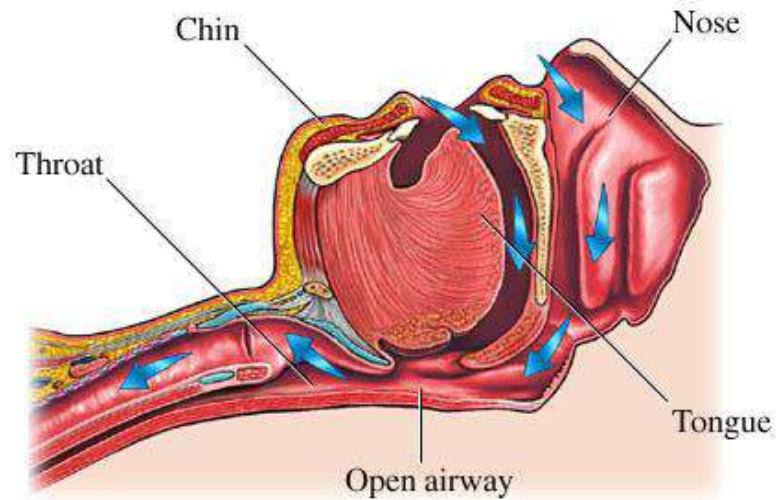




Airway Management In Difficult Situation



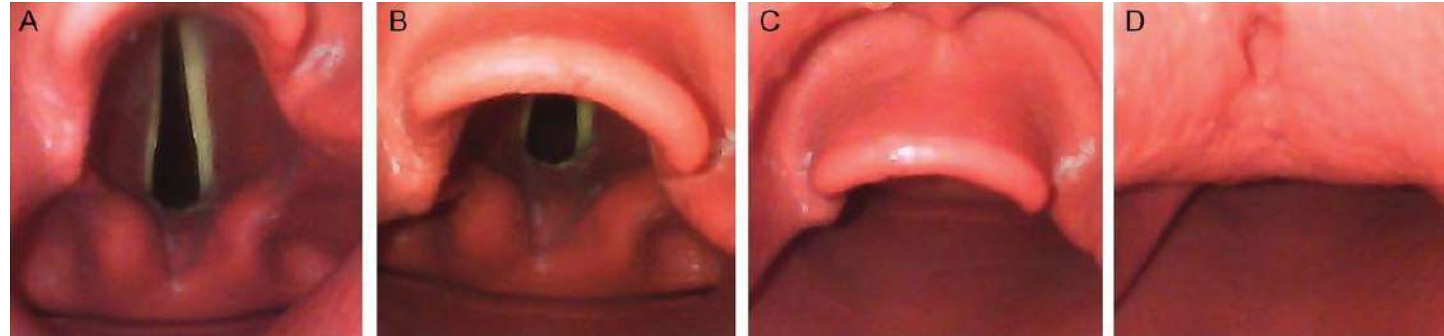
Apirak Thewaritrueangsri, MD
Rawee Jongkongkawutthi, MD

Department of anesthesiology
Naresuan University



What is the problem?

- Uncooperative patient
- Can't see vocal cord
 - Laryngeal view grade $> I$
 - Obscured by Secretion / Blood / Mass
- Seen vocal cord, but can't insert endotracheal tube into vocal cord
 - Can't control tip of ETT to vocal cord
 - Vocal cord edema
- Limited mouth opening or neck movement





Classification Of Difficult Airway



Difficult airway

- The clinical situation in which anticipated or unanticipated difficulty or failure is experienced by a physician trained in anesthesia care



Difficult airway

- The clinical situation in which anticipated or unanticipated difficulty or failure is experienced by a physician trained in anesthesia care



Difficult Facemask Ventilation.

Difficult Laryngoscopy.

Difficult Supraglottic Airway Ventilation.

Difficult or Failed Tracheal Intubation.

Difficult or Failed Invasive Airway.



Classification

- **Difficult Facemask Ventilation**
- Difficult Supraglottic Airway Ventilation.
- Difficult laryngoscopy
- Difficult or Failed Tracheal Intubation
- Difficult or Failed Invasive Airway

Inadequate mask seal

Excessive gas leak

Excessive resistance to the ingress or egress of gas.



Classification

- Difficult Facemask Ventilation
- **Difficult Supraglottic Airway Ventilation.**
- Difficult laryngoscopy
- Difficult or Failed Tracheal Intubation
- Difficult or Failed Invasive Airway

Difficult supraglottic airway placement

Supraglottic airway placement requiring multiple attempts

Inadequate supraglottic airway seal

Excessive gas leak

Excessive resistance to the ingress or egress of gas.



Classification

- Difficult Facemask Ventilation
- Difficult Supraglottic Airway Ventilation.
- **Difficult laryngoscopy**
- Difficult or Failed Tracheal Intubation
- Difficult or Failed Invasive Airway

It is not possible to visualize any portion of the vocal cords after multiple attempts at laryngoscopy.



Classification

- Difficult Facemask Ventilation
- Difficult Supraglottic Airway Ventilation.
- Difficult laryngoscopy
- **Difficult or Failed Tracheal Intubation**
- Difficult or Failed Invasive Airway

Tracheal intubation requires multiple attempts or tracheal intubation fails after multiple attempts.



Classification

- Difficult Facemask Ventilation
- Difficult Supraglottic Airway Ventilation.
- Difficult laryngoscopy
- Difficult or Failed Tracheal Intubation
- **Difficult or Failed Invasive Airway**

Anatomic features or abnormalities reducing or preventing the likelihood of successfully placing an airway into the trachea through the front of the neck.



Difficult airway

- A quick assessment of congenital or acquired *anatomic defects*

Facial, **H**ead, or **N**eck trauma,
Oral bleeding, **R**egurgitated gastric contents,
Frothing of the mouth



Neck mobility, **B**eard, **O**besse, **N**o teeth , **E**lderly,
Sleep apnea/snoring, **R**estricted mouth opening, **O**bstruction,
Distorted airway, **S**tiff lungs or c-spine surgery,
Mass, **T**hyromental distance



Stanford Medicine 25





Guidelines for Management of the Difficult Airway



2022 American Society of Anesthesiologists Practice Guidelines for Management of the Difficult Airway*

Jeffrey L. Apfelbaum, M.D., Carin A. Hagberg, M.D.,
Richard T. Connis, Ph.D., Basem B. Abdelmalak, M.D.,
Madhulika Agarkar, M.P.H., Richard P. Dutton, M.D.,
John E. Fiadjoe, M.D., Robert Greif, M.D.,
P. Allan Klock, Jr., M.D., David Mercier, M.D.,
Sheila N. Myatra, M.D., Ellen P. O'Sullivan, M.D.,
William H. Rosenblatt, M.D.,
Massimiliano Sorbello, M.D.,
Avery Tung, M.D.

