PREOPERATIVE EVALUATION OF THE PEDIATRIC PATIENT



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THE OBJECTIVES

Discuss Preoperative Evaluation of the Pediatric Surgical Patient
Discuss the Complications of Anesthesia in children with Upper Respiratory Infections
Discuss the Safety of anesthetic agents in the pediatric population



Is This Necessary?

- Helps streamline the day of surgery
- Helps eliminate the preventable causes for cancellation of surgery
- Helps to optimize medical management
- Helps familiarize child and parent to the hospital
- Reduce the percentage of cancellations and OR delays
- Enhance the utilization of OR time



Preoperative Evaluation Objectives

Obtaining a medical history
Performing a physical examination
Review of systems
Family history
NPO status
Appropriate laboratory tests



Preoperative Evaluation Maternal Health and Children

Maternal Health	Implications to Children
Diabetes Mellitus	Neonatal hypoglycemia, macrosomia, respiratory distress, cardiac anomalies, vertebral anomalies
Pre eclampsia	IUGR, preterm delivery, high incidence of meconium at birth
Alcohol Abuse	Mental retardation, congenital heart disease, cleft palate, skeletal anomalies
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Preoperative Evaluation An Outline

Birth History:- Full term or pre term baby Determine post conceptual age History of previous hospitalizations and surgical procedures Concurrent medical illnesses History of recent upper respiratory infection Allergies



Birth History The Former Preterm Infant *Problems*

- Pulmonary:- lung immaturity, bronchopulmonary dysplasia, elevated pulmonary vascular resistance
- Airway:- tracheomalacia, subglottic stenosis
- Multiple medications
- Apnea and bradycardia



Postoperative Apnea and Bradycardia

Factors affecting include:-

- Prematurity
- Post conceptual age less than 60 weeks
- Anemia

 History of Apnea and bradycardia, on home apnea and bradycardia monitor



Preoperative evaluation Physical Examination

- Here, importance is given to the evaluation of:-
- Airway
- Dentition
- Heart murmurs
- Neurological status and deficits















Preoperative evaluation Physical Examination

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Preoperative evaluation Miscellaneous workups

Laboratory studies
Radiographs
CT scans/MRI
Cardiac/pulmonary/hematology consults where necessary



Preoperative Evaluation Family History

Aspects of Family History with increased anesthetic risks include Unusual reactions to surgery or anesthesia Malignant Hyperthermia Sickle cell disease Thalassemia Atypical Pseudo cholinesterase Neuromuscular disorders

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Preoperative Evaluation NPO Guidelines

Clear liquids
Breast milk
Infant formula
Nonhuman Milk
Solids

2 hours 4 hours 6 hours 6 hours 8 hours



"My Child Has A Running Nose" URIs and Anesthesia





Upper Respiratory Infections

URIs affect the apparatus of the upper airways Frequently occurring URIs include:

- The Common Cold
- Pharyngitis
- Laryngitis
- Croup
- Epiglottitis
- Sinusitis



"My Child Has A Running Nose" URIs and Anesthesia

URI	Causative Agents
Common Cold	Rhino, Corona, RSV, Para influenza, Influenza, Adenoviruses
Pharyngitis	Viral:- "cold Virus", Coxsackie, Herpes simples, E-B virus.
	Bacterial:- Streptococcus
Laryngitis	Viral:-Influenza, Para influenza, Rhino, Adeno, RSV
	Bacterial:-Streptococcus
	Fungal:-Candida Albicans

"My Child Has A Running Nose" URIs and Anesthesia

URI	Causative Agents
Croup	Para influenza viruses type1,2,3;, Influenza and RSV
Epiglottitis	H. Influenza type B
Sinusitis	S.pneumonia, H influenza, S.aureus, S.pyogenes, fungal infections

Symptoms and signs of URI

Rhinorrhea, nasal congestion Cough with or without expectoration Sneezing Malaise and decreased activity Sore Throat Head ache Fever



URIs and Anesthesia

One URI can progress to another type
Signs and symptoms are similar
Difficult to distinguish these infections from one another
Most children develop 6-10 "colds" a year
Less frequent after age 6



URIs and Anesthesia

Differential diagnosis of URI:Vasomotor rhinitis
Allergic rhinitis



Risk factors for Perioperative Complications

- Copious secretions
- ETT in children <3yrs</p>
- History of prematurity
- Nasal congestion
- Parental smoking
- History of Reactive airway disease
- Surgery involving the airway
- Productive Sputum

Tait et al, Anesthesiology, V 95, No 2, Aug 2000.



