CONTROVERSIES: THE PREGNANT PATIENT FOR NON-OBSTETRIC SURGERY

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(* I have no conflicts to disclose.*)

GOALS & OBJECTIVES

Upon completion of this presentation, participants will be able to:
1. Cite guidelines from ASA and ACOG relating to non-obstetric surgery during pregnancy.
2. Apply physiologic principles and knowledge of teratogenic effects to develop an anesthetic plan.
3. Adapt the broad anesthetic plan to special situations such as trauma, cardiac surgery, neurosurgical procedures, or laparoscopy.

THE PUBLIC IS CONCERNED

“The fetus usually dies from the anesthesia administered to the mother before the procedure begins . . . The intravenous anesthetic administered to the woman during the procedure induces a medical coma in the fetus and eventually a neurological fetal demise.”

Congressional Testimony 6/23/95

ANESTHESIOLOGISTS ARE CONCERNED

A retrospective survey of female veterinarians related preterm delivery (< 37 weeks) to self-reported occupational anesthetic exposures.
• OR 2.5 for those who performed surgery without scavenging anesthetic gases vs. with scavenging
• OR 3.69 for those working > 45 hours a week.
Obstet Gynecol 2009;113:1008
ASA & ACOG JOINT STATEMENT

1. Obtain a consult from obstetrics preop. They should be readily available intraop.
2. No anesthetic agents are teratogenic.
3. FHR monitoring may assist in positioning, cardiorespiratory management, delivery option.
4. Perform surgery where there are obstetric and pediatric services.

JOINT STATEMENT (cont)

5. Pre and post-op FHR is adequate if pre-viable.
6. A qualified individual should be readily available to interpret if continuous FHR monitoring is used.
7. A pregnant woman should never be denied indicated surgery. Postpone elective surgery until after delivery, non-urgent until 2nd trimester.

Controversial Issues Surrounding Anesthetic Care During Pregnancy

Controversy #1. Which patients need to have preoperative pregnancy testing?

Case: A healthy 25-year old woman presents to your operating room for knee arthroscopy on an outpatient basis. Does she need to have a pregnancy test?

50% of pregnancies in the U.S. are unplanned!

Pregnancy Testing

For one year, all women of childbearing age having ambulatory surgical procedures had preoperative pregnancy testing.

- 7/2056 (0.3%) of tests were positive.
- All 7 patients elected to cancel surgery (2 were infertility procedures).
- The estimated cost to diagnose one pregnancy was $2879.

Anesthesiology 1995;83:690
PREGNANCY TESTING
A specialty orthopedic hospital initiated mandatory pregnancy testing. After 1 year:
• 2588 tested—8 positives, but 4 were false positives.
• 3 of the false positives subsequently had a negative serum hCG, and surgery then proceeded.
• 4/2588 (0.15%) of tests were true positives and their surgery was canceled.
• NNT for 1 true positive : 647
• Cost for 1 true positive : $3273
Anesth Analg 2008;106:1127

PREGNANCY TESTING
Discussion: “...should a spontaneous abortion occur after surgery, or the baby be born with a congenital anomaly, this may be attributed to the surgery or anesthetic……screening may decrease litigation, although potential cost savings are difficult to quantify.”
Anesth Analg 2008;106:1127

PREGNANCY TESTING
• A review of 2 years of mandatory pregnancy testing in a freestanding pediatric hospital found that 2.4% of patients 16 and older were positive. None were positive in patients less than 15 years of age, so overall incidence was 1.3%.
• Their conclusions: A policy of mandatory pregnancy testing in patients aged 15 and older is advisable. Specific written consent for the test is not necessary, but proper notification processes must be established.
Anesth Analg 1996;82:4-7

PREGNANCY TESTING
What about ethical and privacy concerns?
1. Can you test without the patient’s consent?
2. If positive, can/should you inform a minor’s parents?
3. Will you cancel the case if they refuse testing?
4. Is testing an entire class of patient (i.e., females) just to protect the provider from liability ethical?
5. How would your anesthetic be any different? Since there are no modern anesthetics that will affect the pregnancy or fetus, testing is not medically indicated.