ANESTHESIOLOGY 2022; 136:31–81

ABSTRACT

The American Society of Anesthesiologists; All India Difficult Airway Association; European Airway Management Society; European Society of Anaesthesiology and Intensive Care; Italian Society of Anesthesiology, Analgesia, Resuscitation and Intensive Care; Learning, Teaching and Investigation Difficult Airway Group; Society for Airway Management; Society for Ambulatory Anesthesia; Society for Head and Neck Anesthesia; Society for Pediatric Anesthesia; Society of Critical Care Anesthesiologists; and the Trauma Anesthesiology Society present an updated report of the Practice Guidelines for Management of the Difficult Airway.

(Anesthesiology 2022; 136:31–81)

HIGHLIGHTS BOX

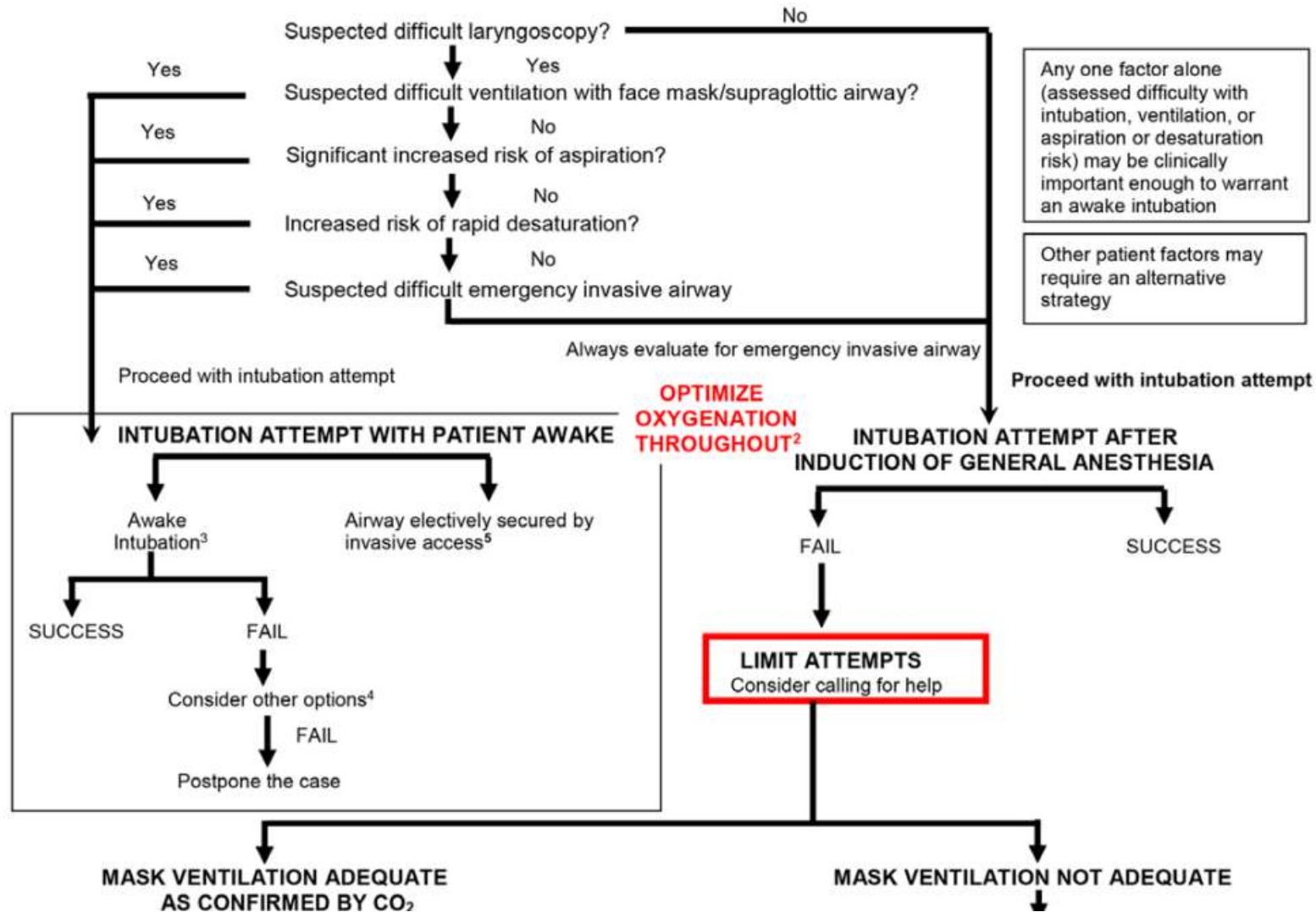
These updated guidelines:

- Replace the “Practice Guidelines for Management of the Difficult Airway: A Report by the American Society of Anesthesiologists Task Force on Management of the Difficult Airway,” adopted by the American Society of Anesthesiologists in 2012 and published in 2013.¹
- Specifically address difficult airway management. The guidelines do not address education, training, or certification requirements for



ASA DIFFICULT AIRWAY ALGORITHM: ADULT PATIENTS

Pre-Intubation: Before attempting intubation, choose between either an awake or post-induction airway strategy. Choice of strategy and technique should be made by the clinician managing the airway.¹



