CPT[®] 76805/CPT[®] 76811) has not yet been performed **or**
- CPT[®] 76816 (if an anatomy ultrasound CPT[®] 76805 or CPT[®] 76811 has previously been performed and at least 2 weeks since anatomy ultrasound)
- Repeat ultrasound is not usually necessary unless there are new indications.
- Note: Above imaging for acute onset abdominal-pelvic pain, NOT for contraction pain/rule out labor. See **Current Preterm Labor (OB-18.3)**

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

Vaginal Bleeding (OB-5.2)

OB.AP.0005.2.A

v1.0.2025

First Trimester

At the time of complaint, can perform:

- Complete first trimester ultrasound CPT[®] 76801 and/or CPT[®] 76817 if complete ultrasound has not yet been performed, and <14 weeks **or**
- CPT[®] 76815 and/or CPT[®] 76817
- Repeat ultrasound is not usually necessary unless there are new indications.

Second and Third Trimesters

At the time of complaint, can perform:

- CPT[®] 76815 and/or CPT[®] 76817 **or**
- CPT[®] 76805/CPT[®] 76811 if indicated (plus CPT[®] 76812 for each additional fetus) if ≥14 weeks, if fetal anatomic scan has not yet been performed, and/or CPT[®] 76817 **or**
- CPT[®] 76816 and/or CPT[®] 76817 if fetal anatomy scan CPT[®] 76805/CPT[®] 76811 has been performed
- Plus CPT[®] 93976 (limited duplex scan) if requested [See **Placental and Cord Abnormalities (OB-21)**].
- BPP (CPT[®] 76818 or CPT[®] 76819) can be considered starting at 26 weeks.
- Repeat ultrasound is not usually necessary unless there are new indications
- For suspected placental abruption, See **Suspected Abruption Placentae (OB-21.5)**
- CPT[®] 76821 if vaginal bleeding with +KB (Kleihauer-Betke) (if feto-maternal hemorrhage – at risk for fetal anemia and hydrops)

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

Ectopic Pregnancy (OB-5.3)

OB.AP.0005.3.A

v1.0.2025

Ectopic Pregnancy

First Trimester

- If there is a history of an ectopic pregnancy or
- If there are abnormally rising hCG titers (non-doubling hCG), or
- If there are signs or symptoms of ectopic pregnancy, e.g. pain and/or bleeding.
- Report:
 - Complete first trimester ultrasound CPT® 76801 and/or CPT® 76817 if complete ultrasound has not yet been performed, and is <14 weeks **or**
 - CPT® 76815 and/or CPT® 76817
 - Plus Color Doppler ultrasonography (CPT® 93976) if an adnexal mass is confirmed
 - If a cornual or (interstitial) ectopic or C-section scar ectopic pregnancy is suspected^{5,6}
 - CPT® 76376 or CPT® 76377, and/or CPT® 93976 as add-on codes
 - MRI Pelvis without contrast (CPT® 72195) if ultrasound is inconclusive.
 - See **3D and 4D Rendering (OB 28.12)** and **Previous C-section or History of Uterine Scar (OB 24.1)**
- If ectopic pregnancy is being treated non-surgically with Methotrexate:
 - If symptomatic, see **Vaginal Bleeding (OB-5.2)** and/or **Acute Abdominal/Pelvic Pain (OB-5.1)** or the imaging guidelines above for ectopic pregnancy
 - If bHCG is not declining appropriately with treatment, see the imaging guidelines above for ectopic pregnancy

Background and Supporting Information

- Cornual or (interstitial) pregnancies pose a significant high morbidity/mortality risk due to massive intraperitoneal bleeding, and are often difficult to diagnose. Conventional sonography still remains the primary diagnostic tool, but 3D US and MRI are being utilized more frequently to aid in earlier detection and treatment.
- CPT® 76815 should never be reported with complete studies CPT® 76801/CPT® 76802, CPT® 76805/CPT® 76810 or CPT® 76811/CPT® 76812 or with CPT® 76816 or BPP (CPT® 76818 and CPT® 76819).

- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation

Spontaneous Abortion/Threatened/ Missed Abortion (OB-5.4)

OB.AP.0005.4.A

v1.0.2025

Spontaneous Abortion/Threatened/Missed Abortion

- To evaluate for threatened or missed abortion:
 - Complete first trimester ultrasound CPT[®] 76801 and/or CPT[®] 76817 if complete ultrasound has not yet been performed, and is <14 weeks **or**
 - CPT[®] 76815 and/or CPT[®] 76817 **or**
 - CPT[®] 76805/CPT[®] 76811 if indicated (plus CPT[®] 76812 for each additional fetus), if ≥14 weeks when complete fetal anatomic scan has not yet been performed, and/or CPT[®] 76817
 - Repeat ultrasound (CPT[®] 76815 and/or CPT[®] 76817) can be repeated weekly if hCG is rising or not falling, or if unable to confirm a viable IUP (fetal pole with cardiac activity)
 - Ultrasound imaging can be repeated earlier than seven days if there are new symptoms
- For complete spontaneous abortion, ultrasound is generally not indicated if there is no pain, or ongoing bleeding, and hCG levels are decreasing.

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation

Hydatidiform Mole (OB-5.5)

OB.AP.0005.5.A

v1.0.2025

Hydatidiform Mole

First, Second and Third Trimester

- Ultrasound can be performed for diagnosis of hydatidiform mole
 - Complete first trimester ultrasound CPT[®] 76801 and/or CPT[®] 76817 if complete ultrasound has not yet been performed, and is <14 weeks, **or**
 - CPT[®] 76815 and/or CPT[®] 76817 **or**
 - CPT[®] 76805/CPT[®] 76811 if indicated (plus CPT[®] 76812 for each additional fetus) if ≥14 weeks, when complete fetal anatomic scan has not yet been performed, and/or CPT[®] 76817
 - CPT[®] 76830 and CPT[®] 76856 or CPT[®] 76857 if hCG titers are not decreasing as expected, or are increasing following treatment, or if there is onset of pain despite falling hCG titers. See **Molar Pregnancy and GTN (PV-16.1)** in the Pelvis Imaging Guidelines
- History of a molar pregnancy, can perform:
 - Complete first trimester ultrasound CPT[®] 76801 if <14 weeks, **or**
 - CPT[®] 76815 and/or CPT[®] 76817 **or**
 - CPT[®] 76805/CPT[®] 76811 if indicated (plus CPT[®] 76812 for each additional fetus) if ≥14 weeks, when complete fetal anatomic scan has not yet been performed, and/or CPT[®] 76817

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

References (OB-5)

v1.0.2025

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Fetal Aneuploidy and Anomaly Screening (OB-6)

Guideline	Page
First Trimester Screening (OB-6.1).....	38
Second Trimester Screening (OB-6.2).....	40
References (OB-6).....	41

First Trimester Screening (OB-6.1)

OB.FA.0006.1.A

v1.0.2025

- First trimester screening includes the assessment of biochemical markers and fetal nuchal translucency (NT) (CPT[®] 76813). An increased Fetal Nuchal Translucency, defined as a NT ≥ 3.0 mm, or $>95^{\text{th}}$ percentile for the crown rump length (CRL), may indicate a fetus with aneuploidy (e.g. Down's syndrome, Trisomy 18) but may also indicate an increased risk for cardiac defects or other structural defects or genetic syndromes in euploid fetuses.
- Nuchal translucency can be performed if CRL 44-83 mm (typically between 10 4/7 and 14 weeks' gestation).
- Indications for a detailed first-trimester fetal anatomic ultrasound [requested as CPT[®] 76801 plus CPT[®] 76813 (and CPT[®] 76802 plus CPT[®] 76814 for each additional fetus)] See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)**

First Trimester Screening:

- Ultrasound CPT[®] 76813 (plus CPT[®] 76814 for each additional fetus) is the initial imaging for first trimester screening, to evaluate fetal nuchal translucency
- If increased Fetal Nuchal Translucency (NT ≥ 3.0 mm or $>95^{\text{th}}$ percentile for the CRL), perform:
 - A detailed first-trimester obstetric ultrasound [requested as CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus)] between 12 weeks 0 days and 13 weeks 6 days,⁹ **and/or**
 - Fetal anatomic ultrasound (CPT[®] 76811) at ≥ 16 weeks
 - Fetal echo (CPT[®] 76825 and/or CPT[®] 76827 and/or CPT[®] 93325) at ≥ 16 weeks
 - Cell-Free DNA (cfDNA), Amniocentesis or CVS can be performed
 - See **High Risk Pregnancy (OB-9)** and **Fetal Echocardiography - Indications for Fetal Conditions (OB-12.2)**
- Fetal NT (CPT[®] 76813) is NOT recommended if cfDNA has already been planned or performed, as they are both screening tools for fetal aneuploidy.
 - Twins and higher order multiples are an exception to this since the sensitivity of cfDNA screening may not be as accurate in this group.¹⁰
 - Fetal NT (CPT[®] 76813) can be performed in twins and higher order multiples even if cfDNA has already been planned or performed. See **Known Dichorionic Multiple Gestations (OB-11.2)** and **Known Monochorionic-Diamniotic or Monochorionic-Monoamniotic Multiple Gestations (OB-11.3)**

- Cell-Free DNA (cfDNA) can be performed any time after 10 weeks gestation and is currently the most sensitive screening test for Down's syndrome per the American College of Medical Genetics and Genomics (99% accurate).
- Those with a positive cfDNA should be offered diagnostic testing (amniocentesis or CVS) and a detailed first-trimester obstetric ultrasound [requested as CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus)] between 12 weeks 0 days and 13 weeks 6 days,⁹ and a detailed anatomy scan (CPT[®] 76811) at ≥16 weeks. See **High Risk Group One – Risk Factors (OB-9.1)**.
 - A “no call” or indeterminate result can occur (risk is higher with maternal obesity), which also has a higher risk of aneuploidy. These individuals should be managed as if positive.

Background and Supporting Information

- CPT[®] 76801 **plus** CPT[®] 76813 [and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus)] when billed together, can also be used to report a detailed late first-trimester obstetric ultrasound examination – performed between 12 weeks 0 days and 13 weeks 6 days
 - This indication-driven detailed first trimester fetal anatomic evaluation is generally performed by those with special skills to perform this study, such as a Maternal Fetal Medicine specialist (Perinatologist), or a Radiologist with advanced training in fetal imaging. It can be performed even if cfDNA has been planned or performed.
- CPT[®] 76813/CPT[®] 76814 for first trimester screening alone, can be performed once per pregnancy, and should be performed only by those certified by the Fetal Medicine Foundation or Nuchal Translucency Quality Review Program (NTQR).
- The use of ultrasound codes (CPT[®] 76801/CPT[®] 76802) should be indication driven and should NOT be routinely done whenever an ultrasound for nuchal translucency (CPT[®] 76813/CPT[®] 76814) is requested. In cases where there is either a maternal and/or fetal indication, then the CPT[®] 76801/CPT[®] 76802 code can indeed be billed along with the nuchal translucency screening (CPT[®] 76813/CPT[®] 76814).
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

Second Trimester Screening (OB-6.2)

OB.FA.0006.2.A

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Second Trimester Screening:

- A fetal anatomy ultrasound (CPT[®] 76805) and/or QUAD screen can be performed during the second trimester to detect fetal aneuploidy, neural tube defects, and other anatomical defects.
 - See **Fetal Anatomic Scan (OB-7.1)**
- If the quad screening is abnormal, a detailed anatomy ultrasound (CPT[®] 76811) can be performed.

Background and Supporting Information

Multiple marker screening is used in the second trimester (15 to 22 6/7 weeks) to screen for aneuploidy as well as open neural tube defects (ONTD).

- Maternal serum alpha-fetoprotein (MSAFP) can be done at 15 to 20 weeks to screen for neural tube defects in those that have had cfDNA or NT screen.
- The “quad” screen (AFP (alpha-fetoprotein), hCG (human chorionic gonadotropin), uE (Unconjugated estriol), dimeric inhibin-A) is the most commonly used test for the second trimester.
- A penta screen (quad screen markers + hyperglycosylated hCG) may be done in lieu of a quad screen.
- Combined, integrated or sequential screening (first and second trimester screening) may also be used and provides a higher detection rate than a single screening.
- Providers often wait for the results of the quad screen before ordering CPT[®] 76805. If the quad screen is abnormal, they may request CPT[®] 76811 in lieu of CPT[®] 76805.

References (OB-6)

v1.0.2025

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Fetal Anatomic Scan/ Cervical Length Screening (OB-7)

Guideline	Page
Fetal Anatomic Scan (OB-7.1).....	43
Fetal Anatomic Scan – Follow-up (OB-7.2).....	44
Cervical Length Screening (OB-7.3).....	45
References (OB-7).....	46

Fetal Anatomic Scan (OB-7.1)

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v1.0.2025

- Though fetal anatomy survey (CPT® 76805/CPT® 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation. This timing allows for a survey of fetal anatomy and an accurate estimation of gestational age.
 - For a **normal/low risk** pregnancy, report a fetal anatomy ultrasound CPT® 76805 if ≥16 weeks.
 - If **high risk** indication is met can report:
 - A detailed fetal anatomy ultrasound (CPT® 76811) if ≥16 weeks^{15,16}
 - These high risk scans indication driven and generally performed by a Maternal Fetal Medicine (MFM) specialist/Perinatologist, or a Radiologist at an AIUM or ACR accredited facility.
 - See **High Risk Pregnancy (OB-9)**
 - For a detailed first-trimester fetal anatomy ultrasound [requested as CPT® 76801 plus CPT® 76813 (and CPT® 76802 plus CPT® 76814 for each additional fetus)] between 12 weeks 0 days and 13 weeks 6 days See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)**

Fetal Anatomic Scan – Follow-up (OB-7.2)

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- Follow-up ultrasounds (CPT[®] 76815 to assess a single item or CPT[®] 76816 if multiple areas to be assessed) can be performed once for incomplete or equivocal finding on initial fetal anatomic scan. This can be performed at any time after incomplete anatomy scan.
- CPT[®] 76816 (should not be performed prior to a CPT[®] 76801 or an anatomy scan CPT[®] 76805 (normal pregnancy) or Detailed anatomy scan CPT[®] 76811 (high risk pregnancy)
- If pregnancy is high risk See **High Risk Pregnancy (OB-9)** or other applicable high risk guideline.
- Detailed anatomy ultrasound CPT[®] 76811 can be performed (if not previously performed) when initial fetal anatomic scan CPT[®] 76805 is abnormal. See **High Risk Pregnancy (OB-9)**
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).

Cervical Length Screening (OB-7.3)

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- Current ACOG and SMFM guidelines state that the method of CL screening in singleton gestations without a prior spontaneous PTB cannot yet be universally mandated. Initial cervical length screening by transabdominal or transperineal ultrasound may limit the need for transvaginal exams
 - Transvaginal ultrasound (CPT[®] 76817) can be performed if:
 - The transabdominal cervical length (CL) appears shortened (≤ 3.6 cm) and transvaginal exam would be needed to guide therapeutic recommendations.
 - If documented screening transabdominal attempt fails to visualize cervix, then CPT[®] 76817 may be performed on a case by case basis.
 - If cervical shortening is identified – See **Cervical Insufficiency (OB-18.1)**
- SMFM currently recommends against routine use of progesterone, pessary or cerclage for the treatment of cervical shortening in twin gestations outside of the context of a clinical trial.
 - Therefore serial measurements of cervical length in multiple gestations is not recommended.

References (OB-7)

v1.0.2025

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Third Trimester Imaging (OB-8)

Guideline	Page
Third Trimester Imaging – Ultrasound (OB-8.1).....	48
Reference (OB-8).....	49

Third Trimester Imaging – Ultrasound (OB-8.1)

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- Imaging in the third trimester is indicated for bleeding, pain, absent fetal heart tones, decreased fetal movement and/or other high-risk indications
 - See specific guidelines based on indication
- For suspected breech position, See **Abnormal Fetal Position/Presentation (OB-14)**

Reference (OB-8)

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1. Practice Bulletin No. 175: Ultrasound in Pregnancy. *Obstetrics & Gynecology*. 2016;128(6):e241-e256. Reaffirmed 2022 doi:10.1097/AOG.0000000000001815

High Risk Pregnancy (OB-9)

Guideline	Page
High Risk General Information (OB-9.0).....	51
High Risk Group One – Risk Factors (OB-9.1).....	53
High Risk Group Two - Ultrasound Findings (OB-9.2).....	59
High Risk Group Three – Pre-pregnancy BMI ≥ 30 kg/m ² (OB-9.3).....	61
High Risk Group Four – Macrosomia (OB-9.4).....	64
High Risk Group Five – Zika and COVID-19 Virus (OB-9.5).....	65
High Risk Group Six - Pre-Gestational or Early Diagnosed (≤ 20 weeks) Diabetes (OB-9.6).....	68
High Risk Group Seven Gestational Diabetes (OB-9.7).....	70
Hypertensive Disorders in Pregnancy (OB-9.8).....	72
History of Spontaneous Pre-Term Delivery/History of PPRM (OB-9.9).....	76
History of Stillbirth (OB-9.10).....	77
Short Interval Pregnancy (≤ 18 months between last delivery and conception of current pregnancy) (OB-9.11).....	78
Detailed First Trimester Fetal Anatomic Scan (OB-9.12).....	79
References (OB-9).....	80

High Risk General Information (OB-9.0)

OB.HR.0009.0.A

v1.0.2025

High Risk Pregnancy General Information:

- Though CPT[®] 76811 can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- The detailed first trimester ultrasound and the detailed fetal anatomic evaluation are indication driven and generally performed by those with special skills to perform this study, such as a Maternal Fetal Medicine specialist (Perinatologist), or a Radiologist with advanced training in fetal imaging. See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)**
- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform the a fetal anatomic ultrasound examination (CPT[®] 76811) **due to geographic or other constraints**, a standard (after first trimester) fetal anatomic ultrasound examination (CPT[®] 76805) can be authorized instead.
- CPT[®] 76805, CPT[®] 76810, CPT[®] 76811, and CPT[®] 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication and/or change in condition.
- Current ACOG and SMFM guidelines state that the method of CL screening in singleton gestations without a prior spontaneous PTB cannot yet be universally mandated. Initial cervical length screening by transabdominal or transperineal ultrasound may limit the need for transvaginal exams.
 - Transvaginal ultrasound (CPT[®] 76817) can be performed if:
 - The transabdominal cervical length (CL) appears shortened (≤ 3.6 cm) and transvaginal exam would be needed to guide therapeutic recommendations.
 - If documented screening transabdominal attempt fails to visualize cervix, then CPT[®] 76817 may be performed on a case by case basis. See **Cervical Length Screening (OB-7.3)**.
 - If cervical shortening is identified – See **Cervical Insufficiency (OB-18.1)**
 - SMFM currently recommends against routine use of progesterone, pessary, or cerclage for the treatment of cervical shortening in twin gestations outside of the context of a clinical trial.
 - Therefore, serial measurements of cervical length in multiple gestations in not recommended.