URI and Pulmonary Function

Pulmonary Function following URI:-Reduced FEV1

- Decreased mucociliary clearance
- Increased airway reactivity
- Increased closing volumes
- Compromised diffusion capacity
- Increased intrapulmonary shunting and worsening ventilation perfusion mismatch



Causes for airway hyper-reactivity

Respiratory epithelial cell damage
Activation of the sub epithelial receptors
Release of Mast cells, substance P
Bronchial smooth muscle stimulation



URI:-Perioperative adverse events

- Laryngospasm
- Airway obstruction
- Croup
- Hypoxemia
- Bronchospasm
- Stridor
- Breath holding



Laryngospasm and URI Schreiner, Anesthesiology vol. 85(3), Sept. '96

Laryngospasm is 2 times more common
More likely to be in the younger age group
More likely to be scheduled for airway surgery

Anesthesia supervised by a less experienced anesthesiologist



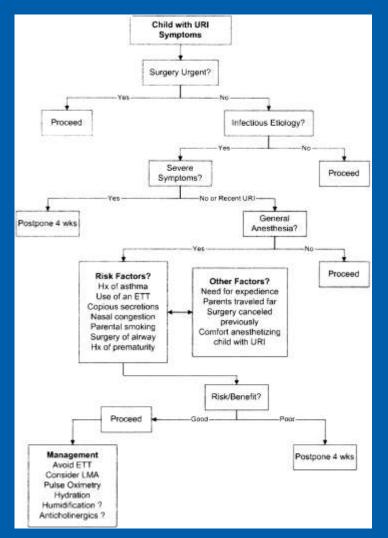
Cardiac surgery and URI Malviya et al (Anesthesiology 98, March '03) Intraoperatively:-Increased airway complications Postoperatively an increased incidence of :-Respiratory complications Bacterial infections Atelectasis Extended stay in the ICU



My Child has a LITTLE cough Should surgery be cancelled?



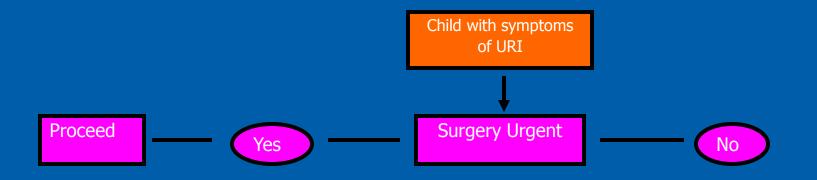
Figure 1. Suggested algorithm for the assessment and anesthetic management of the child with an upper respiratory infection.