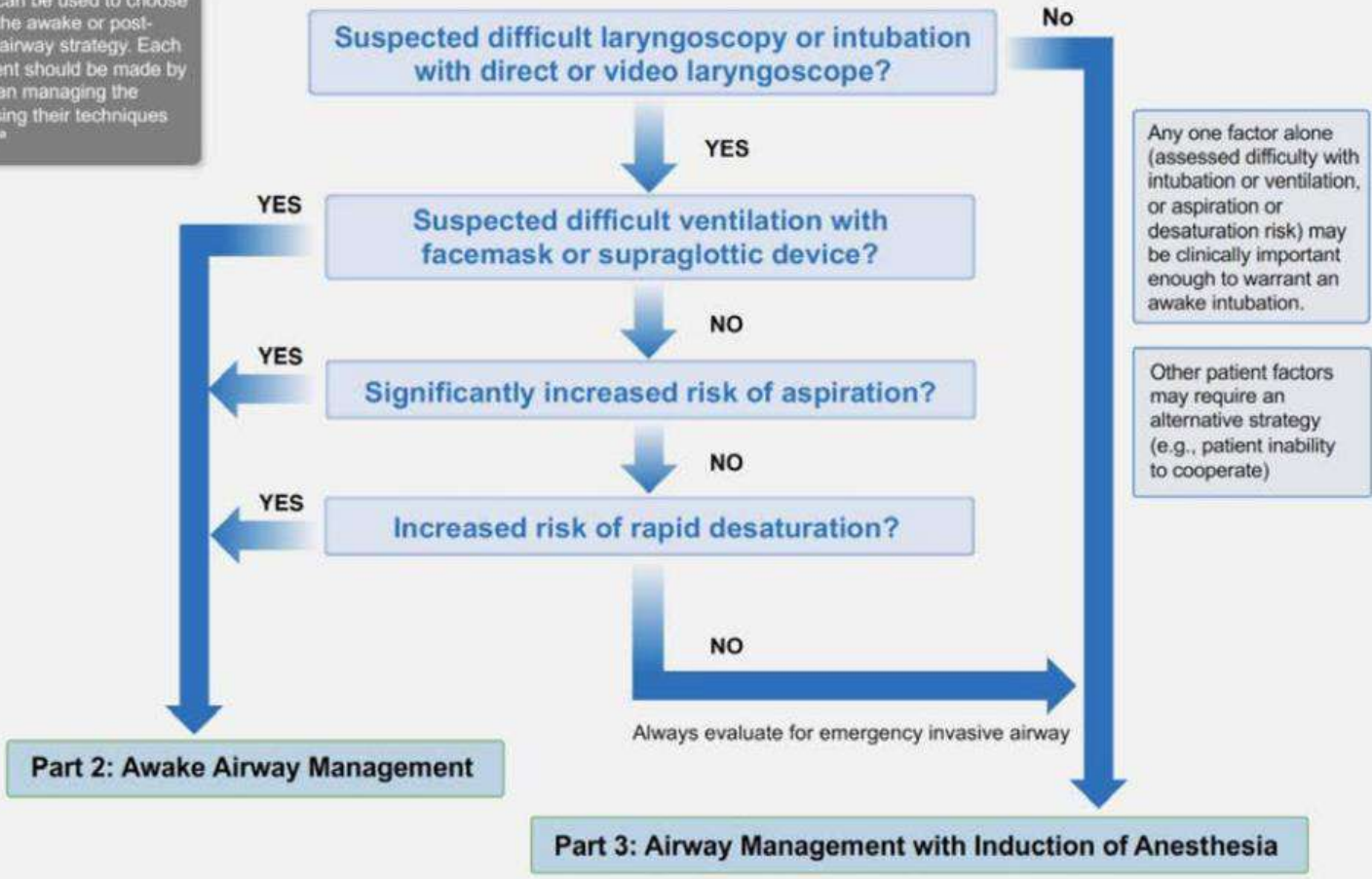
Guidelines for Management of the Difficult Airway



DIFFICULT AIRWAY INFOGRAPHIC: ADULT PATIENTS

Part 1: Pre-Airway Management Decision Making Tool (planning)

This tool can be used to choose between the awake or post-induction airway strategy. Each assessment should be made by the clinician managing the airway, using their techniques of choice.⁹



Guidelines for Management of the Difficult Airway



Preoxygenation Technique

Rises in the alveolar O₂ fraction (FAO₂), reductions in the alveolar nitrogen fraction (FAN₂)

- Face mask
 - Deep Breathing
 - Rapid Breathing at FiO₂=1.0
 - Four Vital Capacities Method
- Transnasal Humidified Rapid Insufflation Ventilator Exchange (THRIVE)
- Oxygen cannula



Preoxygenation Technique

- **Face mask**
 - Deep Breathing
 - Rapid Breathing at $FiO_2 = 1.0$
 - Four Vital Capacities Method
 - Transnasal Humidified Rapid Insufflation Ventilator Exchange (THRIVE)
- Oxygen cannula

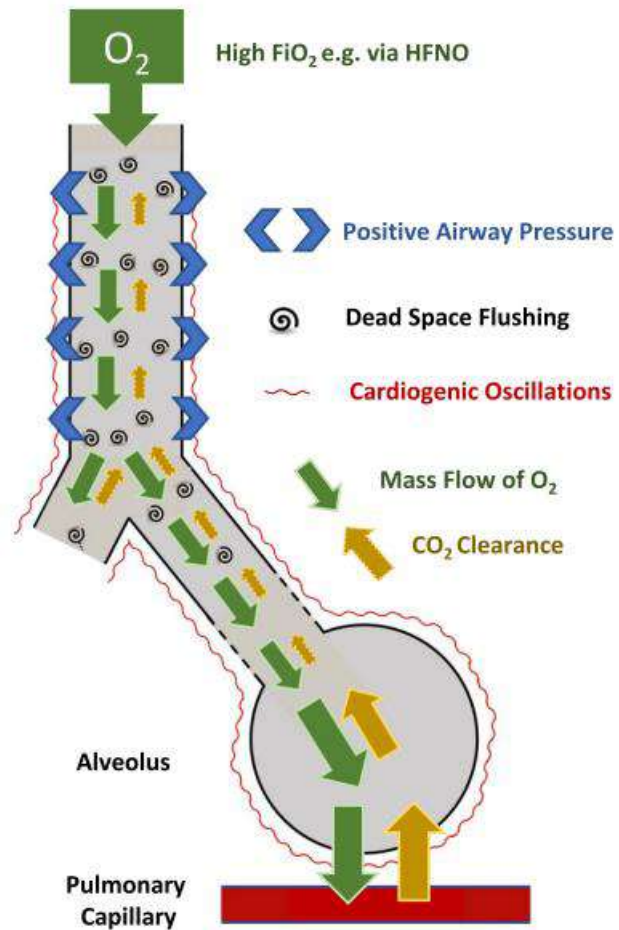


Preoxygenation Technique

- Face mask
 - Deep Breathing
 - Rapid Breathing at $FiO_2 = 1.0$
 - Four Vital Capacities Method
 - Transnasal Humidified Rapid Insufflation Ventilator Exchange (THRIVE)
- Oxygen cannula