PREGNANCY TESTING
Do pregnant women have greater morbidity after surgery than non-pregnant?
• Using the NSQIP database from 2005-9, 857 appendectomy and 436 cholecystectomy cases in pregnancy were reviewed.
• Morbidity was no different than non-pregnant.
• Pregnant women were more likely to be infected at the time of surgery.
Obstet Gynecol 2011;118:1261

PREGNANCY TESTING
From the ASA Practice Advisory for Pre-anesthesia Evaluation:
“The Task Force recognizes that patients may present for anesthesia with early undetected pregnancy. The Task Force believes that the literature is inadequate to inform patients or physicians on whether anesthesia causes harmful effects on early pregnancy. Pregnancy testing may be offered to female patients of childbearing age and for whom the result would alter the patient’s management.”
Anesthesiology 2012; 116: 522
PREGNANCY TESTING

New problem – EMRs allow us to cut-and-paste.

- **Case:** A woman presented for D&C due to a history of heavy menstrual bleeding. Pre-op pregnancy test was reportedly negative. At the time of D&C, an 8 week gestation was identified – a much desired pregnancy. It was later discovered that her preoperative assessment was populated with a previous negative test result, rather than the current positive pregnancy test.

ASA Newsletter; September 2014: 42

Controversy #2. Are benzodiazepines and nitrous oxide teratogenic, or are they safe to use during pregnancy?

Case: A woman at 12 weeks gestation requests general anesthesia for a cervical cerclage placement scheduled for ~ 20 minutes. She is extremely anxious about the procedure and asks you for preoperative sedation.

DOCUMENTED TERATOGENS

ACE Inhibitors | Lead
Alcohol | Lithium
Androgens | Mercury
Antithyroid Drugs | Phenytoin
Chemotherapy | Streptomycin
Cocaine | Thalidomide
Coumadin | Trimethadione
Diethylstilbestrol | Valproic Acid
Isoretinoin

TERATOGENICITY

How long after a drug is marketed does it take to establish safety for use in pregnancy?

- Experts assessed 469 drugs approved by the FDA between 1980 and 2000, reviewing available studies.
- 91% of drugs were still classified as their risk of use during pregnancy being "undetermined".
- Inadequate information is available for women and their physicians to determine risks of most drugs.

Obstet Gynecol 2002;100:465

NITROUS OXIDE

Pregnant rats given nitrous oxide 75% for 24 hours on day 9 of gestation had a 4-fold increase in resorptions (abortions) and a 15-fold increase in anomalies when compared to rats given equi-anesthetic concentrations of xenon.

Science 1980;210:899
**NITROUS OXIDE**

Why might N₂O cause adverse effects?

N₂O inactivates vitamin B₁₂, a coenzyme of methionine synthetase, causing depression of methionine synthetase activity and potentially affecting production of thymidine and DNA.

**NITROUS OXIDE**

However, even very low concentrations of N₂O (<1%) will abolish methionine synthetase activity, yet it requires 24 hours of high N₂O concentrations (75%) to cause teratogenesis.

Are the adverse effects of nitrous oxide biochemical (reduced methionine synthetase activity), or could they be due to hemodynamic effects?

**NITROUS OXIDE**

• N₂O enhances adrenergic tone and causes vasoconstriction.
• Halothane (a sympatholytic) and other volatile anesthetics administered with N₂O protect against major and minor anomalies in rodents.

Teratology 1988;38:121

**ANESTHETIC EXPOSURE**

Self-reported occupational exposures during pregnancy from 7482 nurses in the Nurses’ Health Study II were used to investigate the risk of spontaneous abortion:
• 10% had spontaneous abortions < 20 weeks
• Exposure to anti-neoplastic drugs and sterilizing agents was associated with doubled risk.
• There was no association of early or late abortion with x-ray radiation or anesthetic gases.

Am J Obstet Gynecol 2012;206:327

**NITROUS OXIDE**

The largest retrospective study of exposure to surgery and anesthesia during pregnancy compared 5405 women who had surgery to case controls.
• 54% had GETA, 97% of those had N₂O
• No difference in stillbirth or anomalies, but...
• There were increases in IUGR, prematurity

Am J Obstet Gynecol 1989;161:1178
NITROUS OXIDE: SUMMARY

- Teratogenic effects in animal studies may be due to vasoconstriction and decreased uterine blood flow. Combine N₂O with a sympatholytic agent.
- Adverse human effects have never been documented, even in large outcome studies.
- Studies in modern hospital settings with O.R. scavenging do not show an association with nitrous oxide or volatile anesthetics and adverse pregnancy outcomes.