- CPT[®] 76816 (should not be performed prior to a CPT[®] 76801 or an anatomy scan CPT[®] 76805 (normal pregnancy) or Detailed anatomy scan CPT[®] 76811 (high risk pregnancy))
- Typically all components of the BPP (CPT[®] 76818 and CPT[®] 76819), such as breathing, are not present until ≥ 26 weeks gestation. However, a modified BPP (CPT[®] 76815) can be utilized sooner in certain high risk cases but should not be done prior to 22 weeks. .
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- SMFM suggest that ductus venosus, middle cerebral artery, or uterine artery Doppler use for routine clinical management of early- or late-onset FGR *is not recommended*

High Risk Group One – Risk Factors (OB-9.1)

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High Risk Pregnancy – Risk Factors:

Socio-Demographic Risk Factors (maternal age)

- Age ≥35 years of age at the estimated date of confinement (EDC)

Lifestyle Related Risk Factors (legal or illicit drug/alcohol use)

- Recreational drug (e.g. cocaine, amphetamines, opiates) or excessive alcohol use (≥5 drinks per week) during current pregnancy, or excessive (at least weekly) Marijuana/Cannabinoids/THC use in 2nd and/or 3rd trimesters
 - For 1st trimester marijuana exposure – See **Potentially Teratogenic Medications/ Substances (OB-10.1)**
- Nicotine (≥10 cigarettes a day)
- Other nicotine exposure in pregnancy (e-cigs, vaping, chewing, patch) are also high risk
- Current Maternal IV drug use
- Current use of Suboxone, Subutex, Methadone
- Other polysubstance use

See **Medications/ Exposures Associated with Poor Pregnancy Outcome (OB 10.2)** for imaging recommendations for other high-risk medication or substances not listed above

Health Condition Related Risk Factors (maternal diseases or conditions)

- Anemia severe, <8 grams Hgb or 24% HCT
- Antiphospholipid Syndrome
- Asthma (poorly controlled or steroid dependent)
- Autoimmune disease (e.g. Multiple Sclerosis, Immune Thrombocytopenic Purpura)
- Bariatric surgery

Health Condition Related Risk Factors (maternal diseases or conditions)

- Connective tissue disorders (lupus, RA, scleroderma, Sjogren's, etc.)
- DVT/PE or Maternal thrombophilia (Antiphospholipid Syndrome, Factor V Leiden mutation, Antithrombin III deficiency, Protein C/Protein S deficiency, Prothrombin gene mutation etc.)
- Heart disease (Maternal) – World Health Organization (WHO) Class II or greater
- Hemoglobinopathies (e.g. sickle cell disease, Alpha and Beta thalassemia minor (trait) or major)
- History of endometrial ablation or Uterine Artery embolization
- History of solid organ transplant and/or history of solid organ donation
- Inflammatory Bowel Disease (Ulcerative colitis, Crohn's Disease)
- Liver disease e.g. Hepatitis, Cholestasis of pregnancy (see imaging below)
- Malignancy complicating pregnancy
- Maternal malnutrition (BMI <18.5) or poor weight gain in pregnancy (1 lb per week in 2nd or 3rd trimester if BMI ≤24.9)
- PKU
- Renal disease e.g. glomerulonephritis, persistent protein in the urine, renal insufficiency
- Seizure disorders – on antiepileptic medication
- Thyroid disorder (e.g. hyperthyroidism, poorly controlled hypothyroidism)

Previous pregnancy related risk factors

- Prior pregnancy with adverse outcome (e.g. severe or early onset preeclampsia ≤34 weeks, abruption, accreta, previous uterine dehiscence or rupture, nonimmune hydrops).
- Prior pregnancy with SGA (baby weighing <2500 grams (5.5 pounds/5 lbs 8 oz) at term or less than the 10th percentile of expected weight) or FGR at any gestational age.
- For stillbirth See: **History of Stillbirth (OB-9.10)**

Current pregnancy related risk factors

- Abnormal 1st or 2nd trimester screen (e.g. Abnormal MSAFP; Low PAPP_A; Elevated inhibin A, elevated hCG³¹)
- Known chromosomal abnormalities or abnormal cfDNA
- Genetic Carrier status e.g., Cystic Fibrosis/Known carrier of Spinal Muscular Atrophy (SMA), CF, Tay-Sachs genetic diseases
- Major Fetal anomaly such as gastroschisis, fetal ventriculomegaly, moderate or severe fetal urinary tract dilation defined as ≥ 7 mm at < 28 weeks or > 9 mm at ≥ 28 weeks³², achondroplasias, fetal congenital heart disease, neural tube defect, etc. For sustained fetal arrhythmias. See **Other Causes of Fetal Anemia (OB-16.5)** For Persistent Right Umbilical Vein (PRUV) - See **Persistent Right Umbilical Vein (PRUV) (OB-21.2)**.
- Grand multiparity: must have completed 5 or more pregnancies of greater than 20 weeks gestation, living or stillbirth (does not include current pregnancy; twins count as 1 pregnancy)
- Abnormal Fetal Nuchal Translucency ≥ 3.0 mm or above the 95th percentile for the CRL
- No prenatal care prior to the third trimester
- Short inter-pregnancy interval ≤ 6 months from delivery to conception.^{33,34} [For inter-pregnancy interval > 6 to 18 months – See **Short Interval Pregnancy (≤ 18 months between last delivery and conception of current pregnancy) (OB-9.11)**]
- Pregnancy with retained IUD

Maternal Infections (not exposure)

- Acquired Immune Deficiency Syndrome/HIV Positive
- Chicken Pox/Varicella
- Cytomegalovirus (CMV)
- Malaria
- Known parvovirus in current pregnancy post fetal treatment. See **Exposure to Parvovirus B-19 (OB-16.2)**
- Rubella
- Syphilis, untreated

Maternal Infections (not exposure)

- Toxoplasmosis
- Tuberculosis
- For Zika Virus and COVID-19 Virus See **High Risk Group Five: Zika and COVID-19 Virus (OB-9.5)**

Imaging For Above Conditions

- Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 **and/or** CPT[®] 76817 for a transvaginal ultrasound is indicated
 - See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)** for indications for detailed first trimester fetal anatomic evaluation^{5,6}
- Detailed Fetal Anatomic Scan CPT[®] 76811 if ≥16 weeks
 - Though CPT[®] 76811 can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
- Starting at 22 weeks follow-up growth scans (CPT[®] 76816) every 3 to 6 weeks
- BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815), weekly starting at 32 weeks
- More frequent antepartum fetal surveillance can be performed as stipulated below:
 - Starting at 32 weeks, perform BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815) **up to 2x weekly** for the conditions below:
 - Antiphospholipid Syndrome
 - Maternal Renal Disease (moderate to severe with creatinine >1.4mg/dl)
 - Sickle cell disease
 - Starting **at diagnosis** perform BPP (CPT[®] 76818 or CPT[®] 76819) if ≥26 weeks, or modified BPP (CPT[®] 76815) if ≥22 weeks, **up to 2x weekly**:
 - Intra-hepatic cholestasis of pregnancy (IHCP)
 - Complicated Sickle cell disease (e.g. co-existing hypertension, vaso-occlusive crisis, fetal growth restriction)
 - Complicated SLE (e.g. active lupus nephritis, or recent flares)
 - Major fetal anomaly in the current pregnancy (e.g. gastroschisis, fetal ventriculomegaly, fetal hydronephrosis (>10mm), achondroplasias, fetal congenital heart disease, neural tube defect, sustained fetal arrhythmias)

Background and Supporting Information

Studies that note lower birth weights among offspring exposed to marijuana have noted that these findings were more pronounced among females who used more marijuana, particularly during the first and second trimesters (at least weekly during the pregnancy). CPT® 76811 can be performed, however, given the limited evidence for antenatally detected abnormal growth, serial growth ultrasounds is not indicated in the absence of other findings concerning for growth restriction.

Modified WHO Classification of Heart Disease in Pregnancy
WHO I: Uncomplicated pulmonary stenosis (PS), mitral valve prolapse (MVP), well repaired patent ductus arteriosus (PDA) or persistent anomalous pulmonary venous return (PAPVR)
WHO II: Unrepaired ASD/VSD, repaired tetralogy of fallot (TOF), most arrhythmias
WHO II-II: Mild LV impairment, hypertrophic cardiomyopathy (HCM), valvular heart disease, Marfan without aortic dilation, bicuspid aortic valve (BAV) with Ao <4.5 cm, repaired coarctation of the aorta
WHO III: Mechanical valves, systemic right ventricle (RV), Fontan, cyanotic heart disease, Marfan Aorta 4.0-4.5 cm, Bicuspid Aortic Valve Aorta 4.5-5.0 cm
WHO IV: Pregnancy Contraindicated – native severe coarctation of the aorta, Pulmonary Hypertension, LVEF <30%, NYHA III-IV, severe symptomatic mitral stenosis (MS), Marfan Aorta >4.5; Bicuspid aortic valves Ao >5.0; prior peripartum cardiomyopathy with residual LV impairment.

Recommended Weight Gain During Pregnancy			
Pre-pregnancy weight Category	BMI	Total Wt Gain (lbs)	Recommended Rate of gain in 2 nd and 3 rd trimester (lb/wk)
Underweight	<18.5	28-40	1
Normal weight	18.5-24.9	25-35	1
Overweight	25-29.99	15-25	0.6

Recommended Weight Gain During Pregnancy			
Obese	>30	11-20	0.5

Modified from. ACOG Practice Bulletin No 230. Obesity in Pregnancy, *Obstetrics & Gynecology*: June 2021

High Risk Group Two - Ultrasound Findings (OB-9.2)

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Soft Markers for Aneuploidy (OB-9.2.1)

- If the following 'soft markers' are found in fetus of current pregnancy on routine imaging:
 - Shortened long bones (femur and/or humerus),
 - Mild Pyelectasis^{28,32} (4 mm to <7 mm at 16 to 27 weeks; or 7 mm to <9 mm at ≥28 weeks)
 - For moderate or severe fetal urinary tract dilation (≥7mm at <28 weeks or >9 mm at ≥28 weeks) See **High Risk Group One - Risk factors (OB-9.1)**
 - Echogenic bowel
 - See **Other Ultrasound Findings (OB-9.2.2)** for Fetal Echogenic intra-cardiac focus and/or choroid plexus cyst
- Detailed Fetal anatomic scan ≥16 weeks (CPT[®] 76811).
- One follow-up scan (CPT[®] 76816) in third trimester

Other Ultrasound Findings (OB-9.2.2)

- If an isolated soft marker²⁸ is found in fetus of current pregnancy on routine imaging, including:
 - Choroid plexus cyst, or
 - Echogenic intra-cardiac foci, or
 - Thickened nuchal fold (≥6mm at 15 to 20 weeks), or
 - Absent or hypoplastic nasal bone
 - Report:
 - Detailed fetal anatomic scan (CPT[®] 76811) at ≥16 weeks
- If a Major fetal anomaly is found, or if abnormal cfDNA or amniocentesis results - See **High Risk Group One - Risk factors (OB-9.1)**
- If negative cfDNA or negative amniocentesis – Fetal echo or follow-up ultrasound are not warranted for these isolated findings

- If history of a prior pregnancy with a chromosomal and/or structural congenital anomaly, or
- Current pregnancy with suspected fetal anomaly on initial imaging
- Report:
 - A detailed first-trimester obstetric ultrasound^{5,6} [requested as CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus)] between 12 weeks 0 days and 13 weeks 6 days, **and/or**
 - Detailed fetal anatomic scan (CPT[®] 76811) at ≥16 weeks
- If a Major fetal anomaly is found - See **High Risk Group One - Risk factors (OB-9.1)** and/or **Fetal Echocardiography - Indications for Fetal Conditions (OB 12.2)**

High Risk Group Three – Pre-pregnancy

BMI ≥ 30 kg/m² (OB-9.3)

OB.HR.0009.3.A

v1.0.2025

Pre-pregnancy BMI ≥ 30 kg/m²

- Complete first trimester anatomy ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- A detailed first trimester fetal anatomic evaluation^{5,6} can be performed between 12 weeks 0 days and 13 weeks 6 days. (See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)** for indications)
 - Report: CPT[®] 76801 plus CPT[®] 76813 (and CPT[®] 76802 plus CPT[®] 76814 for each additional fetus) for a detailed first trimester fetal anatomy ultrasound.
 - A detailed first trimester fetal anatomic evaluation can be performed if indicated, even if cfDNA has been planned or performed
- A detailed fetal anatomic scan at ≥ 16 weeks (CPT[®] 76811)
 - Though CPT[®] 76811 can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation

Class I Obesity - Pre-pregnancy BMI 30 to 34.9 (OB-9.3.1)

Class I Obesity (BMI 30-34.9)

- After first and second trimester imaging as outlined above, Report **One** follow-up scan (CPT[®] 76816) between 32 to 36 weeks
 - If unable to clinically assess fundal height due to body habitus a growth scan (CPT[®] 76816) can be considered in the early third trimester with follow up in 4 weeks

Class II Obesity - Pre-pregnancy BMI 35-39.9 (OB-9.3.2)**Class II Obesity (BMI 35-39.9)**

- After first and second trimester imaging as outlined above, Report:
 - Growth scans (CPT[®] 76816) every 4 weeks starting in the third trimester (≥28 weeks)
 - BPP (CPT[®] 76818 or CPT[®] 76819) or a modified BPP (CPT[®] 76815) weekly starting at 36 weeks

Class III Obesity - Pre-pregnancy BMI ≥40 (OB-9.3.3)**Class III Obesity (BMI ≥40)**

- After first and second trimester imaging as outlined above, Report:
 - Growth scans (CPT[®] 76816) every 4 weeks starting in the third trimester (>28 weeks)
 - BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815) weekly starting at 32 weeks

Background and Supporting Information

If pre-pregnancy or early pregnancy weight is over 200 pounds, it can be presumed that BMI is ≥30 kg/m².

Recommended Weight Gain During Pregnancy

Pre-pregnancy weight Category	BMI	Total Wt Gain (lbs)	Recommended Rate of gain in 2nd and 3rd trimester (lb/wk)
Underweight	<18.5	28-40	1
Normal weight	18.5-24.9	25-35	1
Overweight	25-29.99	15-25	0.6
Obese	>30	11-20	0.5

Modified from ACOG Practice Bulletin No 230. Obesity in Pregnancy, *Obstetrics & Gynecology*: June 2021

High Risk Group Four – Macrosomia (OB-9.4)

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Prior Pregnancy with Macrosomia (OB-9.4.1)

Prior pregnancy with macrosomia (baby weighing >4000 grams at term or greater than the 90th percentile of expected weight)

- Report one of the following in the first trimester to establish dates:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - Detailed Fetal Anatomic Scan CPT[®] 76811 if ≥16 weeks
 - Though CPT[®] 76811 can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
- Follow-up scan (CPT[®] 76816) in the third trimester (28-32 weeks) and at ≥35 weeks to plan for delivery^{35,36}

Current Pregnancy with Suspected or Known Macrosomia (OB-9.4.2)

- See **Macrosomia – Large for Dates Current Pregnancy (OB-20.2)**
- See **Unequal Fundal Size and Dates (OB-27)**

High Risk Group Five – Zika and COVID-19 Virus (OB-9.5)

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Zika Virus (OB-9.5.1)

Zika Virus

Suspected exposure without symptoms³⁷

- Report one of the following:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76805/CPT[®] 76811 if otherwise indicated (plus CPT[®] 76810/CPT[®] 76812 for each additional fetus) if anatomy ultrasound has not yet been performed, **or**
 - CPT[®] 76816 if anatomy ultrasound (CPT[®] 76805/CPT[®] 76811) previously performed
 - Though a fetal anatomy scan can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
 - If test positive or if symptoms developed, See below

Zika Virus

Suspected exposure with symptoms or known infection³⁷

- Report one of the following:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76811 if ≥16 weeks when an anatomy ultrasound (CPT[®] 76811) has not yet been performed
 - Though CPT[®] 76811 can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
 - Growth scan, (CPT[®] 76816) every 3 to 4 weeks to monitor for findings such as intracranial calcifications and microcephaly, starting at 16 weeks.
 - CPT[®] 76816 if anatomy ultrasound (CPT[®] 76805/CPT[®] 76811) previously performed
 - Though a fetal anatomy scan can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
 - If fetal anomaly suspected or diagnosed See **High Risk Group One – Risk Factors (OB-9.1)**, if FGR diagnosed, See **Fetal Growth Restriction Current Pregnancy (OB-20.1)**

COVID-19 Virus (OB-9.5.2)

COVID-19 Virus

COVID-19 infection in the current pregnancy³⁹

- Report one of the following:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76805 (CPT[®] 76811 if otherwise indicated), if ≥16 weeks when an anatomy ultrasound (CPT[®] 76805/CPT[®] 76811) has not yet been performed
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
 - Growth scan (CPT[®] 76816) every 3 to 4 weeks starting at 22 weeks [See **High Risk Group One - Risk factors (OB-9.1)**]
 - Starting **at diagnosis** perform weekly BPP (CPT[®] 76818 or CPT[®] 76819) if ≥26 weeks, or modified BPP (CPT[®] 76815) if ≥22 weeks. (See **High Risk Group One - Risk factors (OB-9.1)**)
 - If FGR diagnosed then follow **FGR imaging Fetal Growth Restriction Current Pregnancy (OB-20.1)**

High Risk Group Six - Pre-Gestational or Early Diagnosed (≤ 20 weeks) Diabetes (OB-9.6)

OB.HR.0009.6.A
v1.0.2025

- If diabetes is diagnosed prior to pregnancy or in the first or early second trimester (typically before 20 weeks gestation) with standard diagnostic criteria of: HbA1C $\geq 6.5\%$, fasting plasma glucose ≥ 126 mg/dL, or 2-hour glucose ≥ 200 mg/dL on a 75-g oral glucose tolerance test, then image as below.

Test	When	Frequency	Codes
First Trimester (Dating) Ultrasound	<14 weeks	Once	CPT [®] 76801 is preferred for dating, but if this is unable to be completed then CPT [®] 76815 and/or CPT [®] 76817 for a transvaginal ultrasound is indicated
Detailed first-trimester obstetric ultrasound ^{5,6}	12 to 13+6 weeks	Once	CPT [®] 76801 plus CPT [®] 76813 (and CPT [®] 76802 plus CPT [®] 76814 for each additional fetus)
Fetal anatomic scan	≥ 16 weeks	Once	CPT [®] 76811
Initial Fetal echo	Starting at ≥ 16 weeks	Once	CPT [®] 76825 and/or CPT [®] 76827 and/or CPT [®] 93325
Ultrasound (for fetal growth)	Starting at 22 weeks	Every 2 to 4 weeks	CPT [®] 76816

Test	When	Frequency	Codes
Biophysical Profile (BPP) or modified BPP	Starting at 32 weeks (can start at ≥26 weeks if complicated by additional risk factors (e.g., FGR Oligohydramnios, HTN)	Up to twice weekly	CPT® 76818 (BPP) or CPT® 76819 (BPP) or CPT® 76815 (modified BPP)
Umbilical artery Doppler (if FGR diagnosed)	Upon diagnosis of FGR if ≥22 weeks	Weekly	CPT® 76820

High Risk Group Seven Gestational Diabetes (OB-9.7)

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v1.0.2025

Gestational Diabetes - Diet-Controlled (GDM-A1) (OB-9.7.1)

If patient has gestational diabetes and it is diet controlled:			
Test	When	Frequency	Codes
Fetal anatomic scan	≥16 weeks	Once	CPT® 76805
Ultrasound (for fetal growth)	Once at the time of diagnosis, then starting at 32 weeks	Every 4 weeks	CPT® 76816
Biophysical Profile (BPP) or modified BPP	Starting at 34 weeks	Once weekly if diet controlled.	CPT® 76818 (BPP) or CPT® 76819 (BPP) or CPT® 76815 (modified BPP)

Gestational Diabetes on Medications (GDM-A2) (OB-9.7.2)

If patient has gestational diabetes and is on oral medication or insulin:			
Test	When	Frequency	Codes
Fetal anatomic scan	≥16 weeks	Once	CPT® 76811
Fetal echo (if HbA1C >6%)	Once in the third trimester (≥32 weeks)	Once	CPT® 76825 and/or CPT® 76827 and/or CPT® 93325

If patient has gestational diabetes and is on oral medication or insulin:			
Test	When	Frequency	Codes
Ultrasound (for fetal growth)	Starting at 22 weeks	Every 2 to 4 weeks	CPT® 76816
Biophysical Profile (BPP) or modified BPP	Starting at 32 weeks (can start at ≥26 if complicated by additional risk factors (e.g., FGR Oligohydramnios))	Up to twice weekly	CPT® 76818 (BPP) or CPT® 76819 (BPP) or CPT® 76815 (modified BPP)
Umbilical artery Doppler (if FGR diagnosed)	Upon diagnosis of FGR if ≥22 weeks	Weekly	CPT® 76820

Background and Supporting Information

- If HbA1c levels are >6%, in those with GDM-A2, fetal echocardiogram in the third trimester to assess for ventricular hypertrophy can be performed.