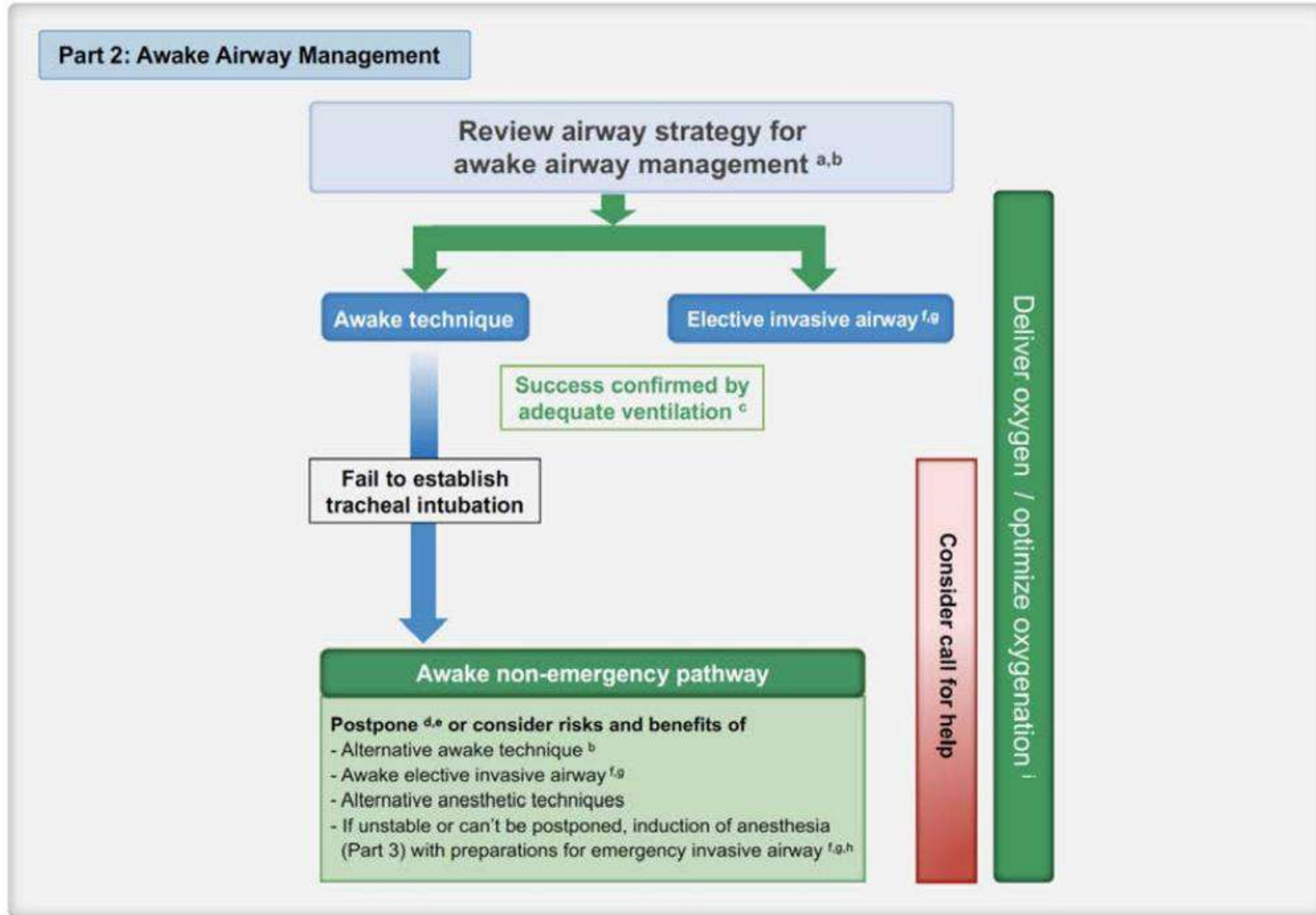
Transnasal Humidified Rapid Insufflation Ventilator Exchange (THRIVE)





Preoxygenation Technique

- Face mask
 - Deep Breathing
 - Rapid Breathing at $FiO_2=1.0$
 - Four Vital Capacities Method
- Transnasal Humidified Rapid Insufflation Ventilator Exchange (THRIVE)
- Oxygen cannula