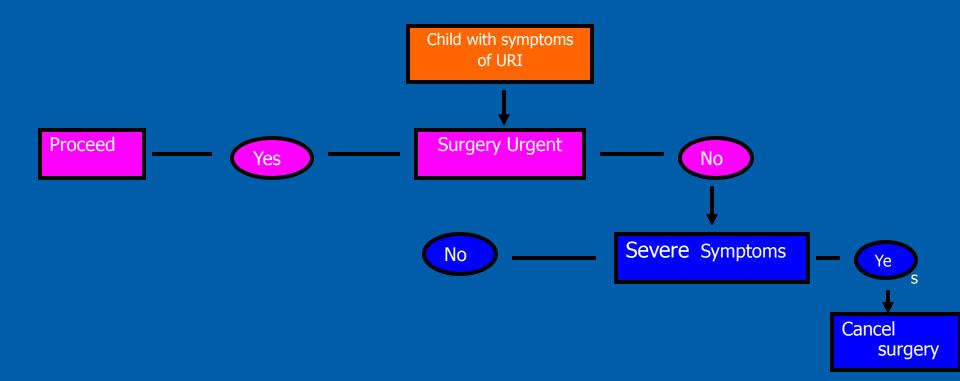


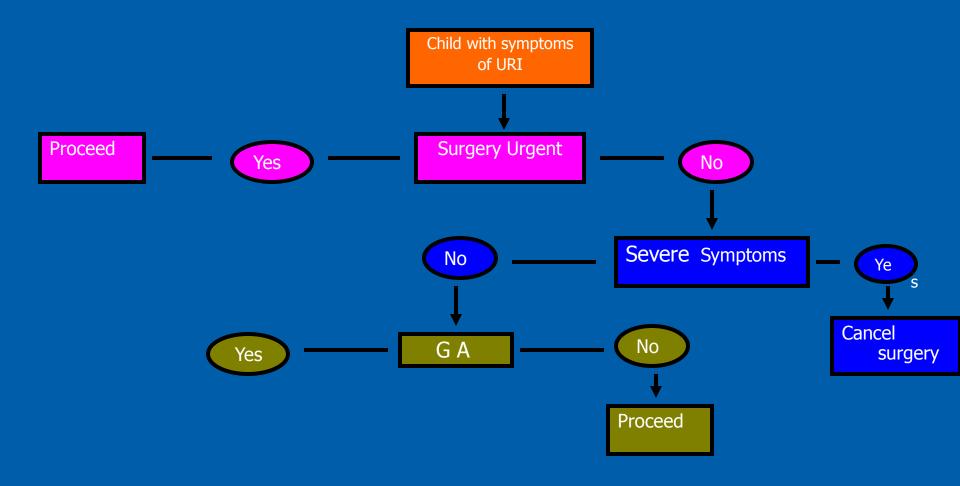
Tait A R, Malviya S Anesth Analg 2005;100:59-65

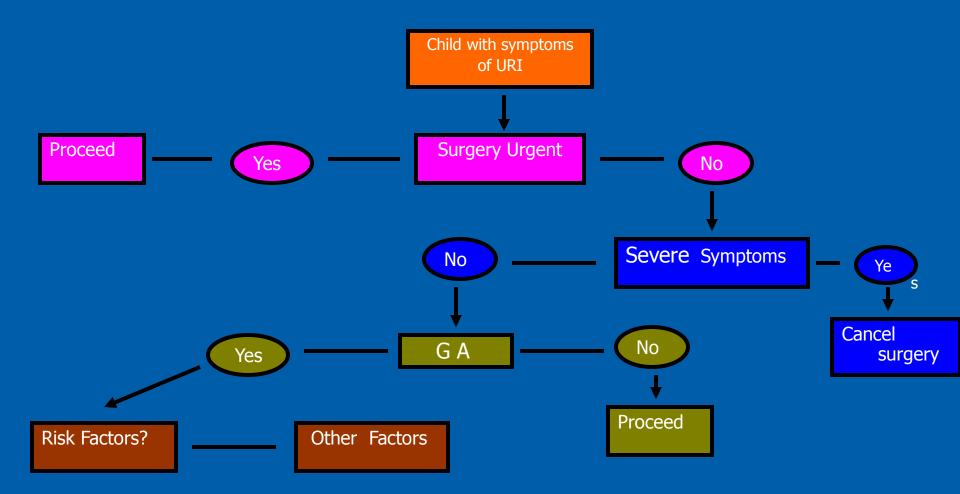
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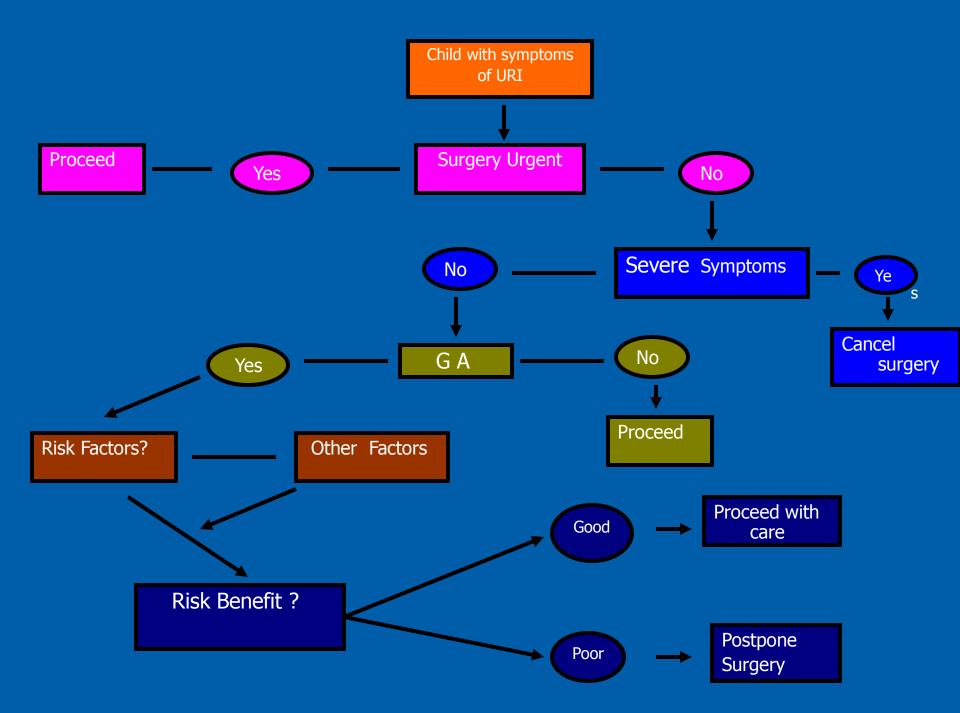
Child with symptoms of URI