Hypertensive Disorders in Pregnancy (OB-9.8)

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Screening in High Risk Groups (OB-9.8.1)

Screening in High Risk Groups

- SMFM state that uterine artery Doppler has limited diagnostic accuracy and clinical utility in predicting FGR, SGA birth, and perinatal mortality. As such, its use for screening in high risk groups *is not recommended*.

Current Chronic Hypertension not on Medication (OB-9.8.2)

Test	When	Frequency	Codes
First Trimester (Dating) Ultrasound	<14 weeks	Once	CPT [®] 76801 is preferred for dating, but if this is unable to be completed then CPT [®] 76815 and/or CPT [®] 76817 for a transvaginal ultrasound is indicated
Fetal anatomic scan	≥16 weeks	Once	CPT [®] 76811
Ultrasound (for fetal growth)	In the third trimester (≥28 weeks)	Every 4-6 weeks	CPT [®] 76816
If blood pressure is elevated from baseline, See Gestational Hypertension (GH, preeclampsia, toxemia) (OB-9.8.4) below			

Current Chronic Hypertension on Medication (OB-9.8.3)

Test	When	Frequency	Codes
First Trimester (Dating) Ultrasounds	<14 weeks	Once	CPT [®] 76801 is preferred for dating, but if this is unable to be completed then CPT [®] 76815 and/or CPT [®] 76817 for a transvaginal ultrasound is indicated
Detailed Fetal Anatomic Scan	≥16 weeks	Once	CPT [®] 76811
Ultrasound (for fetal growth)	Starting at 22 weeks gestation	Every 3 to 4 weeks	CPT [®] 76816
Biophysical profile (BPP) or modified BPP	Starting at 32 weeks If complicated by other risk factors (e.g. DM, FGR Oligohydramnios) can start at ≥26 weeks)	Once weekly If complicated by other risk factors (e.g., FGR Oligohydramnios) twice weekly	CPT [®] 76818 (BPP) or CPT [®] 76819 (BPP) or CPT [®] 76815 (AFI)
Umbilical artery Doppler (if FGR diagnosed) See Fetal Growth Restriction Current Pregnancy (OB-20.1)	Upon diagnosis of FGR if ≥22 weeks	Twice weekly	CPT [®] 76820

Gestational Hypertension (GH, preeclampsia, toxemia) (OB-9.8.4)

Test	When	Frequency	Codes
Fetal anatomic scan	≥16 weeks	Once	CPT® 76805 or CPT 76811 if other high risk issues and if not previously completed
Growth US	Starting at time of diagnosis	Every 3 to 4 weeks If FGR, Oligohydramnios or severe preeclampsia (every 2 to 4 weeks)	CPT® 76816
BPP	Starting at time of diagnosis if ≥26 weeks	Up to twice weekly Hypertension/ pre-eclampsia with severe features - Daily	CPT® 76818 or CPT® 76819
Modified BPP	Starting at time of diagnosis if ≥22 weeks	Up to twice weekly Hypertension/pre-eclampsia with severe features - Daily	CPT® 76815
Umbilical artery Doppler <u>Fetal Growth Restriction Current Pregnancy (OB-20.1)</u>	Starting at time of diagnosis of FGR or Oligohydramnios if ≥22 weeks	Twice weekly	CPT® 76820

Background and Supporting Information

Disorder	Definition
Hypertension in pregnancy	Systolic blood pressure ≥ 140 mm Hg or diastolic BP ≥ 90 mm Hg, or both, measured on two occasions at least 4 hours apart
Severe-range hypertension	Systolic blood pressure ≥ 160 mm Hg or diastolic BP ≥ 110 mm Hg, or both, measured on two occasions at least 4 hours apart
Chronic hypertension	Hypertension diagnosed or present before pregnancy or before 20 weeks of gestation; or hypertension that is diagnosed for the first time during pregnancy and that does not resolve in the postpartum period
Chronic hypertension with superimposed preeclampsia	Preeclampsia in a female with a history of hypertension before pregnancy or before 20 weeks of gestation
Gestational hypertension	Hypertension diagnosed after 20 weeks of gestation, in a female with a previously normal blood pressure.
Preeclampsia	Disorder of pregnancy associated with new-onset hypertension, which occurs most often after 20 weeks of gestation and frequently near term. Although often accompanied by new-onset proteinuria, hypertension and other signs or symptoms of preeclampsia may present in some females in the absence of proteinuria.
Eclampsia	Convulsive manifestation of the hypertensive disorders of pregnancy and is among the more severe manifestations of the disease.

History of Spontaneous Pre-Term Delivery/History of PPROM (OB-9.9)

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v1.0.2025

Spontaneous Preterm Delivery or PPROM <37 0/7 Weeks (OB-9.9.1)

Initial Imaging

- For initial imaging:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- Fetal Anatomy Ultrasound CPT[®] 76811 [plus CPT[®] 76812 for each additional fetus] if ≥16 weeks and a complete fetal anatomic scan has not yet been performed during this pregnancy
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- CPT[®] 76817 and/or CPT[®] 76815 every 1 to 4 weeks, starting at ≥16 0/7 weeks until 24 0/7 weeks
- Starting after the fetal anatomic scan at ≥22 weeks, ultrasound (CPT[®] 76816) can be performed every 3 to 6 weeks until delivery
- If history of Preterm Delivery or PPROM below 37 0/7 weeks in immediately preceding pregnancy starting at 32 weeks, weekly BBP CPT[®] 76818 or CPT[®] 76819 or modified BPP CPT[®] 76815
- If funneling or short cervix ≤25 mm (2.5 cm) is found on a transvaginal ultrasound in a singleton pregnancy See **Cervical Insufficiency (OB-18.1)**
- For current preterm labor See **Current Preterm Labor (OB-18.3)**
- If additional risk factors present see **OB 9.1** for indications for fetal monitoring

History of Stillbirth (OB-9.10)

OB.HR.0009.10.A

v1.0.2025

Initial Imaging

- For initial imaging:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, and/or CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - A detailed first trimester fetal anatomic evaluation^{5,6} can be performed between 12 weeks 0 days and 13 weeks 6 days. (See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)** for indications)
 - Report: CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus) for a detailed first trimester fetal anatomy ultrasound
 - A detailed first trimester fetal anatomic evaluation can be performed if indicated, even if cfDNA has been planned or performed
- Fetal anatomic scan at ≥16 weeks (CPT[®] 76811)
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
- Follow up ultrasound (CPT[®] 76816) every 2 to 4 weeks to assess fetal growth starting at 22 weeks or two weeks before prior pregnancy loss.
- Twice weekly BPP (CPT[®] 76818 or CPT[®] 76819, if ≥26 weeks) **or** modified BPP CPT[®] 76815 (not to be performed prior to 22 weeks) starting at 32 weeks or two weeks before prior pregnancy loss

Background and Supporting Information

- A history of stillbirth is not an indication for fetal echo. Per 2020 ACOG bulletin – there is no mention of recommendation for echo – just a detailed anatomy US. If demised fetus had a confirmed cardiac anomaly on autopsy, or if the detailed anatomy scan on either the demised fetus or the current pregnancy had findings suspicious for cardiac anomaly, then echo may be indicated. See **Fetal Echocardiography - Indications for Fetal Conditions (OB 12.2)**

Short Interval Pregnancy (≤ 18 months between last delivery and conception of current pregnancy) (OB-9.11)

OB.HR.0009.11.A

v1.0.2025

If Inter-Pregnancy interval ≤ 6 months ^{33,34}	
Follow imaging as per High Risk Group One – Risk Factors (OB 9.1)	
If Inter-Pregnancy interval >6 months but ≤ 18 months ^{33,34}	
- Report one of the following to establish dates: - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, and/or CPT[®] 76817 for a transvaginal ultrasound - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated - For fetal anatomy scan report CPT[®] 76805 (CPT[®] 76811 if otherwise indicated), if ≥ 16 weeks - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation. - One follow-up scan, CPT[®] 76816 in the third trimester (28-32 weeks) to assess fetal growth.	

Background and Supporting Information

Inter-pregnancy intervals shorter than 18 months are associated with higher risks of adverse pregnancy outcomes, including preterm delivery, small-for-gestational-age (SGA) birth, and infant mortality. Per ACOG, females should be advised to avoid inter-pregnancy intervals shorter than 6 months due to even more significant risks.³³

Detailed First Trimester Fetal Anatomic Scan (OB-9.12)

OB.HR.0009.12.A
v1.0.2025

- A detailed first trimester fetal anatomic evaluation^{5,6} can be performed, if indicated, between 12 weeks 0 days and 13 weeks 6 days.
 - If indicated report: CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus) [requested together] for a detailed first trimester fetal anatomy ultrasound.
 - A detailed first trimester fetal anatomic scan can be requested even if cfDNA has been planned or performed.
- Indications for a detailed first trimester fetal anatomic ultrasound include but are not limited to conditions below

Detailed First-Trimester Fetal Anatomy Scan Indications
◦ Previous fetus or child with a congenital, genetic, or chromosomal anomaly
◦ Known or suspected fetal abnormality detected by ultrasound in the current pregnancy
◦ Fetus at increased risk for a congenital anomaly based on the following: ▪ 35 years or older at delivery ▪ Maternal pre-gestational diabetes ▪ Pregnancy conceived via in vitro fertilization ▪ Multiple gestation ▪ Teratogen exposure ▪ Enlarged nuchal translucency ▪ Positive screening test results for aneuploidy, including cell-free DNA screening and serum-only or combined first-trimester screening
◦ Other conditions possibly affecting the pregnancy/fetus, including: ▪ Maternal body mass index of 30 kg/m or higher ▪ Placental implantation covering the internal cervical os under a cesarean scar site or cesarean scar pregnancy diagnosed in index gestation

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v1.0.2025

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High Risk Medications and Substances (OB-10)

Guideline	Page
Potentially Teratogenic Medications/Substances (OB-10.1)	84
Medications/ Exposures Associated with Poor Pregnancy Outcome (OB-10.2).....	87
References (OB-10).....	90

Potentially Teratogenic Medications/ Substances (OB-10.1)

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v1.0.2025

- If maternal exposure to any of the below Potentially Teratogenic Medications/ Substances report:
 - Complete first trimester anatomy ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - A detailed first trimester fetal anatomic evaluation can be performed between 12 weeks 0 days and 13 weeks 6 days. (See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)** for indications)
 - Report: CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus) for a detailed first trimester fetal anatomy ultrasound
 - A detailed first trimester fetal anatomic evaluation can be performed if indicated, even if cfDNA has been planned or performed
 - CPT[®] 76811 (detailed fetal anatomy) if ≥16 weeks
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation

Potentially Teratogenic Medications/Substances
Aminoglycosides (amikacin, gentamycin, kanamycin, tobramycin, and other mycins)
Aspirin – only if exposed less than 10 weeks gestation
Benzodiazepines [e.g.,Diazepam (valium), Lorazepam (Ativan), Alprazolam (Xanax) etc.]
Codeine
Ergotamine (e.g. Methergine)
Fluconazole (Diflucan)

Potentially Teratogenic Medications/Substances
Lead (Exposure in early pregnancy)
Leflunomide (first trimester exposure)
Marijuana/Cannabinoids/THC Exposure
Methyl mercury
Oral contraceptives (combined and/or progestin only exposure in the first trimester)
Selective serotonin reuptake inhibitors (SSRI) (e.g., Sertraline (Zoloft), Citalopram (Celexa), Fluoxetine (Prozac) etc.)
Serotonin norepinephrine receptor inhibitors (SNRIs) SNRIs (e.g., Venlafexine (Effexor), Duloxetine (Cymbalta), etc.)
Serotonin modulators e.g. Trazadone, etc.
Tricyclics (e.g., Amitriptyline (Elavil), Imipramine (Tofranil) etc.)
Tetracyclines (e.g., Chlortetracycline, Doxycycline, Methacycline, Sumycin, etc.)

- This is **not** an all-inclusive list.
 - See **High Risk Group One – Risk Factors (OB 9.1)** Health Condition Related Risk Factors (maternal diseases or conditions) or other appropriate guideline for indicated imaging based on disease process being treated.
 - See **Medications/Exposures Associated with Poor Pregnancy Outcome (OB 10.2)**
- Studies that note lower birth weights among offspring exposed to marijuana have noted that these findings were more pronounced among females who used more marijuana, particularly during the first and second trimesters (at least weekly during the pregnancy). CPT[®] 76811 can be performed, however, given the limited evidence for antenatally detected abnormal growth, serial growth ultrasounds is not indicated in the absence of other findings concerning for growth restriction.
- If documented excessive use (at least weekly) of Marijuana/Cannabinoids/THC in 2nd and/ or 3rd trimesters - See **High Risk Group One – Risk Factors (OB 9.1)**

- Other atypical antidepressants like Bupropion, Mirtazapine, Nefazodone and Duloxetine have **not** been linked to an increased risk of fetal anomalies or poor pregnancy outcomes.

Medications/ Exposures Associated with Poor Pregnancy Outcome (OB-10.2)

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- If maternal exposure to any of the Medications or Exposures noted below:
 - Complete first trimester anatomy ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 **and/or** CPT[®] 76817 for a transvaginal ultrasound is indicated
 - A detailed first trimester fetal anatomic evaluation can be performed between 12 weeks 0 days and 13 weeks 6 days. (See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)** for indications)
 - Report: CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus) for a detailed first trimester fetal anatomy ultrasound
 - A detailed first trimester fetal anatomic evaluation can be performed if indicated, even if cfDNA has been planned or performed
 - Further imaging as per **High Risk Group One – Risk Factors (OB 9.1)**

Medications/Substances associated with poor pregnancy outcome
Anti-convulsants (e.g., Dilantin, Lamictal, Phenobarbital, Tegretol, Valproate etc.)
Anti-hypertensive Agents (e.g. ACE inhibitors, Angiotensin II Antagonists, Beta Blockers, etc.)
Anti-neoplastic agents (e.g. Daunorubicin etc.)
Anti-psychotics (e.g. Abilify, Haldol, Latuda, Seroquel, Stelazine Thorazine Zyprexa, etc.)
Carbon monoxide
Corticosteroids (e.g. Prednisone, Cortisone, etc.)
Coumadin/ warfarin

Medications/Substances associated with poor pregnancy outcome
Heparin/ Low Molecular Weight Heparin (ongoing use during pregnancy)
Immune Modulating Drugs (e.g. Azathioprine, Belatacept, Cyclophosphamide, Cyclosporin A, Hydroxychloroquine, Hydroxyurea, Leflunomide, Mycophenolate mofetil, Sirolimus/Everolimus, etc.)
Lithium
Methimazole
Methotrexate
Mifepristone (RU486)
Misoprostol
Monoamine oxidase inhibitors e.g. Phenelzine (Nardil)
Penicillamine
Pregabalin (Lyrica), Gabapentin (Neurontin)
Quinine
Retinoic acid/retinoid medications
Stimulants (e.g. Ritalin, adderal, etc. and other stimulants used to treat ADHD)
Thalidomide

- This is **not** an all-inclusive list. See **High Risk Group One – Risk Factors (OB 9.1)** Health Condition Related (maternal diseases or conditions) or other appropriate guideline for indicated imaging based on disease process being treated.

Background and Supporting Information

- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a fetal anatomic ultrasound examination (CPT® 76811) due to geographic or other constraints, a

standard (after first trimester) fetal anatomic ultrasound examination (CPT[®] 76805) can be authorized instead.