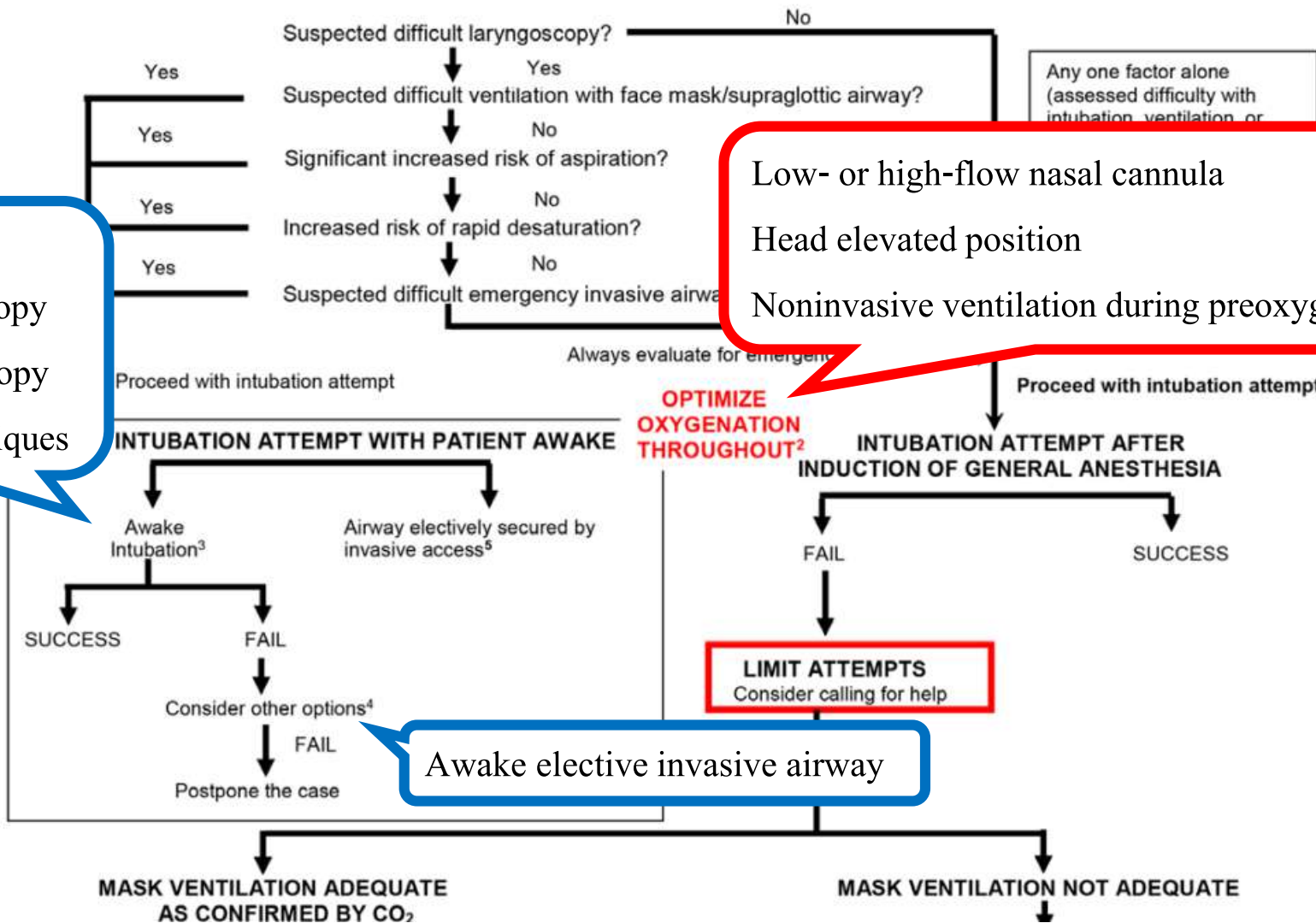


Guidelines for Management of the Difficult Airway



ASA DIFFICULT AIRWAY ALGORITHM: ADULT PATIENTS

Pre-Intubation: Before attempting intubation, choose between either an awake or post-induction airway strategy. Choice of strategy and technique should be made by the clinician managing the airway.¹



Guidelines

- Low- or high-flow nasal cannula
- Head elevated position
- Noninvasive ventilation during preoxygenation

- Fiberoptic
- Video laryngoscopy
- Direct laryngoscopy
- Combined techniques

Management of the Difficult Airway

Awake elective invasive airway

LIMIT ATTEMPTS
Consider calling for help



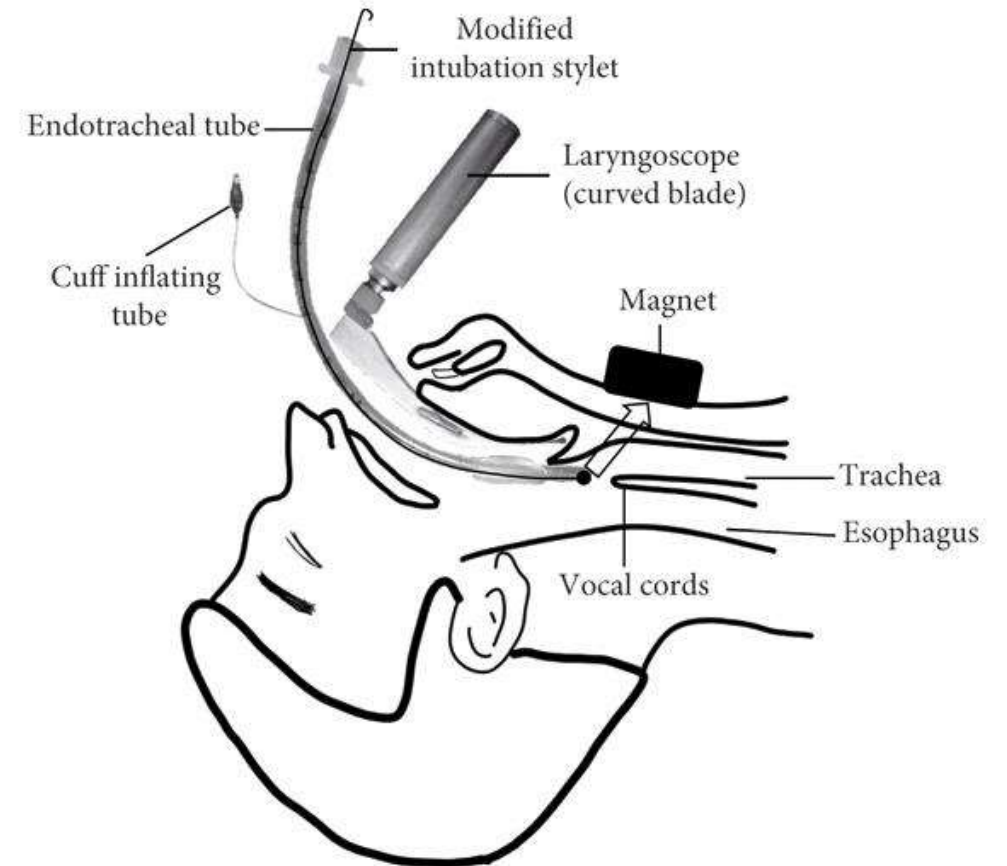
Establish secure airway

- Intubating stylets
- External laryngeal manipulation
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- Intubating supraglottic airway
- Combined techniques



Establish secure airway

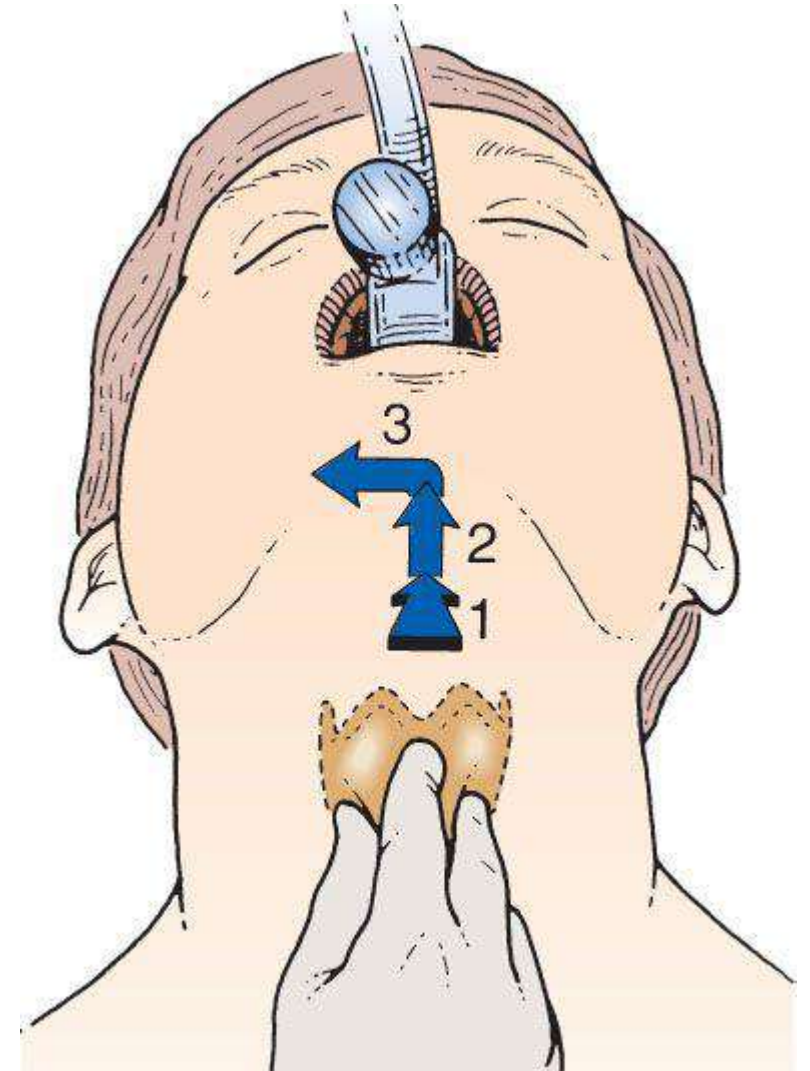
- **Intubating stylets**
- External laryngeal manipulation
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- Intubating supraglottic airway
- Combined techniques