BENZODIAZEPINES

Two studies in 1975 reported an association between maternal exposure to benzodiazepines (Valium® and Librium®) and cleft lip and/or palate.

Int J Epidemiol 1975; 4:37
Lancet 1975; 306:478

But later work refuted these reports…..

BENZODIAZEPINES

- 611 infants with cleft lip and/or palate were compared to 2498 control infants with other birth defects.
- The risk of clefts was no different between women who were or were not exposed to diazepam during first trimester of pregnancy.
- For cleft lip ± palate: RR 0.8 (0.4-1.7).
- For cleft palate alone: RR 0.8 (0.2-2.5).

NEJM 1983; 309:1282

BENZODIAZEPINES

- An NIH-supported prospective study did not find any increased risk of cleft anomalies associated with diazepam use:
  - RR 1.2 versus controls
  - 95% CI 0.17-8.95

NEJM 1984; 311:919

BENZODIAZEPINES

- A meta-analysis of 7 cohort studies involving 1090 infants who were exposed to benzodiazepines found no increased risk of major malformations, or specifically oral clefts (RR 1.19, CI 0.34-4.15).
- “Even when the worst case scenario is assumed, benzodiazepines do not seem to be major human teratogens….”
  - BMJ 1998;317:839

ACOG Clinical Expert Series on Teratogenicity

“Anxiolytics (benzodiazepines): No evidence of significant risk of teratogenicity”

- Initial findings of clefts have not been confirmed by long-term follow-up studies.
- Overall results are reassuring, revealing no adverse effects on neurodevelopment.
- May be beneficial adjunct for hyperemesis (!)

Obstet Gynecol 2009;113:166
**NSAID’s IN PREGNANCY**
Database review of women taking NSAIDs during pregnancy:
- 22% used NSAIDs in first trimester; mainly ibuprofen, aspirin, naproxen
- No association with most anomalies
- Small ↑ risk of a few specific defects
  Am J Obstet Gynecol 2012;206:228

**ONDANSETRON**
Large retrospective review of a Danish database for fetal outcomes after use of ondansetron during pregnancy:
- There was no relationship with stillborn, miscarriage, any birth defects, preterm delivery, or small size
- Usually used during first trimester for severe nausea & vomiting (hyperemesis).

**OPIOIDS**
A Washington State database review of illicit and prescription maternal drug use:
- Rates ↑ from 2000→2008; mainly opioids
- Neonatal withdrawal 3.3 / 1000 births
- Newborns had lower birth weight, longer hospitalizations, more preterm births, feeding difficulties and respiratory issues.
  Obstet Gynecol 2012;119:924

**CANCER CHEMOTHERAPY**
Should aggressive chemo be used when breast cancer is diagnosed during pregnancy?
- 10 women received dose-dense and 99 received conventional chemotherapy
- There was no difference in birth weight, GA at delivery, IUGR, anomalies, maternal or fetal neutropenia.
  Obstet Gynecol 2012;120:1267

**CANCER AT DELIVERY**
21-year review of L&D management of women with cancer in a tertiary center:
- Incidence 0.1%, equally diagnosed before and during pregnancy
- 75% received regional for labor, 22% received general for cesarean
- Life-threatening complications with mediastinal tumors or metastases
  Int J Obstet Anesth 2012;24:524

**Controversy #3. When and how should fetal monitoring be used?**

Case 1: A semi-elective cholecystectomy done at 34 weeks gestation.

Case 2: An emergency femoral thrombectomy at 31 weeks gestation.

Case 3: A series of 5 ECTs performed between 17 and 19 weeks of gestation.
INTRA-OP MONITORING

- Blood pressure (normal or slightly above)
- Oxygenation, ventilation
- Temperature (avoid hyperthermia)
- Blood glucose for longer cases
- Fetal heart rate (FHR) > 24 weeks: intermittent, or continuous if possible
- FHR < 24 weeks: preop and postop check

FETAL MONITORING

- This should not be discussed and decided as a medico-legal issue! It is a medical issue.
- Monitoring may help assess placental perfusion and assure the intrauterine environment is optimized during positioning, induced hypotension, cardiopulmonary bypass, volume shifts / blood loss.
- It provides an important reassurance for the mother.
- But FHR monitoring is imprecise and not predictive of outcome.