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819)
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation

References (OB-10)

v1.0.2025

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Multiple Gestations (OB-11)

Guideline	Page
Suspected Multiple Gestations (OB-11.1).....	92
Known Dichorionic Multiple Gestations (OB-11.2).....	93
Known Monochorionic-Diamniotic or Monochorionic-Monoamniotic Multiple Gestations (OB-11.3).....	96
References (OB-11).....	99

Suspected Multiple Gestations (OB-11.1)

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For Suspected multiple pregnancies:

- Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- CPT[®] 76805 and CPT[®] 76810 for each additional fetus if ≥14 weeks if a dating ultrasound or a complete anatomy ultrasound has not yet been performed during this pregnancy

Known Dichorionic Multiple Gestations (OB-11.2)

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For Known dichorionic multiple pregnancies:

- Complete first trimester anatomy ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- A detailed first trimester fetal anatomic evaluation^{3,4} can be performed between 12 weeks 0 days and 13 weeks 6 days. (See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)** for indications)
 - Report: CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus) for a detailed first trimester fetal anatomy ultrasound
 - A detailed first trimester fetal anatomic evaluation can be performed if indicated, even if cfDNA has been planned or performed
- Fetal NT (CPT[®] 76813) can be performed in twins and higher order multiples even if cfDNA has already been planned or performed.¹ See **First Trimester Screening (OB-6.1)**
- CPT[®] 76811 and CPT[®] 76812 for each additional fetus at ≥16 weeks if a complete detailed anatomic scan (CPT[®] 76811) has not yet been performed
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
- SMFM currently recommends against routine use of progesterone, pessary or cerclage for the treatment of cervical shortening in twin gestations outside of the context of a clinical trial.
 - Therefore serial measurements of cervical length in multiple gestations is not recommended.
- If cervical shortening is identified – See **Cervical Insufficiency (OB-18.1)**
- Growth ultrasound (CPT[®] 76816) can be done every 4 to 6 weeks at ≥14 weeks.
- If otherwise uncomplicated dichorionic twins, perform BPP (CPT[®] 76818 or CPT[®] 76819) **or** modified BPP (CPT[®] 76815) weekly starting at 36 weeks
- If additional risk factors (e.g. diabetes, or hypertensive disease), BPP (CPT[®] 76818 or CPT[®] 76819) **or** modified BPP (CPT[®] 76815) can be performed weekly starting at 32 weeks or sooner (See **High Risk Pregnancy (OB-9)**)

For Known dichorionic multiple pregnancies:

- Fetal loss of one twin during the first trimester does not appear to increase the risk of FGR or preterm delivery in the surviving twin, however, loss of one or more fetus(es) after the first trimester is associated with increased risk for FGR and PTB
 - Dichorionic twin gestations with loss of one fetus after the first trimester (≥ 14 weeks) should be imaged according to **High Risk Group (OB-9.1)**
- If FGR or growth discordance $\geq 20\%$ is diagnosed, can perform:
 - CPT[®] 76816 (growth ultrasound) every 2 to 4 weeks
 - Modified BPP (CPT[®] 76815) up to twice weekly starting at ≥ 22 weeks, or BPP (CPT[®] 76818 or CPT[®] 76819) up to twice weekly starting at ≥ 26 weeks
 - UA Doppler (CPT[®] 76820) weekly (starting at ≥ 22 weeks)
- If Severe FGR (EFW $\leq 3\%$, AC $\leq 3\%$), **OR** Abnormal UA Doppler studies (defined as a PI, RI, or S/D ratio greater than the 95th percentile for gestational age **OR** absent or reversed end-diastolic velocity (AEDV or REDV), **OR** confirmed oligohydramnios:
 - BPP (CPT[®] 76818 or CPT[®] 76819 or CPT[®] 76815) and/or umbilical artery (UA) Doppler (CPT[®] 76820) may be needed more frequently (2-3 times per week, or even daily).
- If IVF dichorionic twins, report an initial fetal echo as CPT[®] 76825 and/or CPT[®] 76827 with or without CPT[®] 93325. Trans-abdominal fetal echo is usually not performed prior to 16 weeks. See **Indications for Maternal Conditions (OB-12.3)**
- If other high risk factors, See **High Risk Pregnancy (OB-9)**

Known Monochorionic-Diamniotic or Monochorionic-Monoamniotic Multiple Gestations (OB-11.3)

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For Known monochorionic-diamniotic or monochorionic-monoamniotic multiple pregnancies

- Complete first trimester anatomy ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- A detailed first trimester fetal anatomic evaluation^{3,4} can be performed between 12 weeks 0 days and 13 weeks 6 days. (See **Detailed First Trimester Fetal Anatomic Scan (OB-9.12)** for indications).
 - Report: CPT[®] 76801 **plus** CPT[®] 76813 (and CPT[®] 76802 **plus** CPT[®] 76814 for each additional fetus) for a detailed first trimester fetal anatomy ultrasound
 - A detailed first trimester fetal anatomic evaluation can be performed if indicated, even if cfDNA has been planned or performed
- Fetal NT (CPT[®] 76813) can be performed in twins and higher order multiples even if cfDNA has already been planned or performed.¹
- CPT[®] 76811 and CPT[®] 76812 for each additional fetus at ≥16 weeks if a complete detailed anatomic scan (CPT[®] 76811) has not yet been performed
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
- Universal cervical length (CL) screening with transvaginal ultrasound (CPT[®] 76817) is NOT recommended in monochorionic twin gestations.
 - Transvaginal ultrasound (CPT[®] 76817) can be performed if the transabdominal cervical length (CL) is ≤3.6 cm. If documented transabdominal attempt fails to visualize cervix, then CPT[®] 76817 may be performed on a case by case basis. See **Cervical Length Screening (OB-7.3)**. If cervical shortening is identified – See **Cervical Insufficiency (OB-18.1)**

For Known monochorionic-diamniotic or monochorionic-monoamniotic multiple pregnancies

- CPT[®] 76816 (growth ultrasound) every 2 to 4 weeks starting at 14 weeks
- Initial Fetal Echo (CPT[®] 76825 and/or CPT[®] 76827) with or without color Doppler (CPT[®] 93325) (usually not performed <16 weeks).
- MCA Doppler (CPT[®] 76821) is indicated every 2 weeks starting at 16 weeks until delivery to monitor for Twin-Twin Transfusions Syndrome (TTTS) and/or Twin Anemia Polycythemia Sequence (TAPS). This can be performed with a limited ultrasound (CPT[®] 76815) or growth ultrasound (CPT[®] 76816).
- Perform BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815) weekly starting at 32 weeks or sooner if additional risk factors (eg. diabetes, or hypertensive disease - See **High Risk Pregnancy (OB-9)**)
- If TTTS is suspected or diagnosed, or if FGR or growth discordance $\geq 20\%$ is diagnosed perform:
 - BPP (CPT[®] 76818 or CPT[®] 76819 (≥ 26 weeks) or CPT[®] 76815) and/or umbilical artery (UA) Doppler (CPT[®] 76820) and/or MCA Doppler (CPT[®] 76821) 2 to 3 times per week (May be needed more frequently on a case-by-case basis)
 - TTTS is diagnosed by the ultrasound findings of polyhydramnios in one twin (the recipient) and oligohydramnios in the other twin (the donor). If AFI is discordant between the twins (low but not < 2 cm in one and/or high but not > 8 cm in the other); weekly imaging (MCA and/or limited US) can be performed
 - If TTTS is diagnosed, follow-up fetal echo (CPT[®] 76826 and/or CPT[®] 76828) with or without color Doppler (CPT[®] 93325) can also be performed as requested. See **Fetal Echocardiography - Coding (OB-12.1)** and **Fetal Echocardiography - Indications for Fetal Conditions (OB-12.2)**
- Fetal loss of one twin during the first trimester does not appear to increase the risk of FGR or preterm delivery in the surviving twin, however, loss of one or more fetus(es) after the first trimester is associated with increased risk for FGR and PTB.
- Monochorionic gestations or higher order Multiple pregnancy with loss of one or more fetus(es) after the first trimester (≥ 14 weeks) should be imaged according to **Multiple Gestations (OB-11.3)**.
- If other high risk factors, See **High Risk Pregnancy (OB-9)**
- Triplets or higher order Multiple Pregnancy receive same imaging as monochorionic-diamniotic twins.

Background and Supporting Information

- The sensitivity of cfDNA screening may not be as accurate in twins and higher order multiples. First trimester screening in twins has a similar detection rate to singleton gestation.
- Birth weight discordance = (larger twin weight minus smaller twin weight) divided larger twin weight × 100.
- Universal CL screening with transvaginal ultrasound (CPT® 76817) is NOT recommended in twin gestations. In addition, Per ACOG - Cerclage placement (prophylactic or rescue) should be avoided in multifetal pregnancies. However, because several studies have shown that a one-time CL measurement ≤20 mm at 18-24 weeks may be an accurate predictor of preterm birth in multiple gestation, and because progesterone therapy might reduce the risk of neonatal morbidity and mortality associated with PTB, then a one-time transvaginal CL assessment can be performed if trans-abdominal CL measures ≤3.6 cm (as with singleton gestation- See **Cervical Length Screening (OB-7.3)**).
- TTTS is diagnosed by the ultrasound findings of polyhydramnios in one twin (the recipient) and oligohydramnios in the other twin (the donor). If AFI is discordant between the twins (low but not <2 cm in one and/or high but not >8 cm in the other); weekly imaging (MCA and/or limited US) can be performed to rule-out developing TTTS.
- There is no evidence that routine assessment with UA Doppler is beneficial in the absence of growth or fluid discordance.¹
- Fetal loss of one twin during the first trimester does not appear to increase the risk of FGR or preterm delivery in the surviving twin, however, loss of one or more fetus(es) after 17 weeks gestation is associated with increased risk for FGR and PTB and should be imaged according to **Multiple Gestations (OB-11)**. Monochorionic twin pregnancies with demise of one twin after 17 weeks have up to an 18% chance of major morbidity or mortality for the remaining fetus.
- Typically all components of the BPP (CPT® 76818 and CPT® 76819), such as breathing, are not present until ≥26 weeks gestation. However, a modified BPP (CPT® 76815) can be performed sooner in certain high risk cases but should not be done prior to 22 weeks.
- CPT® 76815 should never be reported with complete studies CPT® 76801/CPT® 76802, CPT® 76805/CPT® 76810 or CPT® 76811/CPT® 76812 or with CPT® 76816 or BPP (CPT® 76818 and CPT® 76819)
- Though fetal anatomy survey (CPT® 76805/CPT® 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation
- In circumstances where CPT® 76811 cannot be performed See **Ultrasound Code Selection (OB-1.3)**

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v1.0.2025

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Fetal Echocardiography (ECHO) (OB-12)

Guideline	Page
Fetal Echocardiography – Coding (OB-12.1).....	102
Fetal Echocardiography - Indications for Fetal Conditions (OB-12.2).....	103
Fetal Echocardiography - Indications for Maternal Conditions (OB-12.3).....	105
Fetal Echocardiography - Indications for Medication or Drug Exposure (OB-12.4).....	107
References (OB-12).....	108

Fetal Echocardiography – Coding (OB-12.1)

OB.FE.0012.1.A

v1.0.2025

- Supported fetal echocardiography (echo) codes include:
 - Initial Fetal Echo, CPT[®] 76825 and Doppler Echo CPT[®] 76827 are performed only once per fetus/per facility (i.e. Maternal Fetal Medicine versus Pediatric Cardiology request)
 - Follow-up-Fetal echo and/or Follow-up Doppler echo (CPT[®] 76826/CPT[®] 76828)
 - CPT[®] 93325 for Doppler color flow velocity mapping
- An initial fetal echo is usually not performed prior to 16 weeks.
- Doppler echo procedure codes (CPT[®] 76827 or CPT[®] 76828) include the evaluation of veins, arteries, and valves. Guidelines do not support the billing of additional codes (CPT[®] 76820 and/or CPT[®] 76821)

Background and Supporting Information

- The minimal use of color Doppler (CPT[®] 93325) alone, when performed for anatomical structure identification during a standard ultrasound procedure, is not separately reimbursable

Fetal Echocardiography - Indications for Fetal Conditions (OB-12.2)

OB.FE.0012.2.A

v1.0.2025

Initial Fetal echocardiography (CPT® 76825) and/or Doppler echocardiography (CPT® 76827) with or without Doppler color flow velocity mapping (CPT® 93325) can be performed if ≥16 weeks, for the indications listed below (See **Fetal Echocardiography – Coding (OB-12.1)**):

Fetal Echocardiography - Indications for Fetal Conditions

- Known or suspected abnormal fetal cardiac evaluation on fetal anatomic scan.
 - Known or suspected abnormality must be documented as hard copy or acknowledged verbally by provider of known or suspected fetal cardiac evaluation
 - Suboptimal cardiac evaluation alone is not an indication for fetal echogram. If the 4-chamber view is adequate and there is no other suspicion of a cardiac abnormality, a fetal echocardiogram is not considered medically necessary. A follow up ultrasound (CPT® 76815 or CPT® 76816) is indicated for suboptimal visualization. If the follow-up ultrasound fails to show a 4-chamber view or there is suspicion of a cardiac abnormality, fetal echocardiogram is indicated.
- Fetal cardiac arrhythmia; persistent fetal tachycardia or bradycardia
- Major fetal extra-cardiac anomaly
- Congenital heart disease (CHD) in a 1st degree relative of the fetus (i.e. CHD in the mother, father, or sibling of the fetus) or a half-sibling of the fetus
- Known fetal chromosomal abnormalities (fetal aneuploidy) or ultrasound findings of a suspected chromosomal abnormality (excluding soft markers as only ultrasound findings)
 - Early onset FGR (<32 weeks) may be a sign of fetal aneuploidy^{11,12}
- Single umbilical artery
- Chorioangioma or Umbilical cord varix (if suspicion of fetal hydrops)
- Fetal intra-abdominal venous anomaly (persistent right umbilical vein)
- Fetal effusion (pericardial, pleural effusion, ascites, etc.)
- Fetal hydrops, See **Alloimmunization/Rh Isoimmunization/Other Causes of Fetal Anemia/Parvo/Hydrops (OB-16)**
- Monochorionic twins/TTTS

Fetal Echocardiography - Indications for Fetal Conditions

- Abnormal Fetal Nuchal Translucency scan (NT ≥ 3.0 mm or above the 95th percentile for the CRL) during current pregnancy.
- Abnormal ductus venosus waveform⁵
- Fetal echocardiography may be indicated with severe or unexplained polyhydramnios, or if there are also other suspicious findings on an anatomy scan
- Fetal Echo is **NOT** indicated for an isolated soft marker found on routine imaging including:
 - Choroid plexus cyst, or
 - Echogenic intra-cardiac foci, or
 - Thickened nuchal fold (≥ 6 mm at 15 to 20 weeks), or
 - Absent or hypoplastic nasal bone, or
 - Echogenic bowel, or
 - Shortened long bones, or
 - Pyelectasis

Fetal Echocardiography - Indications for Maternal Conditions (OB-12.3)

OB.FE.0012.3.A

v1.0.2025

Initial Fetal echocardiography (CPT[®] 76825) and/or Doppler echocardiography (CPT[®] 76827) with or without Doppler color flow velocity mapping (CPT[®] 93325) can be performed if ≥ 16 weeks, for the indications listed below (See **Fetal Echocardiography – Coding (OB-12.1)**):

Maternal Conditions:

- Maternal pre-gestational DM or early diagnosed GDM (1st or early 2nd trimester)
- Maternal gestational diabetes mellitus, if HbA1C $>6\%$ [in the third trimester (≥ 32 weeks)]
- Maternal connective tissue disease (SLE, RA, Sjogrens) with Anti-Ro/SSA or anti-La/SSB antibodies present
 - Weekly follow-up Fetal echocardiography (CPT[®] 76826) and/or Doppler fetal echocardiography (CPT[®] 76828) or CPT[®] 76815 from the 18th through the 26th week of pregnancy and then every other week until 30 weeks
- Phenylketonuria
- Infections associated with cardiac anomalies (such as parvovirus, Rubella, Cocksackie virus)
- Genetic conditions associated with CHD in a first degree relative of the fetus (e.g. Marfan syndrome, 22q11.2 deletion syndrome (DiGeorge Syndrome) or Noonan syndrome)
- Prior child with CHD born to mother and/or father of the fetus⁵
- Pregnancy conceived by assisted reproductive technology:¹
 - In Vitro Fertilization (IVF)
 - Intracytoplasmic sperm injection (ICSI)¹

Background and Supporting Information

- If diabetes is diagnosed prior to pregnancy or in the first or early second trimester (typically before 20 weeks gestation) with standard diagnostic criteria of: HbA1C $\geq 6.5\%$, fasting plasma glucose ≥ 126 mg/dL, or 2-hour glucose ≥ 200 mg/dL on a 75-g oral glucose tolerance test, then image as above
- For those with GDM on medication, if HbA1c levels are $>6\%$, fetal echocardiogram in the third trimester to assess for ventricular hypertrophy can be performed.

- The Society of Maternal Fetal Medicine states "we recommend that serial fetal echocardiograms for assessment of the PR interval not be routinely performed in patients with anti-SSA or anti-SSB antibodies outside of a clinical trial setting".
- Follow up for known fetal cardiac anomalies is highly individualized and should be based on the individual cardiac lesion. Interval for follow-up echocardiography should be determined by Maternal Fetal Medicine or Pediatric Cardiologist.

Fetal Echocardiography - Indications for Medication or Drug Exposure (OB-12.4)

OB.FE.0012.4.A

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Initial Fetal echocardiography (CPT[®] 76825) and/or Doppler echocardiography (CPT[®] 76827) with or without Doppler color flow velocity mapping (CPT[®] 93325) can be performed if ≥16 weeks, for the indications listed below (See **Fetal Echocardiography – Coding (OB-12.1)**):

- Ace inhibitors
- Alcohol (excessive quantities)
- Anti-seizure medication, e.g. carbamazepine, hydantoin, valproate
- Folate antagonists (methotrexate)
- Lithium
- NSAIDS (Ibuprofen, Indomethacin) 2nd and 3rd trimester
- Paroxetine (Paxil)
- Retinoids (e.g Isotretinoin, Retinoic acid, Vitamin A -over 10,000 IU per day, etc.)
- Thalidomide
- Venlafaxine (Effexor)
- This may not be an all-inclusive list, however, exposure to other potential teratogens associated with cardiac anomalies in the fetus are typically adequately assessed with a detailed fetal anatomy ultrasound. (CPT[®] 76811) (See **Potentially Teratogenic Medications/Substances (OB 10.1)**)

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v1.0.2025

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Fetal MRI (OB-13)

Guideline	Page
Indications for Fetal MRI (OB-13.1).....	110
References (OB-13).....	113

Indications for Fetal MRI (OB-13.1)

OB.MR.0013.1.A

v1.0.2025

ACOG recommendations for imaging during pregnancy and lactation:

- Ultrasonography and magnetic resonance imaging (MRI) are not associated with risk and are the imaging techniques of choice for the pregnant patient, but they should be used prudently and only when use is expected to answer a relevant clinical question or otherwise provide medical benefit to the patient.
- The use of gadolinium contrast with MRI should be limited; it should be used as a contrast agent in a pregnant female only if it significantly improves diagnostic performance and is expected to improve fetal or maternal outcome.