Establish secure airway

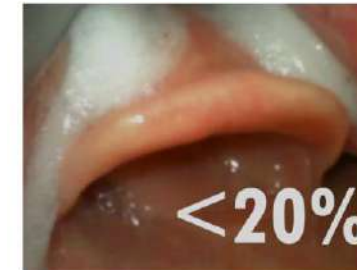
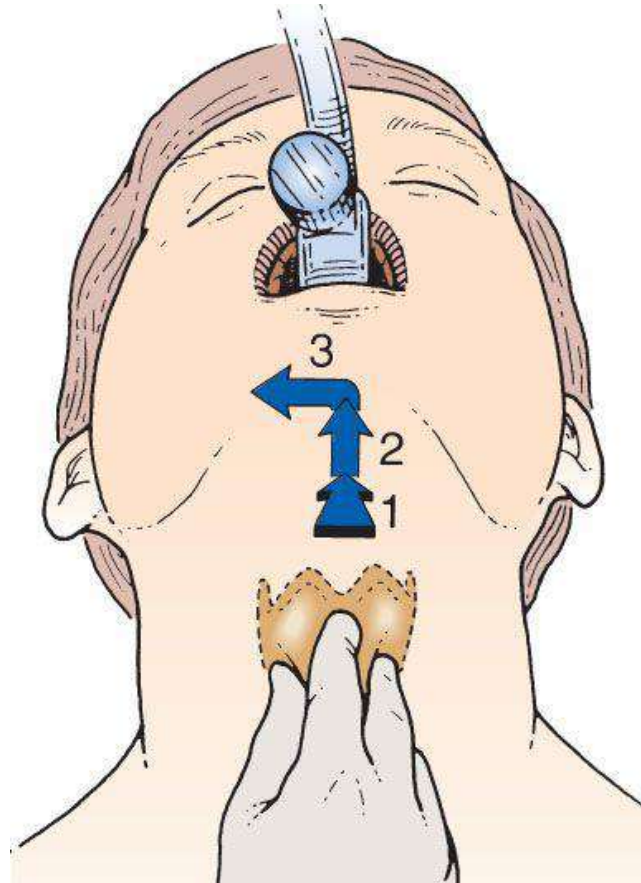
- Intubating stylets
- **External laryngeal manipulation (BURP maneuver)**
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- Intubating supraglottic airway
- Combined techniques



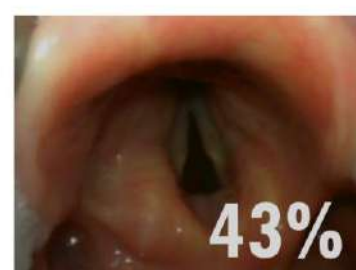
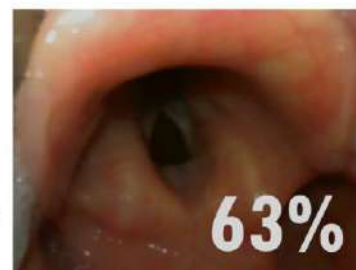
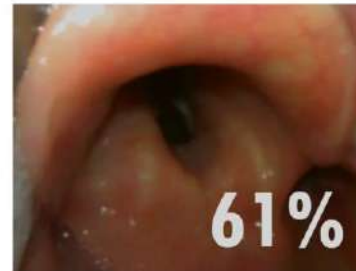


External laryngeal manipulation (BURP maneuver)