CASE 1: FETAL MONITORING

A patient at 34 weeks gestation required cholecystectomy. During skin prep (before any surgical intervention), severe persistent fetal bradycardia occurred. An emergency cesarean was performed and the umbilical cord was tightly coiled and twisted.
- Apgar scores = 1 / 5 / 7
- Umbilical cord pH = 7.17 and 7.18
  Can J Anesth 2003;50:922

CASE 2: FETAL MONITORING

During the 30th week of an uncomplicated pregnancy, a patient underwent femoral thrombectomy under routine GETA. During surgery the fetal monitor showed absent variability, so an emergency cesarean delivery was performed for presumed fetal distress. Umbilical pH was 7.23 (normal), but the child required intubation for prematurity and ICU admission.
  Br J Anaesth 2001; 87:791

CASE 3: FETAL MONITORING

A series of ECTs was required in a woman between 17 and 19 weeks gestation. FHTs checked before and after the first 4 procedures were normal. FHTs monitored during the 5th procedure showed a severe deceleration. No intervention was done due to non-viability. She went on to deliver a normal healthy baby at 38 weeks.
  Acta Anaesthesiol Scand 2003;47:101
**FETAL MONITORING**

What should you do for intra-operative fetal distress?
- ↑ maternal FIO₂ and blood pressure.
- ↑ left uterine displacement or try tilting right.
- Move the surgeons or their retractors.
- Administer a tocolytic (e.g., terbutaline 0.25 mg).
- Document your efforts in the record.
** Consider preop / postop FHR monitoring for most cases, in consultation with the obstetrician.
** Remember: loss of BTBV is normal; decels are not.

**Controversy #4. Should pregnant patients > 24 weeks gestation have surgery in a specialty hospital without L&D coverage?**
(no fetal monitoring, no capability for a C/S, no neonatologists)

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**PERIOPERATIVE BACK-UP**

Case: A woman at 28 weeks gestation was evaluated for deteriorating vision, and a large meningioma was found on MRI. Urgent craniotomy was planned to preserve her sight. The surgery proceeded without fetal monitoring or provision for cesarean delivery as obstetric care was not available at the hospital where neurosurgery was performed.

Can J Anesth 2004;51:573

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**ACOG / ASA JOINT STATEMENT**

"Non-obstetric Surgery During Pregnancy"

“Due to the difficulty of conducting large-scale randomized clinical trials in this population, there are no data to allow for specific recommendations…When non-obstetric surgery is planned, the primary obstetric provider should be notified. If that provider is not at the institution where surgery is to be performed, another obstetric care provider with privileges at that institution should be involved.”

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**ACOG / ASA STATEMENT**

“Surgery should be done at an institution with neonatal and pediatric services; an obstetric provider with cesarean delivery privileges should be readily available, and a qualified individual should be readily available to interpret the fetal heart rate.”

Obstet Gynecol 2011;117:420 (reaffirmed 2013)

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**Controversy #5. What is the best way to manage the EXIT (ex-utero intrapartum treatment) procedure?**

Case: A healthy woman at term has a fetus with a large neck mass found on ultrasound. The mass is compromising its airway and intubation will be required immediately after delivery.
EXIT PROCEDURE
(Ex Utero Intrapartum Treatment)

1. Maternal GETA is induced and then maintained with 2 MAC volatile agent to provide uterine relaxation and fetal anesthesia.
2. After uterine incision and hemostasis, only the head and arm are delivered. A pulse oximeter is placed, IM relaxant and narcotic are given to the fetus. The placenta remains intact.
3. The trachea is intubated or tracheostomy is performed. Surfactant may be administered.
4. Once the airway is secure, volatile agent is discontinued to regain uterine tone, delivery is completed and oxytocics are begun.

Anesthesiology 2011;114:1446

EXIT PROCEDURE

Case report: Fetus with an oral teratoma required EXIT procedure at 25 weeks due to preterm labor. Under general anesthesia, fetus underwent bronchoscopy and tracheostomy while on placental circulation. Delivery and resection followed. The mother was discharged after 4 days.