CPT® Code Guidance

- | |
|--|
| <ul style="list-style-type: none">• Fetal MRI (CPT® 74712); for each additional fetus (CPT® 74713) |
| <ul style="list-style-type: none">• Do not report CPT® 74712 and CPT® 74713 in conjunction with CPT® 72195, CPT® 72196, CPT® 72197 |
| <ul style="list-style-type: none">• If only placenta or maternal pelvis is imaged without fetal imaging, use MRI Pelvis (CPT® 72195) |
- Fetal MRI (CPT® 74712) [plus CPT® 74713 for each additional fetus] optimally performed after 18 to 22 weeks gestation, for assessment of known or suspected fetal abnormalities for counseling, surgical, or delivery planning.
 - There are cases when surgical planning may necessitate imaging earlier than 18 weeks. For those cases where surgery is to be performed prior to 18 weeks and they otherwise meet indications for imaging per this criteria, Fetal MRI may be approved.
 - 3D-4D (CPT® 76376 or CPT® 76377) rendering can be added for surgical planning with diagnosis of complex CHD in the fetus or for surgical planning of other complex fetal malformations⁶.
 - Repeat fetal MRI (CPT® 74712) [plus CPT® 74713 for each additional fetus] later in pregnancy for:
 - Delivery or perinatal surgical planning
 - Fetal MRI indications include but may not be limited to the following:
 - Brain
 - Congenital anomalies
 - Ventriculomegaly
 - Agenesis of the corpus callosum

- Abnormalities of the cavum septum pellucidum
- Holoprosencephaly
- Posterior fossa anomalies
- Malformations of cerebral cortical development
- Microcephaly
- Solid or cystic masses
- Cephalocele
- Screening fetuses with a family risk for brain anomalies
 - Tuberous sclerosis
 - Corpus callosal dysgenesis
 - Malformations of cerebral cortical development
- Vascular abnormalities
 - Vascular malformations
 - Hydranencephaly
 - Intra-uterine cerebrovascular accident (CVA)
- Spine
 - Congenital anomalies
 - Neural tube defects
 - Sacrococcygeal teratomas
 - Caudal regression/sacral agenesis
 - Syringomyelia
 - Vertebral anomalies
- Skull, face, and neck
 - Masses of the face and neck
 - Vascular or lymphatic malformations
 - Hemangiomas
 - Goiter
 - Teratomas
 - Facial clefts
 - Airway obstruction
 - Conditions that may impact parental counseling, prenatal management, delivery planning, and postnatal therapy
- Thorax
 - Masses
 - Congenital pulmonary airway malformations (congenital cystic adenomatoid malformation; sequestration, and congenital lobar emphysema);
 - Congenital diaphragmatic hernia
 - Effusion
 - Mediastinal masses

- Assessment for esophageal atresia
- Volumetric assessment of lung
 - Cases at risk for pulmonary hypoplasia secondary to oligohydramnios, chest mass, or skeletal dysplasias
- Abdomen, retroperitoneal and pelvis
 - Bowel anomalies such as anorectal malformations, or complex bowel obstructions such as with megacystis microcolon hypoperistalsis syndrome
 - Abdominal wall defect
 - Mass
 - Abdominal–pelvic cyst
 - Tumors (e.g. hemangiomas, neuroblastomas, sacrococcygeal teratomas, and suprarenal or renal masses)
 - Complex genitourinary anomalies (e.g. cloaca, prune belly syndrome)
- Congenital Heart Disease (CHD)
- Skeletal dysplasia
- Multiple malformations
- Complications of monochorionic twins/TTTS (eg. Laser treatment of twins, demise of one twin, conjoined twins)
- Any suspected fetal anomaly associated with severe oligohydramnios or anhydramnios

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v1.0.2025

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Abnormal Fetal Position/ Presentation and Pelvimetry (OB-14)

Guideline	Page
Abnormal Fetal Position or Presentation (OB-14.1).....	115
Pelvimetry (OB-14.2).....	116
References (OB-14).....	117

Abnormal Fetal Position or Presentation (OB-14.1)

OB.FP.0014.1.A

v1.0.2025

- To confirm suspected abnormal fetal position or presentation (transverse or breech presentation) at ≥ 36 weeks gestation, report one of the following:
 - CPT[®] 76805 (plus CPT[®] 76810 for each additional fetus) when complete anatomy scan has not yet been performed in the pregnancy **or**
 - CPT[®] 76815 for limited ultrasound to check fetal position **or**
 - CPT[®] 76816 if version is being considered and/or for delivery planning

Background and Supporting Information

- Fetal presentation should be assessed by abdominal palpation (Leopold's) at 36 weeks or later, when presentation is likely to influence the plans for the birth. Routine assessment of presentation by abdominal palpation before 36 weeks is not always accurate. Suspected fetal malpresentation should be confirmed by an ultrasound assessment. An ultrasound can be performed at ≥ 36 weeks gestation to determine fetal position to allow for external cephalic version. Ultrasound to determine fetal position is not necessary prior to 36 weeks gestation unless delivery is imminent.
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).

Pelvimetry (OB-14.2)

OB.FP.0014.2.A**v1.0.2025**

- Pelvimetry (CT or MRI Pelvimetry CPT[®] 72192 or CPT[®] 72195) lacks sufficient evidence to be clinically useful. Current recommendations are that further randomized control studies be performed before it is adapted into routine clinical practice.^{3,4}

References (OB-14)

v1.0.2025

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Adnexal Mass/Uterine Fibroids and Uterine Anomalies (OB-15)

Guideline	Page
Adnexal Mass (OB-15.1).....	119
Uterine Fibroids in Pregnancy (OB-15.2).....	121
Uterine Anomalies in Pregnancy (OB-15.3).....	123
References (OB-15).....	124

Adnexal Mass (OB-15.1)

OB.AM.0015.1.A

v1.0.2025

Adnexal Mass

- For a known or suspected adnexal/pelvic mass, perform:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76805 [plus CPT[®] 76810 if more than one fetus] if a complete fetal anatomic scan has not yet been performed and ≥14 weeks, **or**
 - CPT[®] 76816 if a complete anatomy scan was done previously and/or CPT[®] 76817 if poor visualization of the adnexal mass.
- Following the initial ultrasound, follow up can be done once in each trimester
 - CPT[®] 76805 [plus CPT[®] 76810 for each additional fetus] if a complete fetal anatomic scan has not yet been performed, **or**
 - CPT[®] 76815 or CPT[®] 76816 if a complete ultrasound was previously performed.
 - CPT[®] 76817 if poor visualization of the adnexal mass
- MRI Pelvis (CPT[®] 72195) without contrast can be done for indeterminate findings on ultrasound; for surgical planning and/or for suspected malignancy.
- See **Adnexal Mass/Ovarian Cysts (PV-5)** in the Pelvis Imaging Guidelines

Background and Supporting Information

- The majority of adnexal masses in pregnancy are benign, the most common diagnoses are mature teratomas and corpus luteum or paraovarian cysts. Malignancy is reported in only 1.2-6.8% of pregnant patients with persistent mass.
- Levels of CA-125 are elevated in pregnancy, a low-level elevation in pregnancy is not typically associated with malignancy.
- ACOG recommendations for imaging during pregnancy and lactation:
 - Ultrasonography and magnetic resonance imaging (MRI) are not associated with risk and are the imaging techniques of choice for the pregnant patient, but they should be used prudently and only when use is expected to answer a relevant clinical question or otherwise provide medical benefit to the patient.

- The use of gadolinium contrast with MRI should be limited; it should be used as a contrast agent in a pregnant female only if it significantly improves diagnostic performance and is expected to improve fetal or maternal outcome.

Uterine Fibroids in Pregnancy (OB-15.2)

OB.AM.0015.2.A

v1.0.2025

- If more than one fibroid, total size of all fibroids should be used, i.e. one fibroid at 2 cm and one 3 cm is total of 5 cm and imaging would be indicated as below:
 - Moderate (>5 cm) and large (>10 cm) fibroid(s):
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - Fetal anatomic scan (CPT[®] 76805 or CPT[®] 76811 if other high risk indication. See **High Risk Pregnancy (OB-9)** if ≥16 weeks.
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
 - If the fibroid is in the lower uterine segment or the cervix (cervical fibroid), can perform ultrasound (CPT[®] 76815) and/or transvaginal ultrasound (CPT[®] 76817) every 2 weeks between 16 to 24 weeks, and
 - Follow up ultrasound (CPT[®] 76816) every 3 to 6 weeks, starting at 22 weeks.
- Submucosal fibroid(s) of any size:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - Fetal anatomic scan [CPT[®] 76805 or CPT[®] 76811 if other high risk indication. See **High Risk Pregnancy (OB-9)** if ≥16 weeks
 - If placentation is over a submucosal fibroid:
 - Follow up ultrasound (CPT[®] 76816) every 3 to 6 weeks, starting at 22 weeks

Background and Supporting Information

- Though pregnancy seems to have little or no effect on the overall size of fibroids, Fibroids affect pregnancy and delivery in several ways, with abdominal pain, miscarriage, fetal malpresentation, and difficult delivery being the most frequent complications. These complications relate to preterm labor, placental abruption, fetal growth restriction, and fetal compression syndromes. The risk of preterm labor appears to correlate with the size of the fibroid (over 600 cm³) and/or the presence of multiple fibroids. Placental abruption has been reported to occur

frequently in pregnancies complicated by fibroids, especially with placentation over a fibroid. Fibroid volumes $>200 \text{ cm}^3$ are more commonly associated with fetal growth restriction. Fetal compression syndrome is a direct result of large fibroids and is not commonly found with small fibroids. Finally, malposition or obstructed labor may be associated with fibroids of the lower uterine segment.

Uterine Anomalies in Pregnancy

(OB-15.3)

OB.AM.0015.3.A

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- For uterine septum, uterine didelphys, unicornuate uterus, bicornuate uterus:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76805 or CPT[®] 76811 and/or CPT[®] 76817 at ≥16 weeks
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
 - CPT[®] 76817 and/or CPT[®] 76815 every 2 weeks at 16 to 24 weeks (See **Cervical Insufficiency (OB-18.1)**)
 - CPT[®] 76816 every 3 to 6 weeks starting at ≥22 weeks
 - Starting at 32 weeks, weekly BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815)

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a detailed fetal anatomic ultrasound examination (CPT[®] 76811) due to geographic or other constraints, a standard (after first trimester) fetal ultrasound examination (CPT[®] 76805) can be authorized instead.
- CPT[®] 76811 and CPT[®] 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication and/or change in condition.

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v1.0.2025

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Alloimmunization/Rh Isoimmunization/Other Causes of Fetal Anemia/ Parvo/Hydrops (OB-16)

Guideline	Page
Alloimmunization/Rh Isoimmunization (OB-16.1).....	126
Exposure to Parvovirus B-19 (OB-16.2).....	128
Twin Anemia Polycythemia Sequence (OB-16.3).....	129
Other Fetal Hydrops/Nonimmune Hydrops (OB-16.4).....	130
Other Causes of Fetal Anemia (OB-16.5).....	131
References (OB-16).....	132

Alloimmunization/Rh Isoimmunization (OB-16.1)

OB.AR.0016.1.A

v1.0.2025

Imaging for Alloimmunization/Rh Isoimmunization for any of the following indications:

- When any one of the following maternal antibody titers are $\geq 1:8$ (critical titer):
 - Rhesus antibodies (Cc/Dd/Ee)
 - Anti-Duffy (anti-fya) antibody
 - Anti-Kidd antibody
- If maternal antibody titers are $\geq 1:8$ for other atypical antigens that may be associated with hemolytic disease of the fetus and newborn, e.g. p antigen, MNS series, MSSsMT, Diego, Public antigens, Private antigens.
- Anti-Kell antibody (any antibody titer warrants additional evaluation)
- If evidence of fetal hydrops on previous imaging
- Prior pregnancy associated with HDFN (hemolytic disease of the fetus and newborn)

The following imaging is indicated:

- Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- Detailed Fetal Anatomic Scan (CPT[®] 76811) ≥ 16 weeks
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- CPT[®] 76816 every 2 to 4 weeks to assess fetal growth starting after fetal anatomic scan (CPT[®] 76811)

The following imaging is indicated:

- Fetal middle cerebral artery (MCA) Doppler (CPT[®] 76821) and CPT[®] 76815 every 1 to 2 weeks starting at 16 weeks
- BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815) weekly, starting at 32 weeks or sooner if suspected worsening fetal condition (≥ 26 weeks), e.g. abnormally trending MCA dopplers, suspected hydrops, or polyhydramnios.
- More frequent imaging (MCA Doppler and/or BPP) can be performed if suspected worsening fetal condition (up to 2x weekly), e.g. abnormally trending MCA dopplers, suspected hydrops, or polyhydramnios.

Background and Supporting Information

- Fetal anemia and hydrops may be a result of immune conditions, such as red-cell or Kell alloimmunization, non-immune hydrops caused by parvovirus B19 infection or any other known acquired or congenital causes of fetal anemia.
- Rhesus isoimmunization/alloimmunization is the process through which fetal Rh+ red blood cells enter the circulation of an Rh negative mother causing her to produce antibodies which can cross the placenta and destroy the red blood cells of the current Rh+ fetus and/or in subsequent Rh+ pregnancies.
- Atypical antigens not listed above, may be associated with hemolytic disease of the fetus and newborn and may require fetal assessment as in Alloimmunization/Rh Isoimmunization (OB-16.1) if maternal antibody titers are $\geq 1:8$. Atypical antigens include: MNSs series (M, N, S, s, U, Mi), MSSs-Mta, Diego (Di^a, Di^b), P- PPTj, Public antigen (Yt, En, Co2). Private antigens (Biles, Good, Heibel, Radin, Wright^a, and Z^d). PP1Pk, Far, Good, Lan, LW.
- Peak systolic velocity (PSV) of the fetal middle cerebral artery can be used as a substitute for amniocentesis to evaluate a fetus at risk for anemia due to Rhesus isoimmunization/alloimmunization. Measurements can be initiated as early as 16 weeks of gestation if there is a past history of early severe fetal anemia or very high titers. Because MCA-PSV increases across gestation, results are adjusted for gestational age.
- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a detailed fetal anatomic ultrasound examination (CPT[®] 76811) due to geographic or other constraints, a standard (after first trimester) fetal ultrasound examination (CPT[®] 76805) can be authorized instead.
- CPT[®] 76811 and CPT[®] 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office **and** there is a new medical indication and/or change in condition.

Exposure to Parvovirus B-19 (OB-16.2)

OB.AR.0016.2.A

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- Parvovirus B-19 Exposure (Fifth Disease):
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76811 if not yet performed and ≥16 weeks
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
 - CPT[®] 76816 if >2 weeks since performance of the fetal anatomic scan (CPT[®] 76811)
- Confirmed Parvovirus B-19 Infection (+IgM):
 - Ultrasound (CPT[®] 76816) every 2 to 4 weeks to assess fetal growth starting after performance of the fetal anatomic scan (CPT[®] 76811) and
 - Starting at time of confirmed infection weekly limited ultrasound (CPT[®] 76815) if >22 weeks or weekly BPP (CPT[®] 76818 or CPT[®] 76819) if ≥26 weeks gestation and
 - Fetal middle cerebral artery (MCA) Doppler (CPT[®] 76821) every 1 to 2 weeks, starting at time of confirmed infection (if ≥16 weeks)
 - Continue above imaging for 8 to 12 weeks after initial date of exposure

Twin Anemia Polycythemia Sequence (OB-16.3)

OB.AR.0016.3.A

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- See **Known Monochorionic-Diamniotic or Monochorionic-Monoamniotic Multiple Gestations (OB-11.3)**

Other Fetal Hydrops/Nonimmune Hydrops (OB-16.4)

OB.AR.0016.4.A

v1.0.2025

- Suspected or known hydrops from any cause should be imaged according to **Alloimmunization/Rh Isoimmunization (OB-16.1)**

Other Causes of Fetal Anemia (OB-16.5)

OB.AR.0016.5.A

v1.0.2025

- MCA Doppler (CPT® 76821) assessment can be performed if there is suspected fetal anemia (e.g. fetus with heart failure, hydrops, alloimmunization), in a fetus at high risk for fetal anemia due to other pregnancy complications, e.g. chorioangioma, umbilical vein varix, or finding of sustained fetal tachyarrhythmia or bradyarrhythmia - See **Indications for Fetal Conditions (OB-12.2)**, **Alloimmunization/Rh Isoimmunization (OB-16.1)**, and **Placental and Cord Abnormalities (OB-21)**

Background and Supporting Information

- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a fetal anatomic ultrasound examination (CPT® 76811) due to geographic or other constraints, a standard (after first trimester) fetal anatomic ultrasound examination (CPT® 76805) can be authorized instead.
- CPT® 76811 and CPT® 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication and/or change in condition.
- Typically all components of the BPP (CPT® 76818 and CPT® 76819), such as breathing, are not present until ≥ 26 weeks gestation. However, a modified BPP (CPT® 76815) can be utilized sooner in certain high risk cases but should not be done prior to 22 weeks. .

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v1.0.2025

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Amniotic Fluid Abnormalities/ Oligohydramnios/ Polyhydramnios (OB-17)

Guideline	Page
Amniotic Fluid Abnormalities (OB-17.1).....	134
References (OB-17).....	136

Amniotic Fluid Abnormalities (OB-17.1)

OB.AF.0017.1.A

v1.0.2025

For suspected polyhydramnios or oligohydramnios:

- For example, unequal size and dates or suspected preterm/prelabor rupture of membranes. See **Unequal Fundal Size and Dates (OB-27)** and/or **Preterm/Prelabor Rupture of Membranes (OB-23)**
 - CPT® 76815 for quick look for AFI check

For **confirmed** diagnosis of **polyhydramnios**: AFI ≥ 24 cm or maximum vertical pocket (MVP) ≥ 8 cm.

- CPT® 76811 (Detailed Fetal Anatomy) at diagnosis, if not previously performed
- CPT® 76816 starting at ≥ 22 weeks
 - Every 3 - 4 weeks for mild polyhydramnios (AFI 24 - 29.9 cm or MVP 8 - 9.9 cm)
 - Every 2 weeks for severe polyhydramnios (AFI ≥ 30 cm or MVP ≥ 10 cm)
- CPT® 76815 weekly for antepartum fetal surveillance starting at ≥ 22 weeks
- BPP (CPT® 76818 or CPT® 76819) or modified BPP (CPT® 76815) for AFI with NST starting at 26 weeks
 - Weekly for mild polyhydramnios
 - Twice weekly for severe polyhydramnios

For **confirmed** diagnosis of **oligohydramnios**: AFI ≤ 5 cm or maximum vertical pocket ≤ 2 cm

- CPT® 76811 if not already performed; or
- CPT® 76816 every 2 to 4 weeks starting at ≥ 22 weeks
- CPT® 76815 weekly for antepartum fetal surveillance from 22 - 26 weeks
- CPT® 76818 or CPT® 76819 or a modified BPP (CPT® 76815), weekly, starting at 26 weeks
- CPT® 76820 weekly starting at time of diagnosis if ≥ 22 weeks

Background and Supporting Information

- Polyhydramnios can be an early presenting finding of fetal hydrops associated with fetal anemia. Middle cerebral artery Doppler is commonly used to diagnose whether this fetal anemia is present or not. See **Alloimmunization/Rh Isoimmunization (OB-16.1)**

- Polyhydramnios may also present as a finding of cardiac dysfunction, fetal arrhythmias or cardiac malformation. Fetal echocardiography may be indicated if there are abnormal findings on an anatomy scan. See **Fetal Echocardiography (ECHO) (OB-12)**
- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a fetal anatomic ultrasound examination (CPT[®] 76811) due to geographic or other constraints, a standard (after first trimester) fetal anatomic ultrasound examination (CPT[®] 76805) can be authorized instead.
- CPT[®] 76811 and CPT[®] 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication and/or change in condition.
- Typically all components of the BPP (CPT[®] 76818 and CPT[®] 76819), such as breathing, are not present until ≥ 26 weeks gestation. However, a modified BPP (CPT[®] 76815) can be utilized sooner in certain high risk cases but should not be done prior to viability (22 weeks).
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

References (OB-17)

v1.0.2025

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Cervical Insufficiency/ Current Preterm Labor (OB-18)

Guideline	Page
Cervical Insufficiency (OB-18.1).....	138
Cerclage in Place in Current Pregnancy (OB-18.2).....	139
Current Preterm Labor (OB-18.3).....	140
References (OB-18).....	141

Cervical Insufficiency (OB-18.1)

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v1.0.2025

- For any of the following:
 - History of cervical insufficiency (defined as one or more 2nd trimester loss (14 to 24 weeks gestation) related to painless cervical dilation.)
 - History of cerclage in prior pregnancy
 - History of prior precipitous delivery
 - Surgical trauma to cervix, e.g.
 - Over dilation of cervix during a termination of pregnancy
 - Cervical obstetrical laceration from a previous delivery
 - History of extensive or multiple prior cold-knife conization or Loop Electrosurgical Excision Procedures (LEEP)
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
 - CPT[®] 76805 or CPT[®] 76811 if other high risk factors [plus CPT[®] 76810/CPT[®] 76812 for each additional fetus] **and/or** CPT[®] 76817 at ≥16 weeks if a complete fetal anatomic scan has not yet been performed during this pregnancy.
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
 - CPT[®] 76815 **and/or** CPT[®] 76817 every 1 to 4 weeks from 16 0/7 to 24 0/7 weeks
 - CPT[®] 76816 can be performed every 3 to 6 weeks starting after the fetal anatomic scan at 16 weeks
 - Note: Lower uterine segment or cervical fibroid may also require CL screening with (CPT[®] 76815) and/or transvaginal ultrasound (CPT[®] 76817) every 2 weeks between 16 to 24 weeks (See **Uterine Fibroids in Pregnancy (OB-15.2)**)
-
- If funneling or short cervix ≤25 mm (2.5 cm) is found on a transvaginal ultrasound
 - CPT[®] 76816 can be performed after a complete ultrasound every 3 to 4 weeks **and/or**
 - CPT[®] 76817 and/or CPT[®] 76815 every 1 to 2 weeks until 32 weeks
 - Note: CPT[®] 76815 should not be done on same date of service as CPT[®] 76816
 - Starting at 32 weeks, BPP (CPT[®] 76818 or CPT[®] 76819) or a modified BPP (CPT[®] 76815) can be performed once weekly.