- B – Backward
- U – Upward
- RP – Rightward pressure



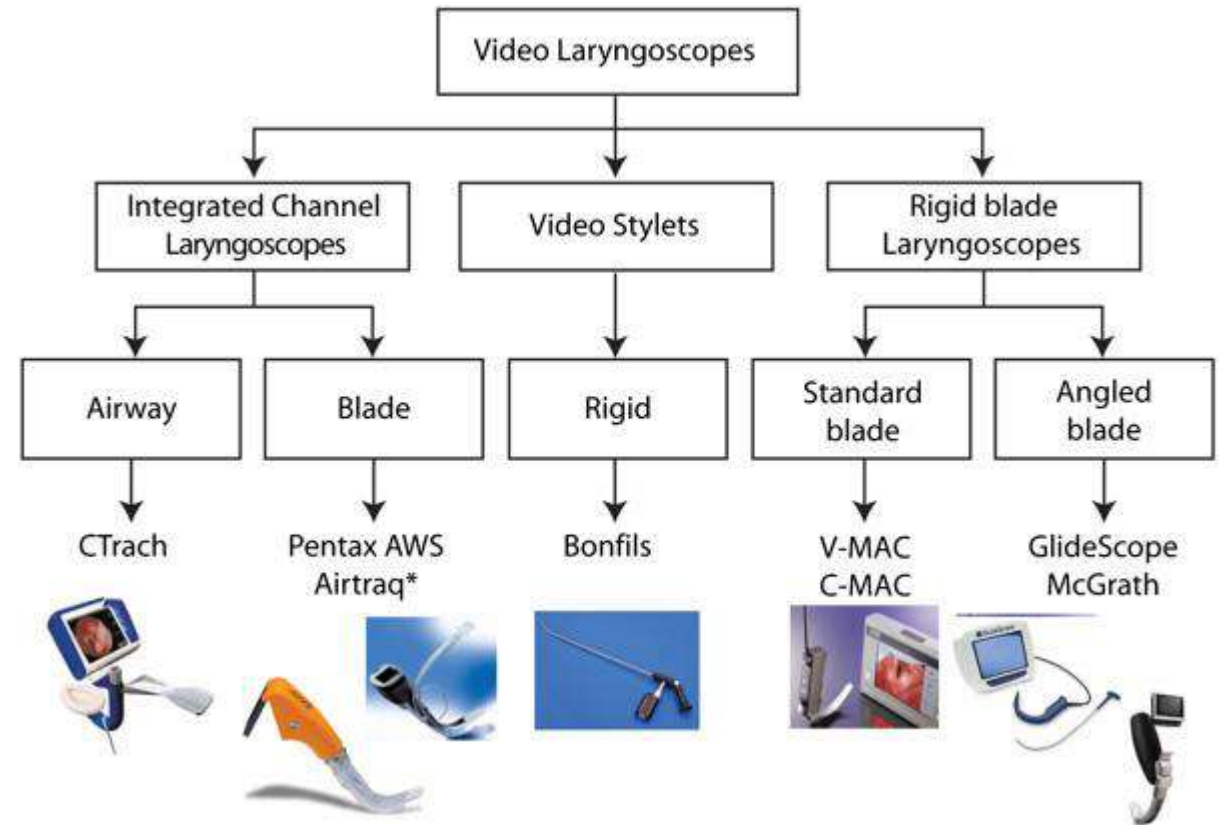
ELM
→
% IMPVMT





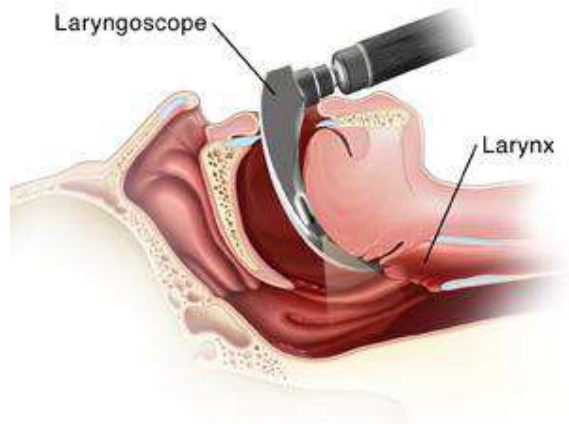
Establish secure airway

- Intubating stylets
- External laryngeal manipulation
- **Video-assisted laryngoscopy**
- Alternative laryngoscope blades
- Intubating supraglottic airway
- Combined techniques





Direct VS Video-assisted laryngoscopy





Video-assisted laryngoscopy

Indications and advantages

- Unnecessary to align airway axes (oral-pharyngeal-laryngeal)
- Improved glottic visualization, (limited mouth opening or neck mobility)
- Allows others to view the screen and/or help
- facilitate ETI (e.g., redirect cricoid pressure, acquire other airway devices)
- Less cervical manipulation
- Possible awake assessment/intubation
- Can provide an official record.

Disadvantages

- Difficulty in passing ETT despite improved glottic visualization (especially with angulated blade)
- Possible increased intubation time; variable learning curve
- Potential for false sense of security and lack of preparation for difficult airway
- Two-dimensional view with loss of depth perception;
- Obscured view by fogging and secretions on camera lens



Video-assisted laryngoscopy

Indications and advantages

- Unnecessary to align airway axes (oral-pharyngeal-laryngeal)
- Improved glottic visualization, (limited mouth opening or neck mobility)
- Allows others to view the screen and/or help facilitate ETI (e.g., redirect cricoid pressure, acquire other airway devices)
- Less cervical manipulation
- Possible awake assessment/intubation
- Can provide an official record.

Disadvantages

- Difficulty in passing ETT despite improved glottic visualization (especially with angulated blade)
- Possible increased intubation time; variable learning curve
- Potential for false sense of security and lack of preparation for difficult airway
- Two-dimensional view with loss of depth perception;
- Obscured view by fogging and secretions on camera lens



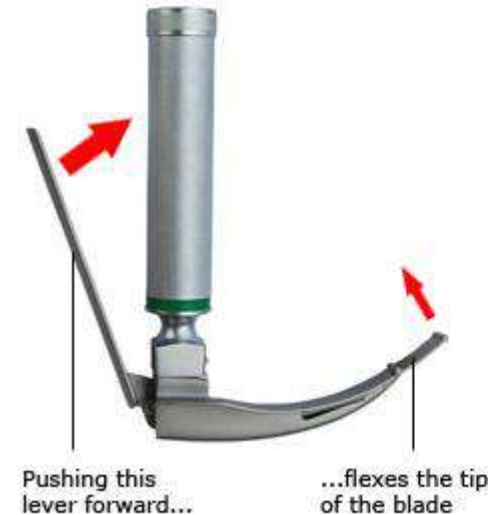
Establish secure airway

- Intubating stylets
- External laryngeal manipulation
- Video-assisted laryngoscopy
- **Alternative laryngoscope blades**
- Intubating supraglottic airway
- Combined techniques

Alternative laryngoscope blades

McCoy laryngoscope

The flexible tip helps view an anterior larynx by elevating the epiglottis



Miller laryngoscope blades

Straight blade (straight line view, better if poor mouth opening)





Establish secure airway

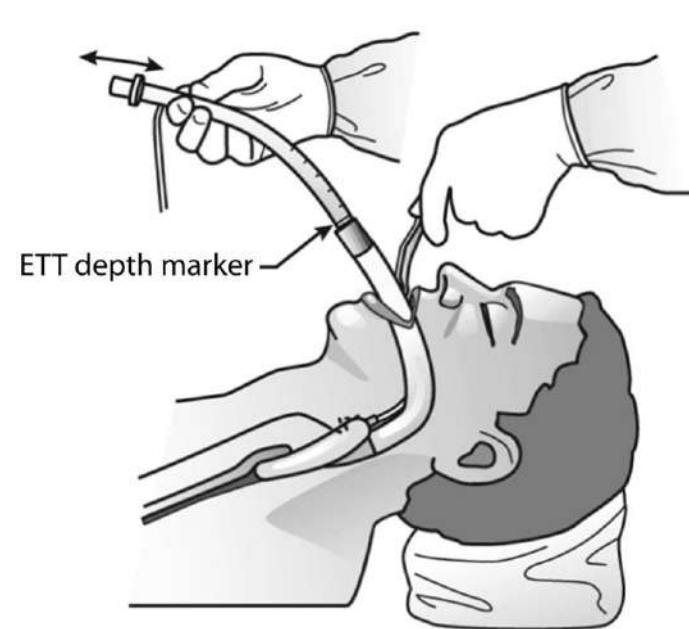
- Intubating stylets
- External laryngeal manipulation
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- **Intubating supraglottic airway**
- Combined techniques