Obstet Gynecol 2012;119:466
FETAL SURGERY

Fetal / prenatal repair of myelomeningocele may be preferable to neonatal repair. Issues to consider for open fetal surgeries:
• Ethical considerations
• Inducing profound uterine relaxation
• Vigilance for maternal and fetal blood loss
• Fetal monitoring and possible resuscitation
• Postoperative tocolysis and analgesia

Anesthesiology 2013; 118: 1211

FETAL SURGERY

Fetal endoscopic tracheal occlusion is used to treat severe CDH.
• ↑ survival: 54% vs. 5%
• Resulted in improvement in fetal lung size and pulmonary vascularity
• Response in lung growth 4 weeks after occlusion is predictive of neonatal survival.

Obstet Gynecol 2012;119:93

FETAL SURGERY

Using a maternal-fetal sheep model, anesthesia with high-dose desflurane was compared to low-dose des + propofol / remifentanil.
• High-dose des → more maternal hypotension, fetal acidosis, and ↓ uterine blood flow.
• Even with maternal normotension, UBF still declined and fetal acidosis persisted.
• No difference in fetal cardiac function.

Anesthesiology 2013; 118: 796

Controversy #6. Is there a “best” anesthetic during pregnancy to protect the fetal brain from neurotoxicity?

Do anesthetic drugs cause developing neurons to commit suicide (apoptosis)?

Case: A healthy well-educated woman requires emergency appendectomy at 26 weeks gestation. Based on information she obtained on the internet, she questions you about the effect of your general anesthetics on her fetus’ developing brain.
**ANIMAL STUDIES**

In a simulated clinical scenario:
- 7-day old rats (0-6 months in humans) received 6 hours of general anesthesia: midazolam, nitrous oxide, isoflurane.
- Postop, the animals had memory / learning impairments, apoptotic neurodegeneration, and hippocampal synaptic function deficits.

J Neuroscience 2003;23:876

**ANIMAL STUDIES**

Are the adverse effects attributable to the direct effects of anesthetics, or are they the result of factors we would not see clinically; eg. high doses over long periods, acidosis, hypoxia, starvation?

Problems extrapolating animals to humans:
- Inter-species differences
- Simulating normal O.R. conditions
- Adequate monitoring

**PRIMATE STUDIES**

Are non-human primates (versus rats) also susceptible to anesthetic toxicity?
- Rhesus macaques received 5 hours of 1 MAC isoflurane while ventilated.
- There was a 13-fold increase in neuro-apoptosis in exposed animals, largely concentrated in the cerebral cortex.

Anesthesiology 2010;112:834

**ANIMAL STUDIES**

Editorial view:
“...it is not clear which anesthetic technique might be least toxic, nor has any general anesthetic been convincingly shown to be more toxic…certainly non-urgent surgery should continue to be postponed until after pregnancy. Considerations should be made to using regional anesthesia when possible.”

Anesthesiology 2011;114:479

**IS REGIONAL BETTER?**

A retrospective study of adnexal mass surgery during pregnancy compared 137 women having general anesthesia with 71 having regional anesthesia.

The overall incidence of preterm labor was higher in the surgical than control groups. Regional had significantly higher incidence (30%) than the general anesthetic group (6%).


**ANESTHETIC TOXICITY**

What about exposure of the fetus in-utero?
- Non-obstetric surgery and fetal interventions often use GETA at high concentrations for longer than C/S, and all lipophilic anesthetics can be measured in the fetal brain.
- 2nd trimester: rapid fetal brain development
- Animal exposure → neuronal cell death and behavioral abnormalities; same for humans?

ANESTHETIC TOXICITY
Do children exposed to anesthesia in infancy have deficits in school performance?
• Mean composite scores on academic achievement tests did not appear different.
• However, 14% scored below 5th %ile, even when other CNS problems or risk factors during infancy could be ruled out.
• There was a negative association between duration of anesthesia and test scores (longer = lower).
  Anesthesiology 2012;117:494

ANESTHETIC TOXICITY
Analysis comparing 321 children age 10 who were exposed to anesthesia under age 3:
• Battery of neuro-psych tests administered.
  • ↑ language disability (RR 1.87)
  • ↑ abstract reasoning deficits (RR 1.69)
  • Disability in language and cognition (RR 2.41)
  • Risks persisted even with only 1 exposure.
  Pediatrics 2012;130:476

ANESTHETIC TOXICITY
Do children age 12 who were exposed to GA for minor procedures during infancy show differences in academic achievement from those who were not exposed? Yes.
• No difference in standardized test scores
• Diagnosed learning disability was 15% vs 4% with an OR 4.5
  Anesth Analg 2013; 117: 1419