Cerclage in Place in Current Pregnancy (OB-18.2)

OB.CI.0018.2.A

v1.0.2025

- Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, **and/or** CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- CPT[®] 76805 or CPT[®] 76811 if other high risk factors [plus CPT[®] 76810/CPT[®] 76812 for each additional fetus] and/or CPT[®] 76817 once if a complete detailed fetal anatomic scan has not been performed
- CPT[®] 76816 can be performed every 3 to 6 weeks starting after the fetal anatomic scan at 16 weeks.
- If a **prophylactic** (history-indicated) cerclage is placed in the current pregnancy (usually done early in the second trimester due to a prior 2nd trimester loss after painless cervical dilation at ≤24 weeks gestation)
 - CPT[®] 76815 **and/or** CPT[®] 76817 can be performed once between 16 to 24 weeks to determine if cervical shortening is present
 - If cervical shortening or funneling is noted See **Cervical Insufficiency (OB-18.1)**.
 - Further CL surveillance CPT[®] 76815 **and/or** CPT[®] 76817 every 2 weeks is not indicated.
- If a **rescue** cerclage (exam indicated) was placed due to cervical shortening or dilation found by ultrasound or physical exam in the current pregnancy
 - Transvaginal (CPT[®] 76817 and/or CPT[®] 76815) every 2 weeks, starting at ≥16 weeks until 32 weeks.
- If funneling or further shortening of the cervix is found See **Cervical Insufficiency (OB 18.1)**

Current Preterm Labor (OB-18.3)

OB.CI.0018.3.A

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- Preterm labor in current pregnancy (contractions **PLUS** cervical change at <37 weeks gestation), can perform once when symptomatic
 - CPT[®] 76805 or CPT[®] 76811 if other high risk factors [plus CPT[®] 76810/CPT[®] 76812 for each additional fetus] if a complete fetal anatomic scan has not yet been performed during this pregnancy; **or**
 - CPT[®] 76815 **or** CPT[®] 76816, if a complete fetal anatomic scan was performed previously, (CPT[®] 76816 if more than 2 weeks since last growth), **and/or** CPT[®] 76817
- Biophysical profile (BPP) (CPT[®] 76818 **or** CPT[®] 76819) **or** modified BPP (CPT[®] 76815), once when symptomatic if ≥26 weeks.
- For history of pre-term labor, See **History of Spontaneous Pre-Term Delivery/History of PPROM (OB-9.9)**

Background and Supporting Information

- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a fetal anatomic ultrasound examination (CPT[®] 76811) due to geographic or other constraints, a standard (after first trimester) fetal anatomic ultrasound examination (CPT[®] 76805) can be authorized instead.
- CPT[®] 76811 and CPT[®] 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication and/or change in condition.
- Typically all components of the BPP (CPT[®] 76818 and CPT[®] 76819), such as breathing, are not present until ≥26 weeks gestation. However, a modified BPP (CPT[®] 76815) can be utilized sooner in certain high risk cases but should not be done prior to 22 weeks.
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

References (OB-18)

v1.0.2025

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No Fetal Heart Tones/ Decreased Fetal Movement (OB-19)

Guideline	Page
No Fetal Heart Tones (OB-19.1).....	143
Decreased Fetal Movement (OB-19.2).....	144
References (OB-19).....	145

No Fetal Heart Tones (OB-19.1)

OB.FT.0019.1.A
v1.0.2025

- Prior to considering ultrasound, fetal heart tones (FHT) should be assessed with a hand-held doppler device.

First Trimester
• If ≥12 weeks gestation and unable to obtain FHT using a hand-held doppler, or • If <12 weeks gestation, in the setting of absent fetal heart tones, only if accompanied by other maternal signs or symptoms (such as cramping, vaginal bleeding, etc.) or • If <12 weeks and FHT had previously been heard using a fetal hand-held doppler, but now are unable to be heard by this method, regardless of symptoms • Report one of the following: ◦ Complete first trimester anatomy ultrasound CPT[®] 76801 (plus CPT[®] 76802 for each additional fetus) and/or CPT[®] 76817 if a complete ultrasound has not yet been performed; or ◦ CPT[®] 76815 and/or CPT[®] 76817
Second and Third Trimester
Report one of the following: • CPT[®] 76815 for limited ultrasound or • CPT[®] 76805 (plus CPT[®] 76810 for each additional fetus) if ≥14 weeks, when complete fetal anatomic scan CPT[®] 76805 has not yet been performed or • CPT[®] 76816 if ≥14 weeks and an anatomy ultrasound (CPT[®] 76801 or CPT[®] 76805/CPT[®] 76811) was performed previously

Decreased Fetal Movement (OB-19.2)

OB.FT.0019.2.A

v1.0.2025

- One of the following can be performed at the time of the complaint of decreased fetal movement:
 - Limited ultrasound or modified BPP (CPT[®] 76815) or
 - If ≥ 26 weeks BPP (CPT[®] 76818 or CPT[®] 76819) or a modified BPP (CPT[®] 76815) can be performed. See **Biophysical Profile (BPP) (OB-28.8)**
 - CPT[®] 76816 for delivery planning, e.g. if BPP/Modified BPP is abnormal.
 - Subsequent/repeat ultrasound is not usually necessary unless there are new indications or if BPP is abnormal.

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).

References (OB-19)

v1.0.2025

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Fetal Growth Problems (FGR and Macrosomia) (OB-20)

Guideline	Page
Fetal Growth Restriction Current Pregnancy (OB-20.1).....	147
Macrosomia – Large for Dates Current Pregnancy (OB-20.2).....	149
References (OB-20).....	150

Fetal Growth Restriction Current Pregnancy (OB-20.1)

OB.FG.0020.1.A

v1.0.2025

- The ACOG definition of Fetal Growth Restriction (FGR): Estimated or actual weight of the fetus $\leq 10^{\text{th}}$ percentile for gestational age, and/or Abdominal Circumference $\leq 10^{\text{th}}$ percentile.

For Suspected FGR:

- If there is a $\geq 3\text{cm}$ difference in fundal height measurement and gestational age, **or**
- FGR is suspected based on reduced fetal growth velocity, defined as a fall in the EFW or AC of 50% or more (e.g. from 70% to 20% ect.).¹²
- Report **one** of the following:
 - CPT[®] 76805/CPT[®] 76811 if otherwise indicated (plus CPT[®] 76810/CPT[®] 76812 for each additional fetus) if anatomy ultrasound has not yet been performed **or**
 - CPT[®] 76816 if anatomy ultrasound (CPT[®] 76805/CPT[®] 76811) previously performed
- For clinical situations that have a higher probability of FGR such as maternal hypertension, maternal diabetes, previous stillbirth, etc. See **High Risk Pregnancy (OB-9)**, or the specific guidelines for these clinical entities for guidance regarding follow-up ultrasounds to assess fetal growth

For Known FGR:

- Detailed Fetal Anatomic Scan (CPT[®] 76811) at diagnosis if not already performed
- Starting at diagnosis, if ≥ 16 weeks gestation, follow up ultrasound (CPT[®] 76816) can be performed every 2 to 4 weeks if complete anatomy ultrasound previously performed
- Starting at 22 weeks, a modified BPP (CPT[®] 76815) can be performed once or twice weekly, **or**
- Starting at 26 weeks, BPP (CPT[®] 76818 or CPT[®] 76819) **or** a modified BPP (CPT[®] 76815) can be performed once or twice weekly, **and**
- Starting at 22 weeks Umbilical artery (UA) Doppler (CPT[®] 76820) can be performed weekly.
- If FGR is diagnosed in the current ultrasound, BPP (CPT[®] 76818 or CPT[®] 76819) can be performed if ≥ 26 weeks, and/or UA Doppler (CPT[®] 76820) if ≥ 22 weeks.

For Known FGR:

- If Severe FGR (EFW $\leq 3\%$, AC $\leq 3\%$), **OR** Abnormal UA Doppler studies (defined as a PI, RI, or S/D ratio greater than the 95th percentile for gestational age **OR** absent or reversed end-diastolic velocity (AEDV or REDV), **OR** confirmed oligohydramnios:
 - BPP (CPT[®] 76818 or CPT[®] 76819 or CPT[®] 76815) and/or umbilical artery (UA) Doppler (CPT[®] 76820) can be performed twice weekly.⁵
- Per SMFM guidelines: Ductus venosus, middle cerebral artery, or uterine artery Doppler use for routine clinical management of early- or late-onset FGR is not recommended

Background and Supporting Information

- An abnormal umbilical artery Doppler is defined as a PI, RI, or S/D ratio greater than the 95th percentile for gestational age or an absent or reversed end-diastolic velocity (AEDV or REDV). Those with REDV are usually hospitalized for closer surveillance and delivery planning.
- Fetuses with early onset FGR are at significant risk for Intrauterine Fetal Demise (IUFD) and the rate of fetal growth prior to 23 weeks and Umbilical Artery (UA) Doppler findings may aid in counseling patients in this peri-viability period (Termination of Pregnancy (TOP) versus attempt at prolonging pregnancy till viability).
- In circumstances where CPT[®] 76811 cannot be performed See **Ultrasound Code Selection (OB-1.3)**
- Typically all components of the BPP (CPT[®] 76818 and CPT[®] 76819), such as breathing, are not present until ≥ 26 weeks gestation. However, a modified BPP (CPT[®] 76815) can be utilized sooner in certain high risk cases but should not be done prior to 22 weeks.
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

Macrosomia – Large for Dates Current Pregnancy (OB-20.2)

OB.FG.0020.2.A

v1.0.2025

- The ACOG definition of macrosomia: Estimated fetal weight of greater than 4000 grams (DM) or 4500 grams (non-DM); $\geq 90^{\text{th}}$ percentile for gestational age.
- For history of a macrosomia See **Prior Pregnancy with Macrosomia (OB-9.4.1)**

For Suspected Macrosomia

- At ≥ 22 weeks gestation, if there is a ≥ 3 week difference in fundal height and gestational age, or if the estimated fetal weight is $\geq 90^{\text{th}}$ percentile for gestational age¹¹, report **one** of the following:
 - CPT[®] 76805 [plus CPT[®] 76810 for each additional fetus] if a complete fetal anatomic scan has not yet been performed **or**
 - CPT[®] 76816 if a complete ultrasound was done previously
- See **Unequal Fundal Size and Dates (OB-27.1)**

For Known Macrosomia $\geq 90^{\text{th}}$ percentile

Report:

- CPT[®] 76816 at ≥ 35 weeks for delivery planning (if more than 2 weeks since last growth).¹¹
- In a low risk pregnancy, ultrasound is generally not indicated to estimate fetal weight before 30 weeks gestation. As such, repeat imaging is generally not necessary unless needed to plan for delivery or if there are other high risk indications.
- Additional imaging recommendations are usually guided by the cause of the fetal macrosomia (obesity, DM, etc.) See appropriate guideline for indication
- Per SMFM guidelines: Ductus venosus, middle cerebral artery, or uterine artery Doppler use for routine clinical management of early- or late-onset FGR is not recommended

References (OB-20)

v1.0.2025

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Placental and Cord Abnormalities (OB-21)

Guideline	Page
Single Umbilical Artery (Two Vessel Cord) (OB-21.1).....	152
Persistent Right Umbilical Vein (PRUV) (OB-21.2).....	153
Placental/Cord Abnormalities (OB-21.3).....	154
Subchorionic Hematoma/Hemorrhage (Placental Hematoma) (OB-21.4).....	157
Suspected Abruption Placentae (OB-21.5).....	158
Previa (Placenta Previa and Vasa Previa) (OB-21.6).....	159
Placenta Accreta Spectrum (PAS): Accreta, Increta, Percreta (OB-21.7).....	161
References (OB-21).....	163

Single Umbilical Artery (Two Vessel Cord) (OB-21.1)

OB.UA.0021.1.A

v1.0.2025

If a single umbilical artery is found on initial imaging:	
- Detailed anatomic ultrasound at ≥ 16 weeks - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.	CPT [®] 76811
Fetal echocardiogram (if ≥ 16 weeks)	CPT [®] 76825 and/or CPT [®] 76827 and/or CPT [®] 93325
Follow-up ultrasound to evaluate fetal growth at ≥ 28 weeks and then every 3 to 6 weeks if more than one clinical high-risk factors are documented	CPT [®] 76816
Weekly BPP or modified BPP starting at 36 weeks	CPT [®] 76818 or CPT [®] 76819 (BPP) or modified BPP CPT [®] 76815

Background and Supporting Information

- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a fetal anatomic ultrasound examination (CPT[®] 76811) due to geographic or other constraints, a standard (after first trimester) fetal anatomic ultrasound examination (CPT[®] 76805) can be authorized instead.
- CPT[®] 76811 and CPT[®] 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication and/or change in condition.

Persistent Right Umbilical Vein (PRUV) (OB-21.2)

OB.UA.0021.2.A
v1.0.2025

PRUV – is a variant of the usual intra-abdominal umbilical venous connection. It may be associated with other fetal defects.

If a PRUV is found on initial imaging:	
- Detailed anatomic ultrasound at ≥16 weeks - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.	CPT [®] 76811
Fetal echocardiogram (if ≥16 weeks)	CPT [®] 76825 and/or CPT [®] 76827 and/or CPT [®] 93325
Follow-up ultrasound to evaluate fetal growth at ≥22 weeks and then every 3 to 6 weeks if more than one clinical high-risk factors are documented	CPT [®] 76816
Weekly BPP or modified BPP starting at 32 weeks	CPT [®] 76818 or CPT [®] 76819 (BPP) or modified BPP CPT [®] 76815

Placental/Cord Abnormalities (OB-21.3)

OB.UA.0021.3.A

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Placental/Cord Abnormalities (OB-21.3.1)

Circumvallate Placenta

Placental hemangioma

Succenturiate placenta or accessory lobe

Hypo/Hyper-coiled Umbilical Cord

Marginal Cord Insertion

Umbilical cord cyst

Velamentous Cord Insertion

- Fetal anatomic scan can be performed after 16 weeks (CPT[®] 76805/CPT[®] 76811) with or without CPT[®] 93976 (limited duplex scan of arterial and venous)
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- Ultrasound CPT[®] 76817 can be performed to evaluate the placenta and/or cord in relation to the cervix
- Ultrasound (CPT[®] 76816) with or without CPT[®] 93976 (limited duplex scan) every 3-6 weeks starting at 28 weeks until delivery
- Weekly BPP or modified BPP (CPT[®] 76818/CPT[®] 76819 or CPT[®] 76815) starting at 32 weeks

Background and Supporting Information

- **Hypo/Hyper-coiled umbilical cord** - Several studies have reported an increased frequency of adverse pregnancy outcome, including congenital anomalies, growth restriction, fetal heart rate abnormalities, preterm birth, and intrauterine death in pregnancies with both hypocoiled and hypercoiled umbilical cord.

Other Placental/Cord Abnormalities (OB-21.3.2)

Amniotic Sheet/ Amniotic Band (Uterine Synechiae)

- Fetal anatomic scan can be performed after 16 weeks (CPT[®] 76805)

- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- One follow-up Ultrasound (CPT[®] 76816) can be performed in the 3rd trimester to ensure that the band is not restricting fetal growth or movement.
- **Amniotic Band Syndrome** is a completely different entity, associated with an increased risk of fetal anomalies and poor outcome. If Amniotic band syndrome is suspected or diagnosed, image as in **High Risk Pregnancy (OB-9)**.

Chorioangioma

Umbilical cord varix

- Detailed fetal anatomic scan can be performed after 16 weeks (CPT[®] 76811) with or without CPT[®] 93976 (limited duplex scan).
 - Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- Ultrasound (CPT[®] 76816) with or without CPT[®] 93976 (limited duplex scan) every 3-6 weeks starting at the time of diagnosis until delivery
- Weekly BPP (CPT[®] 76818 or CPT[®] 76819) or a modified BPP (CPT[®] 76815) starting at 32 weeks
- Both chorioangioma and UVV can be associated with fetal anemia and/or low output heart failure. As such, MCA Dopplers (CPT[®] 76821) may be indicated on a case-by-case basis, e.g. If turbulence develops within the UVV
- If suspected or known hydrops, Fetal ECHO (CPT[®] 76825, CPT[®] 76827, CPT[®] 93325) may be indicated. See **Fetal Echocardiography - Indications for Fetal Conditions (OB-12.2)**. If fetal hydrops develops then image as per **Alloimmunization/Rh Isoimmunization (OB-16.1)**

Background and Supporting Information

- Amniotic Bands visualized on ultrasound are often due to uterine synechiae (intrauterine adhesions), residual gestation sac of a demised twin, fibrin strands s/p bleeding, chorioamniotic separation or may be noted with a circumvallate placenta. In general, they are benign entities and are not associated with adverse pregnancy outcome.
- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform the more desirable fetal and maternal ultrasound with detailed fetal anatomic examination (CPT[®] 76811) due to geographic or other constraints, a standard (after first trimester) fetal and maternal ultrasound (CPT[®] 76805) can be authorized instead.

- CPT[®] 76811 and CPT[®] 76812 should only be used once per pregnancy unless the mother changes to a new medical caregiver at a new office and there is a new medical indication and/or change in condition

Subchorionic Hematoma/Hemorrhage (Placental Hematoma) (OB-21.4)

OB.UA.0021.4.A

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Subchorionic Hematoma/Hemorrhage (Placental Hematoma)

- Ultrasound can be performed for follow-up of a known subchorionic hematoma or placental hematoma
 - CPT[®] 76815 and/or CPT[®] 76817 if the last ultrasound was performed ≥ 7 days ago or
 - CPT[®] 76816 and/or CPT[®] 76817 if a complete ultrasound scan was performed ≥ 2 weeks ago
- Imaging can be repeated earlier than seven days if there are new or worsening symptoms such as an increasing amount of vaginal bleeding or increasing cramping or pain.
- No further imaging is needed if the follow-up ultrasound shows that the hemorrhage has resolved.
- If pregnancy is in second or third trimester follow **Suspected Abruptio Placentae (OB-21.5)**

Suspected Abruption Placentae (OB-21.5)

OB.UA.0021.5.A
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Suspected Abruption Placentae
Second and Third Trimesters
- For suspected abruption placentae: - CPT[®] 76805 [plus CPT[®] 76810 for each additional fetus] or CPT[®] 76811 if confirmed,¹⁶ and/or CPT[®] 76817 if a complete fetal anatomic scan has not yet been performed during this pregnancy, with or without CPT[®] 93976 (limited duplex scan) - CPT[®] 76815 for limited ultrasound and/or CPT[®] 76817, or - CPT[®] 76816 if anatomy ultrasound was done previously, and/or CPT[®] 76817 - Vaginal bleeding with +KB (Kleihauer-Betke) – feto-maternal hemorrhage – at risk for fetal anemia and hydrops CPT[®] 76821 can be performed.
- Ultrasound is appropriate to follow-up a known abruption: - CPT[®] 76815 or CPT[®] 76816 if a complete ultrasound was done previously and/or CPT[®] 76817. - The number and frequency of follow-up ultrasounds will depend on the degree of abruption and the presence or absence of ongoing signs and symptoms.