In Conclusion---Upper respiratory infection and anesthesia

What seems like a little cough to me may not mean the same to my colleagues. And, when this cough is associated with other signs of infection like fever, sputum which is greenish-yellow, wheezes, lethargy, ----- it most definitely is not a Little Cough.



Is it safe for my baby to have anesthesia ?





Pain And Its Effects In The Human Neonate And Fetus

Anand KJ, Hickey PR, N Engl J Med, 1987
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Cutaneous Sensory Perception
MYELINATION
Nerve Tracts in the Spinal Internal Capsule Cord and Brain Stem Corona Radiata
CORTICAL MATURATION
Neuronal Migration
Dendritic Arborization
Synaptogenesis with Thalamocortical Fibres
EEG PATTERNS
Intermittent
Pattern I EEG Synchronous
Pottern 2 EEG
Pattern 3 EEG
Pattern 4 EEG
Cortical Evoked Potentials

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Randomised Trial Of Fentanyl Anaesthesia In Preterm Babies Undergoing Surgery

Anand, Sippell and Aynsley-Green Lancet 1987

Premature neonates for ligation of PDA

- Randomized
 - Nitrous oxide and curare with fentanyl (10 µg/kg)
 - Nitrous oxide and curare
- No fentanyl group
 - Metabolic acidosis
 - Hyperglycemia
 - Post operative
 - Ventilator requirements
 - Bradycardia
 - Intraventricular hemorrhages
 - hypotension



What We Think That We Know Now

- Based on Anand's early work, Infants have a profound physiological response to surgical trauma
- This response has an impact on outcome
- Poor outcomes can be reduced by the judicious use of anesthetics
- Bottom line: Infants have pain



Commonly used Anesthetic Agents

Table 1. Receptor Activity of Commonly Used Anesthetic Agents^a

Anesthetic agent	NMDA antagonist	GABA-mimetic	μ -Opioid agonis	
Volatile anesthetics				
Halothane	-/0	+++	0	
Isoflurane	-/0	+++	0	
Desflurane	-/0	+++	0 0 0	
Enflurane	-/0	+++	0	
Sevoflurane	-/0	+++	0	
Injectable anesthetics				
Propofol	0	+++	0	
Barbiturates	0	+++	0 0 0 0	
Etomidate	0	+++	0	
Benzodiazepines	0	+++	0	
Ketamine	1000	-/0	0	
Medical gases				
Nitrous oxide		+++	0	
Opioid analgesics				
Morphine	-/0	0	+++	
Methadone		0	+++	
Meperidine	-/0	0	+++	
Fentanyl	-/0	0	+++	
Other sedative hypnotics				
Chloral hydrate		+++	0	
Trichloroethanol		+++	0 0 0	
Ethanol		+++	0	

Key: (---); strong antagonism within the clinically relevant range based on available in vitro data; (+++); strong potentiation within the clinically relevant range based on available in vitro data; (+/0), little antagonism within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (+/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentiation within the clinically relevant range based on available in vitro data; (-/0), little potentint on vitro data; (-/0), little potentiation witron data; (-/

NMDA = N-methyl-p-aspartate; GABA = y-aminobutytic acid.

⁴ Receptor activity as noted in the table does not imply mechanism of action; rather it indicates potential to neurological injury. The table was adapted, in large part, from ref and Harrison (9) and by Dilger (10); for more detail, please refer these review articles.



Blockade of NMDA Receptors and Apoptotic Neurodegeneration in the Developing Brain

Chrysanthy Ikonomidou,* Friederike Bosch, Michael Miksa, Petra Bittigau, Jessica Vöckler, Krikor Dikranian, Tanya I. Tenkova, Vanya Stefovska, Lechoslaw Turski, John W. Olney

 Programmed cell death occurs during normal development
 Blockade of NMDA glutamate receptors produce widespread apoptotic neurodegeneration in developing rat brain

• Suggested that glutamate acting at NMDA receptors controls neuronal survival

• Question: Does this have relevance to human neurodevelopmental disorders?