ANESTHETIC TOXICITY
Expert report and consensus statement:
1. Pre-clinical studies indicate general anesthetics are modulators of neuronal development and function.
2. Need clinical studies (vs. animal models)
3. Need strategies to avoid or limit brain injury in pediatric and geriatric patients; GA effects are not entirely reversible.
  Br J Anaesth 2013; 111: 143

ANESTHETIC TOXICITY
1. Non-primate models indicate ↑ programmed cell death (apoptosis) after exposure to general anesthetics.
2. Primate models confirm + associated neurocognitive deficits.
3. Retrospective clinical data is inconclusive and prospective studies are ongoing → changes to pediatric anesthesia practice are premature.
  Br J Anaesth 2013; 110: i53

ANESTHETIC TOXICITY
How do we interpret observational studies?
• What is the population receiving anesthesia?
• Who is actually included in the analysis?
• What is the definition of anesthetic exposure?
• What is the comparison group?
• What is the outcome measure?
• How are the data analyzed?
• What is the clinical relevance?
  Anesthesiology 2012;117:459
ANESTHETIC TOXICITY
Summary of what we know:
• Single anesthetics may not have an effect.
• Repeated exposures do show an effect.
• Effects persist after adjustments for co-morbidities.
• Learning (reasoning), speech and language are affected but not behavior.
• Observational studies are prone to bias, confounding, etc. but RCTs for this question are not possible or ethical. Prospective trials are ongoing.
AAP 2012 Nat’l Conference / Medscape, 10/25/12

ANESTHETIC MANAGEMENT OF THE PREGNANT SURGICAL PATIENT

COMMON SURGERIES
The most common indications for surgery unrelated to pregnancy:
1. Appendicitis, 1:2000 pregnancies
2. Cholecystitis, 1:6000 pregnancies
3. Maternal trauma
4. Maternal malignancies

PREOP ASSESSMENT
• Is my patient pregnant? Document LMP on record. Offer pregnancy testing.
• Operate during second trimester if possible. Less risk of early spontaneous miscarriage. Theoretical risks of teratogenicity are avoided. 3rd trimester ↓ introp visibility, ↑ preterm labor
• Reassure her about risks to fetus or pregnancy.
• Educate her about uterine displacement, symptoms of preterm labor.

PREOP MEDICATIONS
• Sedation may be beneficial. Narcotics Benzodiazepines
• Consider aspiration prophylaxis. Antacid Metoclopramide H-2 receptor blocker

INDOMETHACIN TOCOLYSIS
Usual dose 50 mg loading, 25 mg q 6 hours PO or PR
Implications Maternal Fetal
Platelet function Necrotizing enterocolitis
GI symptoms Oligohydramnios
Renal insufficiency Closure of fetal ductus
CHOICE OF ANESTHETIC

• There is no evidence in humans (yet) that any drug or anesthetic technique is dangerous to the fetus (stay tuned on the neurotoxicity issue).
• Choose the safest anesthetic for the mother’s condition, and modify for the physiologic changes of pregnancy.
• Avoid hypoxia and hypotension!

GENERAL ANESTHESIA

• Full pre-oxygenation / denitrogenation
• Rapid sequence induction, smaller ETT
• First trimester: use tried and true drugs
• Nitrous oxide versus FIO₂
• ET CO₂ 28-32; avoid hyperventilation
• Inhalational agents < 2.0 MAC
• Slow or no reversal of relaxants
• Compression stockings

NEURAXIAL ANESTHESIA

• Advantage of minimizing drug exposure
  First trimester
  Fetal monitoring
• Prevent hypotension
  Adequate fluid replacement
  Uterine displacement
• Decrease neuraxial local anesthetic dose by 30%.
• Choose ephedrine vs. phenylephrine based on maternal heart rate.
• Utilize for postoperative pain control.

POSTOPERATIVE CARE

• Continue monitoring fetal heart rate and uterine activity. Provide L&D nursing expertise.
• Maintain maternal oxygenation and LUD.
• Notify Pediatrics if the fetus is a viable gestational age > 24 weeks.
• Use neuraxial narcotics or regional blocks for pain management if possible to encourage early ambulation.
• Use thromboembolism prophylaxis.