Previa (Placenta Previa and Vasa Previa) (OB-21.6)

OB.UA.0021.6.A

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Placenta Previa (OB-21.6.1)

Placenta Previa
Second and Third Trimesters
- For suspected placenta previa (placental edge covers the internal cervical os) one of the following ultrasound can be performed: - CPT[®] 76805 [plus CPT[®] 76810 for each additional fetus] and/or CPT[®] 76817 if a complete fetal anatomic scan has not yet been performed during this pregnancy with or without CPT[®] 93976 (limited duplex scan) or - CPT[®] 76815 for limited ultrasound and/or CPT[®] 76817 with or without CPT[®] 93976 (limited duplex scan) or - CPT[®] 76816 if a complete ultrasound was done previously and/or CPT[®] 76817 for a transvaginal ultrasound with or without CPT[®] 93976 (limited duplex scan) - For known placenta previa (placental edge covers the internal cervical os) or low lying placenta (placental edge <2 cm from internal os): - One routine follow-up ultrasound can be performed in the 3rd trimester (CPT[®] 76815 or CPT[®] 76816 and/or CPT[®] 76817) - If placenta previa or low lying placenta is still present, one follow-up ultrasound (CPT[®] 76815 or CPT[®] 76816 and/or CPT[®] 76817) can be performed in 3-4 weeks - If persistent placenta previa (placental edge covers the internal cervical os), BPP (CPT[®] 76818/CPT[®] 76819 or modified BPP (CPT[®] 76815) weekly, starting at 32 weeks - Follow-up ultrasound can be performed at any time if bleeding occurs BPP (CPT[®] 76818 or CPT[®] 76819) or CPT[®] 76815 or CPT[®] 76816 if a complete ultrasound was done previously and/or CPT[®] 76817)

Background and Supporting Information

- For pregnancies beyond 16 weeks, if the placental edge is ≥ 2 cm away from the internal os, the placental location should be reported as normal.

- If the placental edge is <2 cm from the internal os but not covering the internal os, it should be labeled as low lying.
- If the placental edge covers the internal cervical os, the placenta should be labeled as a placenta previa.
- "There is no evidence to guide the optimal time of subsequent imaging in pregnancies thought to have placenta previa. In stable patients it is reasonable to perform a follow-up ultrasonogram at approximately 32 weeks of gestation and an additional study at 36 weeks of gestation (if the previa persists) to determine the optimal route and timing of delivery. There is no clear benefit from more frequent ultrasonograms (eg, every 4 weeks) in stable cases."¹⁷

Vasa Previa (OB-21.6.2)

- Vasa previa occurs when fetal blood vessels that are unprotected by the umbilical cord or placenta run through the amniotic membranes and cross over the internal cervical os.

If a Vasa Previa is found on initial imaging:	
Detailed anatomic ultrasound at ≥16 weeks • Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.	CPT [®] 76811 and/or ¹⁹ CPT [®] 76817
Follow-up growth ultrasound every 2 to 4 weeks starting at ≥22 weeks	CPT [®] 76816 and/or ¹⁹ CPT [®] 76817
Once vasa previa is confirmed cervical length screening every 2 to 4 weeks starting at 28 weeks	CPT [®] 76817 and CPT [®] 76816 or CPT [®] 76815
BPP or modified BPP weekly starting at 32 weeks (can be performed earlier and/or more frequently if worsening fetal condition suspected)	CPT [®] 76818 or CPT [®] 76819 (BPP) or CPT [®] 76815

Placenta Accreta Spectrum (PAS): Accreta, Increta, Percreta (OB-21.7)

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v1.0.2025

- PAS includes placenta accreta, placenta increta or placenta percreta.

Suspected PAS (OB-21.7.1)

- For **suspected** PAS:
 - CPT[®] 76811 or CPT[®] 76805 and/or CPT[®] 76817 if a complete fetal anatomic scan has not yet been performed. Can be performed with or without CPT[®] 93976 (limited duplex scan) **or**
 - CPT[®] 76816 (if a complete ultrasound was done previously) or CPT[®] 76815 and/or CPT[®] 76817 with or without CPT[®] 93976 (limited duplex scan)
 - MRI Pelvis without contrast (CPT[®] 72195) if inconclusive or equivocal ultrasound

Known PAS (OB-21.7.2)

- For **known** PAS:
 - Follow up growth ultrasounds can be performed every 2 to 4 weeks (CPT[®] 76816 if a complete ultrasound was done previously and/or CPT[®] 76817)
 - BPP (CPT[®] 76818 or CPT[®] 76819) or a modified BPP (CPT[®] 76815) weekly, starting at 32 weeks or sooner if indicated (other high-risk concerns)
 - Follow-up ultrasound can be performed at any time if bleeding occurs (CPT[®] 76815 and/or CPT[®] 76817)
 - MRI Pelvis without contrast (CPT[®] 72195) if the ultrasound is indeterminate or advanced imaging is needed for surgical planning. MRI Pelvis without contrast (CPT[®] 72195) is the appropriate code if only placenta or maternal pelvis is imaged without fetal imaging
 - The uterus, tubes and ovaries arise out of the pelvis and are considered pelvic organs. If the uterus rises out of the pelvic cavity, the imaging field can be determined on scout films. Imaging of the abdomen is not supported for problems suspected to arise from the pelvis.
 - The scout images (CT) and localizer images (MRI) are used to define the imaging field that is relevant to anatomical structures of clinical interest. The imaging field is defined by this clinical question, not by the imaging procedure code. The imaging code indicates the general anatomical region but does not define the specific imaging protocol or sequences.

Background and Supporting Information

- ACOG recommendations for imaging during pregnancy and lactation:
 - Ultrasonography and magnetic resonance imaging (MRI) are not associated with risk and are the imaging techniques of choice for the pregnant patient, but they should be used prudently and only when use is expected to answer a relevant clinical question or otherwise provide medical benefit to the patient.
 - The use of gadolinium contrast with MRI should be limited; it should be used as a contrast agent in a pregnant female only if it significantly improves diagnostic performance and is expected to improve fetal or maternal outcome.

References (OB-21)

v1.0.2025

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Late-term/Post-term Pregnancy (OB-22)

Guideline	Page
Late-term/Post-term Pregnancy (OB-22.1).....	166
References (OB-22).....	167

Late-term/Post-term Pregnancy (OB-22.1)

OB.LT.0022.1.A

v1.0.2025

- Ultrasound is supported at ≥ 41 weeks gestation
 - CPT[®] 76816 (if ≥ 2 weeks since last follow up ultrasound).
 - Twice weekly BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP CPT[®] 76815

Background and Supporting Information

- In late-term/post-term pregnancies, uterine artery Doppler velocimetry (CPT[®] 93976) has not been found to be useful. Per SMFM - uterine artery Doppler has limited diagnostic accuracy and clinical utility in predicting FGR, SGA birth, and perinatal mortality.

References (OB-22)

v1.0.2025

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Preterm/Prelabor Rupture of Membranes (OB-23)

Guideline	Page
Current Preterm/Prelabor Rupture of Membranes (PPROM) (OB-23.1).....	169
Current Prelabor Rupture of Membranes (PROM) (OB-23.2).....	170
References (OB-23).....	171

Current Preterm/Prelabor Rupture of Membranes (PPROM) (OB-23.1)

OB.RM.0023.1.A

v1.0.2025

- If suspected preterm/prelabor rupture of membranes (<37 weeks) perform:
 - One time CPT[®] 76815 for quick look for AFI check, **OR**
 - If confirmed oligohydramnios or PPRM CPT[®] 76816 or CPT[®] 76811 if detailed anatomy scan not previously performed¹ for delivery planning
 - This will likely result in a hospital admission for evaluation and monitoring until delivery. Only in rare cases is outpatient monitoring performed.
- See **Amniotic Fluid Abnormalities (OB-17.1)**

Current Prelabor Rupture of Membranes (PROM) (OB-23.2)

OB.RM.0023.2.A

v1.0.2025

- If ≥ 37 weeks and PROM is suspected
 - One time CPT[®] 76815 for quick look for AFI check, **OR**
 - CPT[®] 76816 (If confirmed oligohydramnios or PROM), for delivery planning
 - This will likely result in a hospital admission for delivery

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).

References (OB-23)

v1.0.2025

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3. ACOG Practice Bulletin No. 217: Prelabor Rupture of Membranes. *Obstetrics & Gynecology*. March 2020;135:e80–e97. Reaffirmed 2023. doi: 10.1097/AOG.0000000000003701
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Previous C-section or History of Uterine Scar (OB-24)

Guideline	Page
Previous C-section or History of Uterine Scar (OB-24.1).....	173
References (OB-24).....	175

Previous C-section or History of Uterine Scar (OB-24.1)

OB.CS.0024.1.A

v1.0.2025

Previous Cesarean section and/or uterine scar

- Report one of the following:
 - Complete first trimester ultrasound CPT[®] 76801 [plus CPT[®] 76802 for each additional fetus] if <14 weeks and a complete ultrasound has not yet been performed, and/or CPT[®] 76817 for a transvaginal ultrasound
 - CPT[®] 76801 is preferred for dating, but if this is unable to be completed then CPT[®] 76815 and/or CPT[®] 76817 for a transvaginal ultrasound is indicated
- For a normal/low risk pregnancy report a fetal anatomy ultrasound CPT[®] 76805 if ≥16 weeks. If pregnancy is high risk report a detailed fetal anatomy ultrasound CPT[®] 76811 if ≥16 weeks – See **High Risk Pregnancy (OB-9)**
- Growth scan (CPT[®] 76816) in the early third trimester (between 28-32 weeks) **and**
- CPT[®] 76816 once in the late third trimester for delivery planning (36-38 weeks)
- Transvaginal ultrasound, CPT[®] 76817 if poor visualization of the lower uterine segment or if uterine wall thinning (dehiscence) is suspected.
- If a cesarean scar pregnancy⁵ is suspected, can perform:
 - CPT[®] 76817 with or without CPT[®] 93976 (limited duplex scan) and/or
 - CPT[®] 76376 or CPT[®] 76377
 - MRI Pelvis without contrast (CPT[®] 72195) if inconclusive or equivocal ultrasound
- See **3D and 4D Rendering (OB 28.12)** and **Suspected PAS (OB-21.7.1)**

Background and Supporting Information

- Cesarean scar pregnancy (CSP) is a complication in which an early pregnancy implants in the fibrous tissue of a prior uterine scar. Because of the fibrous nature of scar tissue, a CSP is at risk for dehiscence, placenta accreta and hemorrhage. Females who consider pregnancy after a CSP are at significant risk of recurrence.
- A combination of transvaginal and transabdominal, combined with color Doppler ultrasound imaging are recommended for diagnosis.
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).

- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.

References (OB-24)

v1.0.2025

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Termination of Pregnancy – Imaging (OB-25)

Guideline	Page
Imaging for Planned Pregnancy Termination (OB-25.1).....	177
References (OB-25)	178

Imaging for Planned Pregnancy Termination (OB-25.1)

OB.PT.0025.1.A

v1.0.2025

- State and Federal legislation supersede these guidelines, see individual state regulations. For states in which pregnancy termination is supported by law, imaging as follows:
- For a planned pregnancy termination, ultrasound can be performed to determine intrauterine pregnancy and gestational age.
 - Complete first trimester anatomy ultrasound CPT® 76801 if <14 weeks and/or CPT® 76817, or
 - CPT® 76815 and/or CPT® 76817, or
 - If ≥14 weeks, CPT® 76805 (or CPT® 76811 if otherwise indicated, e.g. termination planned for chromosomal or structural fetal anomaly) can be performed (there may be State mandated imaging prior to termination).
- Routine follow-up is not usually necessary after uncomplicated medical abortion.
- For suspected medical abortion complication See **Spontaneous Abortion/Threatened/Missed Abortion (OB 5.4)**

Background and Supporting Information

- In general, most ultrasounds are indicated for planned pregnancy termination regardless of clinical information provided. Imaging may be indicated to confirm EGA, placenta location, and/or fetal anomalies.
- Though routine follow-up is not usually necessary after uncomplicated medical abortion, if otherwise medically indicated or preferred by the patient, follow-up ultrasound assessment after a medically induced termination can be considered
- CPT® 76815 should never be reported with complete studies CPT® 76801/CPT® 76802, CPT® 76805/CPT® 76810 or CPT® 76811/CPT® 76812 or with CPT® 76816 or BPP (CPT® 76818 and CPT® 76819).
- Though fetal anatomy survey (CPT® 76805/CPT® 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation

References (OB-25)

v1.0.2025

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Trauma (OB-26)

Guideline	Page
Trauma – Imaging (OB-26.1).....	180
References (OB-26).....	182

Trauma – Imaging (OB-26.1)

OB.TR.0026.1.A

v1.0.2025

Prior to 13 weeks:

- Blunt trauma in the first trimester (prior to 13 weeks) generally does not cause pregnancy loss with the exception of profound hypotension:
 - No imaging is indicated unless there is cramping and/or bleeding.

Between 13-20 weeks gestation:

- CPT[®] 76815 and/or CPT[®] 76817 can be performed, **or**
- Complete first trimester anatomy ultrasound CPT[®] 76801 and/or CPT[®] 76817 if complete ultrasound has not yet been performed, and is <14 weeks **or**
- CPT[®] 76805 (plus CPT[®] 76810 for each additional fetus) (or CPT[®] 76811/CPT[®] 76812 if otherwise indicated) if ≥14 weeks, when complete fetal anatomic scan has not yet been performed

After 20 weeks:

- CPT[®] 76805 (or CPT[®] 76811 plus CPT[®] 76812 for each additional fetus if otherwise indicated) when a fetal anatomic scan has not yet been performed, **or**
- CPT[®] 76815 and/or CPT[®] 76817 **or**
- CPT[®] 76816
- Additionally, starting at 26 weeks, BPP (CPT[®] 76818 or CPT[®] 76819) or modified BPP (CPT[®] 76815) can be considered
- CPT[®] 76821 if vaginal bleeding with +KB (Kleihauer-Betke) (if feto-maternal hemorrhage - at risk for fetal anemia and hydrops)
- Other advanced imaging may be indicated. See **Blunt Abdominal Trauma (AB-10.1)**

Background and Supporting Information

- Typically all components of the BPP (CPT[®] 76818 and CPT[®] 76819), such as breathing, are not present until ≥26 weeks gestation. However, a modified BPP (CPT[®] 76815) can be utilized sooner in certain high risk cases but should not be done prior to 22 weeks.
- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).

- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation.
- ACOG recommendations for imaging during pregnancy and lactation:
 - Ultrasonography and magnetic resonance imaging (MRI) are not associated with risk and are the imaging techniques of choice for the pregnant patient, but they should be used prudently and only when use is expected to answer a relevant clinical question or otherwise provide medical benefit to the patient.
 - With few exceptions, radiation exposure through radiography (Xrays), computed tomography (CT) scan, or nuclear medicine imaging techniques is at a dose much lower than the exposure associated with fetal harm. If these techniques are necessary in addition to ultrasound or MRI or are more readily available for the diagnosis in question, they should not be withheld from a pregnant patient.
 - The use of gadolinium contrast with MRI should be limited; it should be used as a contrast agent in a pregnant female only if it significantly improves diagnostic performance and is expected to improve fetal or maternal outcome.
 - With regards to iodinated IV contrast media, “it is generally recommended that contrast only be used if absolutely required to obtain additional diagnostic information that will affect the care of the fetus or woman during pregnancy”.

References (OB-26)

v1.0.2025

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Unequal Fundal Size and Dates (OB-27)

Guideline	Page
Unequal Fundal Size and Dates (OB-27.1).....	184
References (OB-27).....	185

Unequal Fundal Size and Dates (OB-27.1)

OB.US.0027.1.A

v1.0.2025

Unequal fundal size is defined as ≥ 3 cm difference between fundal height measurement and gestational age (weeks), at ≥ 22 weeks gestation

- One ultrasound can be performed (CPT[®] 76805) if complete fetal anatomic scan is planned and has not been performed **or**
- CPT[®] 76816 if complete anatomy scan or detailed anatomy ultrasound (CPT[®] 76805/CPT[®] 76811) has been done previously
- Where fundus cannot be adequately palpated such as in obesity, leiomyomas, multiple gestations, See appropriate chapter

Background and Supporting Information

- CPT[®] 76815 should never be reported with complete studies CPT[®] 76801/CPT[®] 76802, CPT[®] 76805/CPT[®] 76810 or CPT[®] 76811/CPT[®] 76812 or with CPT[®] 76816 or BPP (CPT[®] 76818 and CPT[®] 76819).
- Though fetal anatomy survey (CPT[®] 76805/CPT[®] 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation

References (OB-27)

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Procedure Coding Basics for Established Pregnancy (OB-28)

Guideline	Page
Procedure Coding Basics for Established Pregnancy General Considerations (OB-28.1).....	187
Required Elements for Complete First Trimester Ultrasound (OB-28.2).....	188
Required Elements for Second or Third Trimester Fetal Anatomic Evaluation Ultrasound (OB-28.3).....	189
Required Elements for a Detailed Fetal Anatomic Evaluation Ultrasound (OB-28.4).....	191
Fetal Nuchal Translucency (OB-28.5).....	193
Limited and Follow-up Studies (OB-28.6).....	194
Obstetric Transvaginal Ultrasound (OB-28.7).....	195
Biophysical Profile (BPP) (OB-28.8).....	196
Fetal Doppler (OB-28.9).....	197
Duplex Scan (OB-28.10).....	198
Fetal Echocardiography (OB-28.11).....	199
3D and 4D Rendering (OB-28.12).....	200
Required Elements for a Detailed First Trimester Fetal Anatomic Evaluation Ultrasound (OB-28.13).....	201
References (OB-28).....	203

Procedure Coding Basics for Established Pregnancy General Considerations (OB-28.1)

OB.PC.0028.1.A**v1.0.2025**

- All obstetric ultrasound studies require permanently recorded images which may be stored on film or in a Picture Archiving and Communication System (PACS). Obstetric ultrasound services may not be billed without image recording.
- Ultrasound procedure codes include the preparation of a required final written report which should be included in the patient's medical record.
 - Each procedure code has specific required elements which are described in this section.
 - The report should document the results of the evaluation of each element or the reason any element is non-visualized.
 - Documentation of less than the required elements requires the billing of the "limited" code for that anatomic region.
 - Only one (1) limited exam should be billed per encounter.
- The use of a hand-held or any Doppler device that does not create a hard-copy output is considered part of the physical examination and is not separately billable.