Early Exposure to Common Anesthetic Agents Causes Widespread Neurodegeneration in the Developing Rat Brain and Persistent Learning Deficits Jevtovic-Todorovic et al, J Neurosci, 2003

- Triple cocktail (P7 rat pups)
 - Midazolam
 - Isoflurane
 - Nitrous oxide
- Histological changes
 - neurodegeneration
- Functional changes
 - Memory impairments
 - Learning impairments



Anesthesiology 2005; 102:866-8

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Of Mice and Men: Should We Extrapolate Rodent Experimental Data to the Care of Human Neonates?

Sulpicio G. Soriano, M.D.,* Kanwaljeet J. S. Anand, M.B.B.S., D.Phil., Cynthia R. Rovnaghi, M.S., Paul R. Hickey, M.D. * Children's Hospital Boston and Harvard Medical School, Boston,



Problems with Rodent Experimental Paradigm

Duration of exposure to drugs

7 rat days \cong 27 human months 6 hour \cong 1 month



Problems with Rodent Experimental Paradigm

Duration of exposure to drugs
Lack of precise physiological monitoring



Physiological Monitoring





Problems with Rodent Experimental Paradigm

Duration of exposure to drugs
 Lack of precise physiological monitoring
 End-organ perfusion
 Varying anesthetic depth
 MAC values

Target-controlled infusions



Problems with Rodent Experimental Paradigm

Duration of exposure to drugs
 Lack of precise physiological monitoring
 Interspecies variation



Problems with Rodent Experimental Paradigm

Duration of exposure to drugs

- Lack of precise physiological monitoring
- Interspecies variation
 - Dose-response
 - Drug metabolism
 - Peak susceptibility



■ 2008 ANESTHESIOLOGY/FAER SESSION: ANESTHESIA AND THE DEVELOPING BRAIN: IMPLICATIONS FOR OBSTETRICS AND PEDIATRICS

Anesthesiology 2009; 110:796-804

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Early Exposure to Anesthesia and Learning Disabilities in a Population-based Birth Cohort

Robert T. Wilder, M.D., Ph.D.,* Randall P. Flick, M.D., M.P.H.,† Juraj Sprung, M.D., Ph.D.,‡ Slavica K. Katusic, M.D., William J. Barbaresi, M.D., Christopher Mickelson, M.D., # Stephen J. Gleich, M.D.,** Darrell R. Schroeder, M.S., Amy L. Weaver, M.S., †† David O. Warner, M.D.,



Early exposure to Anesthesia and Learning Disabilities

- Population based birth cohort study
- Children born between 01/1976 2/1982
- Risk factor was exposure to general anesthesia <4yrs of age (total of 593)
- Tested for learning disabilities
- Higher incidence of LD

in those exposed to 2 or more anesthetics
 in those receiving anesthesia >120mins.



Initiatives by the Pediatric Anesthesia Community

Neonatal Drug Development Initiative Workshops
 Society for Pediatric Anesthesia

 Awareness-educational programs
 Registry and complications database

 American Society of Anesthesiology

 Increased awareness
 Funding investigators

 Multi-center Randomized Control Trial



GAS Study

Multi-national, multi-center collaborative group

- Andrew Davidson, Melbourne, Australia
- Mary Ellen McCann, Boston, USA
- Neil Morton, Glasgow, Scotland
- Randomized controlled equivalence trial
 - Inguinal hernia in infants
 - Spinal (bupivacaine) versus general (sevoflurane) anesthesia
 - Neurodevelopmental assessments at 2 and 5 years

Timeline	2007	2008	2009	2010	2011	2012	2013	2014
Recruit	×	×	×					
Yr 2 assessme nt			×	×	×			
Yr 5 assessme nt						×	× UKH	× ealthCare

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Is it safe for my baby to have anesthesia?

For now:

- Drug induced apoptosis is a valid concern
- Much needed area of research
- Does it affect just preterms or neonates or are adults vulnerable too?
- Is the plasticity of human brain able to overcome these insults
- Until we have more convincing data on this subject it is prudent that we continue with our current level of care



Is it safe for my baby to have anesthesia ?

Remember:1.Majority have been animal studies2. Questionable intraop monitoring3. Limited data in primates