SPECIFIC SURGICAL SITUATIONS
**TRAUMA**

- A leading cause of maternal death, especially MVA without use of seat belts.
- Fetal loss is due to maternal death or placental abruption.
- Need early ultrasound in E.R. to determine gestational age and viability.
- Perform all necessary diagnostic tests on the mother with shielding as necessary.
- Maternal ↑ blood volume may mask blood loss.

Obstet Gynecol 2009;114:147

**TRAUMA**

What are the risks of radiation exposure?

- ACOG has stated: “no single diagnostic x-ray procedure results in radiation exposure to a degree that would threaten the well-being of the developing fetus.”
- Teratogenic risks are not increased with < 5 rad exposure (eg. a head CT < 1 rad).
- Ultrasound and MRI are safe alternatives.

Anesth Analg 2010;110:863

**TRAUMA**

Indications for emergent C/S:

- Stable mother, viable fetus in distress
- Uterine rupture
- Gravid uterus interfering with repairs
- Mother unsalvageable, fetus viable

If the fetus is previable or dead, focus on optimizing the mother. She will tolerate vaginal delivery at a later time better than an emergent laparotomy.

**NEUROSURGERY**

- Intracranial aneurysms or AVM may require repair in this age group.
- Usual anesthetic techniques can be used.
- Fetal monitoring is remote from the field and may be beneficial in some cases, eg. aggressive diuresis, hyperventilation, bleeding and fluid shifts.

Anesth Analg 2008;107:193

**NEUROSURGERY**

Successful endovascular treatment of acutely ruptured intracranial aneurysms in pregnancy:

- 32 wks gestation with HA and vomiting. CT and MRI show SAH and aneurysm: C/S → angio → embolization with coils.
- 22 wks gestation with HA, vomiting, LOC. CT shows SAH: GETA → angio → occlusion with coils using fetal shielding → SVD at term.

Am J Obstet Gynecol 2001;185:1261

**CARDIAC BYPASS**

Pregnant patients who had cardiopulmonary bypass procedures were reviewed:

- Preterm birth or IUFD were associated with emergent procedures, maternal co-morbidities, and early gestational age.
- Recommendations: normothermic, high-flow bypass, postponing until 2nd trimester.
- Elective delivery before CPB should be considered if the fetus is viable.

Ann Thorac Surg 2011;91:1191
### GOALS DURING CPB

- High pump flows (>2.5 L/min/m²)
- High MAP > 65 mmHg
- Hematocrit > 28%
- Normothermic CPB (limit < 32°C)
- Pulsatile flow?
- Optimize CO₂, acid-base, glucose
- Use continuous fetal HR monitoring

### LAPAROSCOPY

Symptomatic cholelithiasis during pregnancy is not rare, but medical versus surgical management has been controversial. Case control study:

- 38% of medical patients had relapses. Each relapse accounted for additional 5 inpatient days.
- Compared to medical management, surgery patients had less preterm labor, fewer premature deliveries, and fewer days in-hospital.


### LAPAROSCOPY

Is laparoscopy better for fetal outcome than an open procedure?

- There are no outcome differences between laparoscopy and laparotomy in maternal complications or fetal outcome.
- Laparoscopy patients (the mothers) had longer operative times but shorter hospital stays, less parenteral narcotics, and earlier resumption of a regular diet.

Clin Obstet Gynecol 2009;52:557

### LAPAROSCOPY

Following laparoscopy (n=2181) or laparotomy (n=1522) performed between the 4th and 20th weeks of gestation, there were no differences in:

- Infant survival to one year
- Rate of fetal malformations
- Birth weight
- Gestational duration
- Growth restriction

There was an increased risk of low birth weight < 2500 gm, delivery before 37 weeks, and growth restriction when comparing the operated groups to the general population.

Am J Obstet Gynecol 1997; 177:673

### GOALS FOR LAPAROSCOPY

- Consider an open technique to enter abdomen.
- Maintain normal end-tidal CO₂, consider blood gas monitoring to rule out respiratory acidosis.
- Keep inflation pressure < 15 mmHg.
- Laparoscopic techniques can be used in any trimester of pregnancy.
- Maintain uterine displacement and monitor the fetus if feasible.
- Use compression devices for DVT prophylaxis.


### SUMMARY

Approach the pregnant surgical patient with respect, rather than apprehension.

Recognize her fears related to her pregnancy.

Doing what is best for the mother will almost always be best for the fetus and the outcome of the pregnancy.