Required Elements for Complete First Trimester Ultrasound (OB-28.2)

OB.PC.0028.2.A

v1.0.2025

CPT® Code Guidance

- Complete First Trimester Ultrasound [CPT® 76801 and CPT® 76802 (for each additional fetus)] can be performed up to and including 13 6/7 weeks gestation and is defined in CPT® as including the following elements:
 - Number and size of gestational sacs and fetuses
 - Survey of visible fetal anatomic structures and placental evaluation when possible
 - Qualitative assessment of amniotic fluid volume/gestational sac shape
 - Examination of maternal uterus and adnexa
- Complete First Trimester Ultrasound (CPT® 76801 and CPT® 76802) should only be reported once per pregnancy/per practice/facility unless the mother changes to a new medical caregiver at a new practice/facility and there is a new medical indication for ultrasound.

Required Elements for Second or Third Trimester Fetal Anatomic Evaluation Ultrasound (OB-28.3)

OB.PC.0028.3.A

v1.0.2025

CPT® Code Guidance

- A complete second or third trimester fetal anatomic evaluation ultrasound (CPT® 76805 and CPT® 76810 for each additional fetus) is defined in CPT® as including the following elements:
 - **Head, face, and neck:** Lateral cerebral ventricles; Choroid plexus; Midline falx; Cavum septum pellucidum; Cerebellum; Cistern magna; Upper lip: A measurement of the nuchal fold may be helpful during a specific age interval to assess the risk of aneuploidy
 - **Chest/Heart:** Four-chamber view; Left and Right ventricular outflow tracts
 - **Abdomen:** Stomach (presence, size, and situs); Kidneys; Urinary bladder; Umbilical cord insertion site into the fetal abdomen and number of vessels
 - **Spine:** Cervical, thoracic, lumbar, and sacral spine
 - **Extremities:** Legs and arms
 - **Genitalia:** (In multiple gestations and when medically indicated)
 - **Placenta:** Location; Relationship to internal os; Appearance; Placental cord insertion (when possible) and overall standard evaluation
 - Fetal number and Presentation
 - Qualitative or semi-qualitative estimate of amniotic fluid
 - **Maternal anatomy:** Cervix (transvaginal if cervical length on transabdominal ultrasound appears shortened (≤ 3.6 cm) and transvaginal exam would be needed to guide therapeutic recommendations), Uterus, and Adnexa See **Cervical Length Screening (OB-7.3)**
 - **Fetal Biometry:** Biparietal diameter, Head circumference, Femur length, Abdominal circumference, and Fetal weight estimate.

CPT® Code Guidance

- CPT® 76805 and CPT® 76810 **should only be used once per pregnancy per practice/facility** unless the mother changes to a new medical caregiver at a new practice/facility and there is a new medical indication for ultrasound.
- Though fetal anatomy survey (CPT® 76805/CPT® 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation. This timing allows for a survey of fetal anatomy and an accurate estimation of gestational age
- Follow-up studies to CPT® 76805/CPT® 76810 should be coded as CPT® 76815 or CPT® 76816.

Required Elements for a Detailed Fetal Anatomic Evaluation Ultrasound (OB-28.4)

OB.PC.0028.4.A

v1.0.2025

CPT® Code Guidance

- Detailed fetal anatomy scan (CPT® 76811 and CPT® 76812 for each additional fetus) is generally performed by those with special skills to perform this study, such as Maternal Fetal Medicine specialists (Perinatologists), or Radiologists **with advanced training in fetal imaging**.
- CPT® 76811 and CPT® 76812 are defined in CPT® as including all of the requirements listed for CPT® 76805 and CPT® 76810. In addition, the report must document detailed anatomic evaluation of the following elements:
 - Head, face, and neck:** 3rd and 4th ventricles; Lateral ventricles; Cerebellar lobes, vermis, and cisterna magna; Corpus callosum; Integrity and shape of cranial vault; Brain parenchyma; Neck; Profile; Coronal face (nose/lips/lenses); Palate, maxilla, mandible, and tongue; Ear position and size; Orbits
 - Chest/Heart:** Aortic arch; Superior and inferior vena cava; 3-vessel view; 3-vessel and trachea view; Lungs; Integrity of diaphragm; Ribs
 - Abdomen:** Small and large bowel; Adrenal glands; Gallbladder; Liver; Renal arteries; Spleen; Integrity of abdominal wall
 - Spine:** Integrity of spine and overlying soft tissue; Shape and curvature
 - Extremities:** Number: architecture and position; Hands; Feet; Digits: number and position
 - Genitalia: Gender
 - Placenta:** Masses; Placental cord insertion; Accessory/succenturiate lobe with location of connecting vascular supply to primary placenta
 - Biometry:** Cerebellum; Inner and outer orbital diameters; Nuchal thickness (16 to 20 wk); Nasal bone measurement (15 to 22 wk); Humerus; Ulna/radius; Tibia/fibula
 - Maternal Anatomy:** Cervix (transvaginal if cervical length on transabdominal ultrasound appears shortened (≤ 3.6 cm) and transvaginal exam would be needed to guide therapeutic recommendations); Uterus; Adnexa See **Cervical Length Screening (OB-7.3)**

CPT® Code Guidance

- CPT® 76811 and CPT® 76812 **should only be used once per pregnancy per practice/ facility** unless the mother changes to a new medical caregiver at a new facility and there is a new medical indication for ultrasound.
- Though fetal anatomy survey (CPT® 76805/CPT® 76811) can be performed as early as 14 weeks gestation, per ACOG, in the absence of other specific indications, it is optimally performed at 18 to 22 weeks of gestation. This timing allows for a survey of fetal anatomy and an accurate estimation of gestational age
- Follow-up studies to CPT® 76811/CPT® 76812 should be coded as CPT® 76815 or CPT® 76816.
- In circumstances where the individual is deemed to have an increased risk for a fetal abnormality and does not have access to a provider who can perform a fetal anatomic ultrasound examination (CPT® 76811) due to geographic or other constraints, a standard (after first trimester) fetal anatomic ultrasound examination (CPT® 76805) can be authorized instead.

Fetal Nuchal Translucency (OB-28.5)

OB.PC.0028.5.A

v1.0.2025

CPT® Code Guidance

- CPT® 76813 and CPT® 76814 (for each additional fetus) describe ultrasound measurement of the clear (translucent) space at the back of the fetal neck.
 - The first trimester screening can be performed if CRL measures 44-83 mm (typically between 10 4/7 and 14 weeks' gestation).
 - Abnormal Fetal Nuchal Translucency scan (NT $\geq$ 3 mm or above the 95th percentile for the CRL) suggests an increased risk for aneuploidy, or may be a marker for cardiac defects, abdominal wall defects, diaphragmatic hernia, and genetic syndromes in euploid fetuses during current pregnancy.
 - The sonographer performing the study and/or the physician interpreting the study must be credentialed by the Maternal Fetal Medicine Foundation or Nuchal Translucency Quality Review Program (NTQR)
-
- The use of ultrasound codes (CPT® 76801/CPT® 76802) should be indication driven and should not be routinely done whenever an ultrasound for nuchal translucency (CPT® 76813/CPT® 76814) is requested. In cases where there is either a maternal and/or fetal indication, then the CPT® 76801 code can be billed along with the nuchal translucency screening (CPT® 76813/CPT® 76814).
 - Fetal Nuchal Translucency (CPT® 76813 and CPT® 76814) should only be reported once per pregnancy

Background and Supporting Information

- The first trimester screening is typically done between 11 and 13 6/7 weeks but can be performed if the crown rump length (CRL) measures between 44-83 mm (typically at gestational age range 10 4/7 to 14 weeks)
- **Required elements of the CPT® 76813 ultrasound code include:**
 - Fetal crown-rump measurement
 - Observation of fetal cardiac activity
 - Observation of the embryo at high magnification until the embryonic neck is in a neutral position and spontaneous embryonic movement allows for differentiation between the outer edge of the nuchal skin and the amnion
 - Measurement of the largest distance between the inner borders of the fetal nuchal translucency

Limited and Follow-up Studies (OB-28.6)

OB.PC.0028.6.A

v1.0.2025

CPT® Code Guidance

- **CPT® 76815** describes a **limited** or “quick look” study used when medically indicated to report one or more of the elements listed in the code definition, i.e. “fetal heartbeat”, placental location, for viability/dating (when indicated), or fluid check (re: modified BPP which is NST with CPT® 76815)
 - Reported only once, regardless of the number of fetuses, and only once per date of service
 - CPT® 76815 should never be reported with complete studies CPT® 76801/CPT® 76802, CPT® 76805/CPT® 76810 or CPT® 76811/CPT® 76812 or with CPT® 76816 or BPP (CPT® 76818 and CPT® 76819).
- **CPT® 76816** describes a **follow-up** ultrasound (eg, re-evaluation of fetal size by measuring standard growth parameters and amniotic fluid volume, re-evaluation of organ system(s) suspected or confirmed to be abnormal on a previous scan), trans-abdominal approach, per fetus.
 - The use of this CPT code is reserved for subsequent follow up ultrasound only; i.e. complete ultrasound (CPT® 76801, or CPT® 76805/CPT® 76811) must have been performed previously.
 - Components include: Focused assessment of fetal biometry, amniotic fluid volume and/or a detailed re-examination of a specific organ or system known or suspected to be abnormal on prior ultrasound.
 - CPT® 76816 should be reported once per fetus evaluated in follow-up.
 - CPT® 76816 should never be reported with complete studies (CPT® 76801, CPT® 76802, CPT® 76805/CPT® 76810, or CPT® 76811/CPT® 76810) or with a limited study (CPT® 76815).
 - CPT® 76816 should not be performed prior to a CPT® 76801 and/or an anatomy scan CPT® 76805 (normal pregnancy) or Detailed anatomy scan CPT® 76811 (high risk pregnancy), and is typically not performed before 14 weeks gestation.

Obstetric Transvaginal Ultrasound (OB-28.7)

OB.PC.0028.7.A
v1.0.2025

CPT® Code Guidance
• CPT® 76817 is used to report an obstetrical transvaginal ultrasound for cervical length assessment (when indicated), or in certain circumstances with poor visualization by transabdominal US assessment.
• CPT® 76817 is reported only once regardless of the number of fetuses.

Biophysical Profile (BPP) (OB-28.8)

OB.PC.0028.8.A
v1.0.2025

CPT® Code Guidance
• CPT® 76818 includes non-stress testing. Supported ≥26 weeks.
• CPT® 76819 does not include the non-stress testing portion. Supported ≥26 weeks
• CPT® 76815 is a modified BPP – includes NST and AFI. Supported ≥22 weeks
• BPPs performed on more than one fetus, should be reported separately.
• Modified BPP (CPT® 76815) should be reported only once, regardless of the number of fetuses, and only once per date of service.
• Modified BPP (CPT® 76815) should never be reported with complete studies, CPT® 76805/CPT® 76810 or CPT® 76811/CPT® 76812 or with CPT® 76816 or BPP (CPT® 76818 and CPT® 76819).

Background and Supporting Information

- The BPP is designed to predict the presence or absence of fetal asphyxia and, ultimately the risk of fetal death. A BPP is scored out of 10 with each parameter given a score of 0 or 2. A score ≥8 is considered normal. The following parameters are evaluated:
 - Fetal breathing movements
 - Gross fetal body movements
 - Fetal tone
 - Amniotic fluid volume, at least one vertical pocket 2 x 2 cm
 - Reactive FHR (non-stress testing portion)
- CPT® 76815 is used to assess the AFI portion of a modified BPP (NST + AFI)
- Typically all components of the BPP (CPT® 76818 and CPT® 76819), such as breathing, are not present until ≥26 weeks gestation. However, a modified BPP (CPT® 76815) can be utilized sooner in certain high risk cases but should not be done prior to 22.
- If BPP ≤6, repeat BPP in ≤24 hours

Fetal Doppler (OB-28.9)

OB.PC.0028.9.A

v1.0.2025

CPT® Code Guidance

- CPT® 76820 describes Doppler velocimetry of the umbilical artery (UA Doppler)
 - Utilized for known FGR; See **Fetal Growth Restriction Current Pregnancy (OB-20.1)** and known oligohydramnios See **Amniotic Fluid Abnormalities (OB-17.1)**, and is typically performed ≥ 22 weeks gestation.
 - Can be performed with known twin to twin transfusion or known discordant twins (See **Multiple Gestations (OB-11)**)
 - Its use to predict preeclampsia, and stillbirth is considered investigational.
- CPT® 76821 describes Doppler velocimetry of the middle cerebral artery (MCA Doppler).
 - Performed to evaluate a fetus at risk for anemia due to Rhesus isoimmunization/alloimmunization (as a substitute for amniocentesis), Twin anemia polycythemia sequence and non-immune hydrops caused by parvovirus B19 infection or any other known acquired or congenital cause of fetal anemia. See **Alloimmunization/Rh Isoimmunization (OB-16.1)**, **Other Causes of Fetal Anemia (OB-16.5)**, and **Multiple Gestations (OB-11)**
- MCA Doppler (CPT® 76821) assessment can be performed if the fetus is at high risk for fetal anemia due to other pregnancy complications, e.g. chorioangioma, umbilical vein varix, or finding of sustained fetal tachyarrhythmia or bradyarrhythmia or a known congenital heart defect with suspected heart failure in the fetus
 - See **Indications for Fetal Conditions (OB-12.2)**, **Alloimmunization/Rh Isoimmunization (OB-16.1)**, and **Placental and Cord Abnormalities (OB-21)**
- SMFM suggest that ductus venosus, middle cerebral artery, or uterine artery Doppler use for routine clinical management of early- or late-onset FGR *is not recommended*

Duplex Scan (OB-28.10)

OB.PC.0028.10.A**v1.0.2025**

- A Duplex scan describes an ultrasonic scanning procedure for characterizing the pattern and direction of blood flow in arteries and veins. It produces real-time images integrating a B-mode two dimensional vascular structure, Doppler spectral analysis, and color flow Doppler imaging.
- **CPT® 93976** describes a limited duplex scan and is used during pregnancy to report uterine artery Doppler studies (done to report fetal umbilical-placental flow evaluation, accreta or other placental or cord abnormalities).
 - **CPT® 93975** describes a complete duplex scan. This code is **NOT** used in obstetrical imaging.
- The minimal use of color Doppler alone, when performed for anatomical structure identification, during a standard ultrasound procedure, is not separately reimbursable.
- SMFM state that uterine artery Doppler has limited diagnostic accuracy and clinical utility in predicting FGR, SGA birth, and perinatal mortality. As such, its use for screening in high risk groups is not recommended

Fetal Echocardiography (OB-28.11)

OB.PC.0028.11.A
v1.0.2025

CPT® Code Guidance
- CPT® 76825 describes a complete fetal echocardiography. - CPT® 76827 describes a complete Doppler echocardiography, fetal, pulsed wave and/or continuous wave with spectral display - It is usually billed along with CPT® 76825
- CPT® 76826 describes a follow-up or repeat fetal echocardiogram - It should never be billed with CPT® 76825 or more than once per fetus on any date of service - CPT® 76828 is a follow-up or repeat Doppler fetal echocardiogram
- Procedure code (CPT® 76827 or CPT® 76828) includes the evaluation of veins, arteries, and valves, and covers Doppler evaluation of the ductus venosus, ductus arteriosus, and PR Interval measurement as well as other vessels. Guidelines do not support the additional billing of CPT® 76820 and/or CPT® 76821. - It is inappropriate to report codes CPT® 76825 – CPT® 76828 for the routine monitoring of fetal heart tones using a hand-held or any Doppler device that does not create a hard-copy output. Such fetal heart tone monitoring is considered part of the physical examination and is not separately billable.
- CPT® 93325 is used to report color mapping in conjunction with fetal echocardiography procedures CPT® 76825 – CPT® 76828. - The use of color Doppler (CPT® 93325) alone, when performed for anatomical structure identification, during a standard ultrasound procedure, is not separately reimbursable.

3D and 4D Rendering (OB-28.12)

OB.PC.0028.12.A

v1.0.2025

- There is currently insufficient data to generate appropriateness criteria for the use of 3D and 4D rendering in conjunction with Obstetrical ultrasound imaging.
 - Per ACOG, despite the technical advantages of 3-dimensional ultrasonography, proof of a clinical advantage of 3-dimensional ultrasonography in prenatal diagnosis, in general, is still lacking.
- However, 3D-4D (CPT[®] 76376 or CPT[®] 76377) rendering can be considered in certain situations of abnormal pregnancy implantation like suspected C-section scar pregnancies or suspected cornual (interstitial) ectopic pregnancy, or to locate an IUD. See **Locate an Intrauterine Device (IUD) (OB-3.1)**, **Previous C-section or History of Uterine Scar (OB 24.1)**, **Ectopic Pregnancy (OB 5.3)**, and **Uterine Anomalies (PV-14.1)** in the Pelvis Imaging Guidelines
- 3D-4D (CPT[®] 76376 or CPT[®] 76377) rendering can be used for surgical planning with diagnosis of complex CHD in the fetus or for surgical planning of other complex fetal malformations.¹⁶

Required Elements for a Detailed First Trimester Fetal Anatomic Evaluation Ultrasound (OB-28.13)

OB.PC.0028.13.A
v1.0.2025

CPT® Code Guidance
Detailed first trimester fetal anatomy scan, CPT® 76801 plus CPT® 76813 (and CPT® 76802 plus CPT® 76814 for each additional fetus) can be performed between 12 weeks 0 days and 13 weeks 6 days when indicated (See Detailed First Trimester Fetal Anatomic Scan (OB-9.12) for indications).
A Detailed first trimester fetal anatomy scan must document detailed anatomic evaluation of the following elements: General: Cardiac activity; Number of fetuses and gestational sacs Head, face, and neck: Cranial bones (calvarium); falx cerebri; choroid plexus; transthalamic flax cerebri; Thalami; Posterior fossa; brain stem; 4th ventricles; cisterns magna; Corpus callosum; nasal bone; profile; maxilla; mandible; evaluation for cystic hygroma, dilated jugular; lymphatic sacs; nuchal translucency evaluation Chest/Heart: Cardiac position and axis; 4-chamber view; 3- vessel and trachea view; Lungs; Integrity of diaphragm; Ribs Abdomen: Stomach; Liver; Cord insertion into abdominal wall; Bladder; Umbilical arteries; Integrity of abdominal wall Spine: Integrity of spine and overlying soft tissue; Shape and curvature Extremities: Number: architecture and position; Hands; Feet: number and position Placenta: Position; Umbilical cord insertion; Echo texture Biometry: Crown-rump length Maternal Anatomy: Myometrial masses; Mulllein duct anomalies; Uterus; Ovaries; Adnexa and cul-de-sac

CPT® Code Guidance

- A Detailed first trimester fetal anatomy scan, CPT® 76801 plus CPT® 76812 (and CPT® 76802 plus CPT® 76814 for each additional fetus) should only be reported once per pregnancy/per practice/facility unless the mother changes to a new medical caregiver at a new practice/facility and there is a new medical indication for ultrasound.
- It is generally performed by those with special skills to perform this study, such as Maternal Fetal Medicine specialists (Perinatologists), or Radiologists **with advanced training in fetal imaging**.

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v1.0.2025

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