Establish secure airway

- Intubating stylets
- External laryngeal manipulation
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- **Intubating supraglottic airway**
- Combined techniques





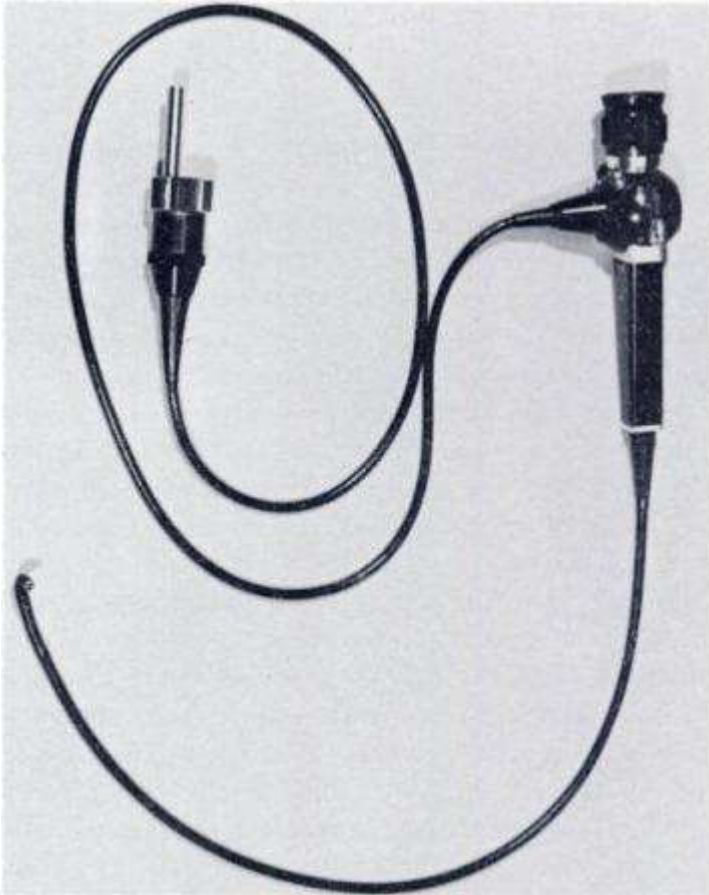
Establish secure airway

- Intubating stylets
- External laryngeal manipulation
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- Intubating supraglottic airway
- **Combined techniques**





Flexible Bronchoscope Intubation





Flexible Bronchoscope Intubation

- Both anticipated and unanticipated difficult airways
- Awake, sedated, and anesthetized patients.
- Orotracheal and nasotracheal routes
- Insulation of these fibers by a glass layer with a different optical density enables transmission by internal reflection of light



Flexible Bronchoscope Intubation

Indications and advantages

- Limited mouth opening
- Abnormal airway anatomy/mass obstructing direct visualization of vocal cords
- Unstable cervical spine
- Airway trauma requiring visualization of larynx and trachea prior to intubation
- Prone/Lateral position requiring rescue intubation

Contraindications and disadvantages

- Blood or secretion in the airway, severe maxillofacial injury
- Need for rapid control of the airway
- Clinician inexperienced
- Coagulopathy (risk of epistaxis)
- Allergy to local anesthetics
- Refusal or uncooperative patient



Flexible Bronchoscope Intubation

Indications and advantages

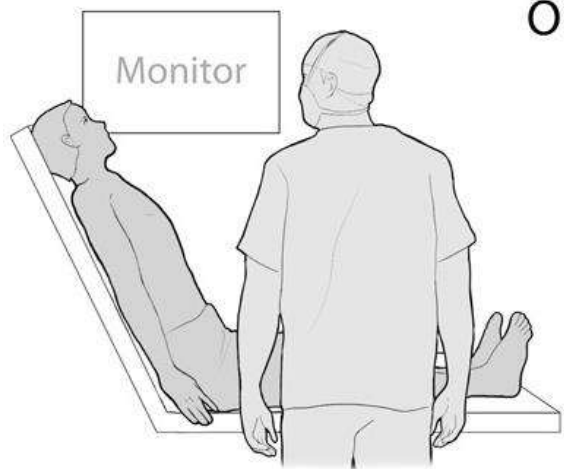
- Limited mouth opening
- Abnormal airway anatomy/mass obstructing direct visualization of vocal cords
- Unstable cervical spine
- Airway trauma requiring visualization of larynx and trachea prior to intubation
- Prone/Lateral position requiring rescue intubation

Contraindications and disadvantages

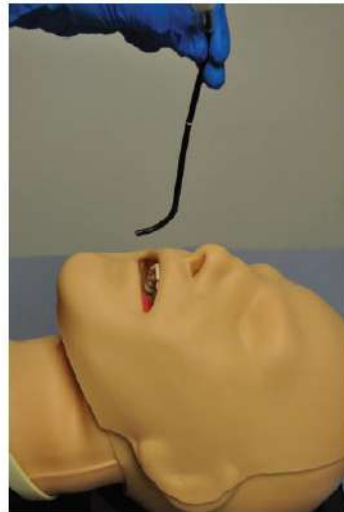
- Blood or secretion in the airway, severe maxillofacial injury
- Need for rapid control of the airway
- Clinician inexperienced
- Coagulopathy (risk of epistaxis)
- Allergy to local anesthetics
- Refusal or uncooperative patient



Positioning

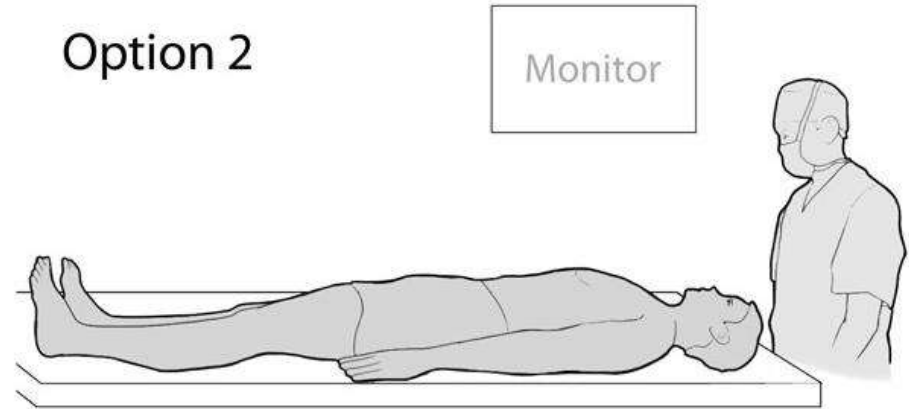


Option 1

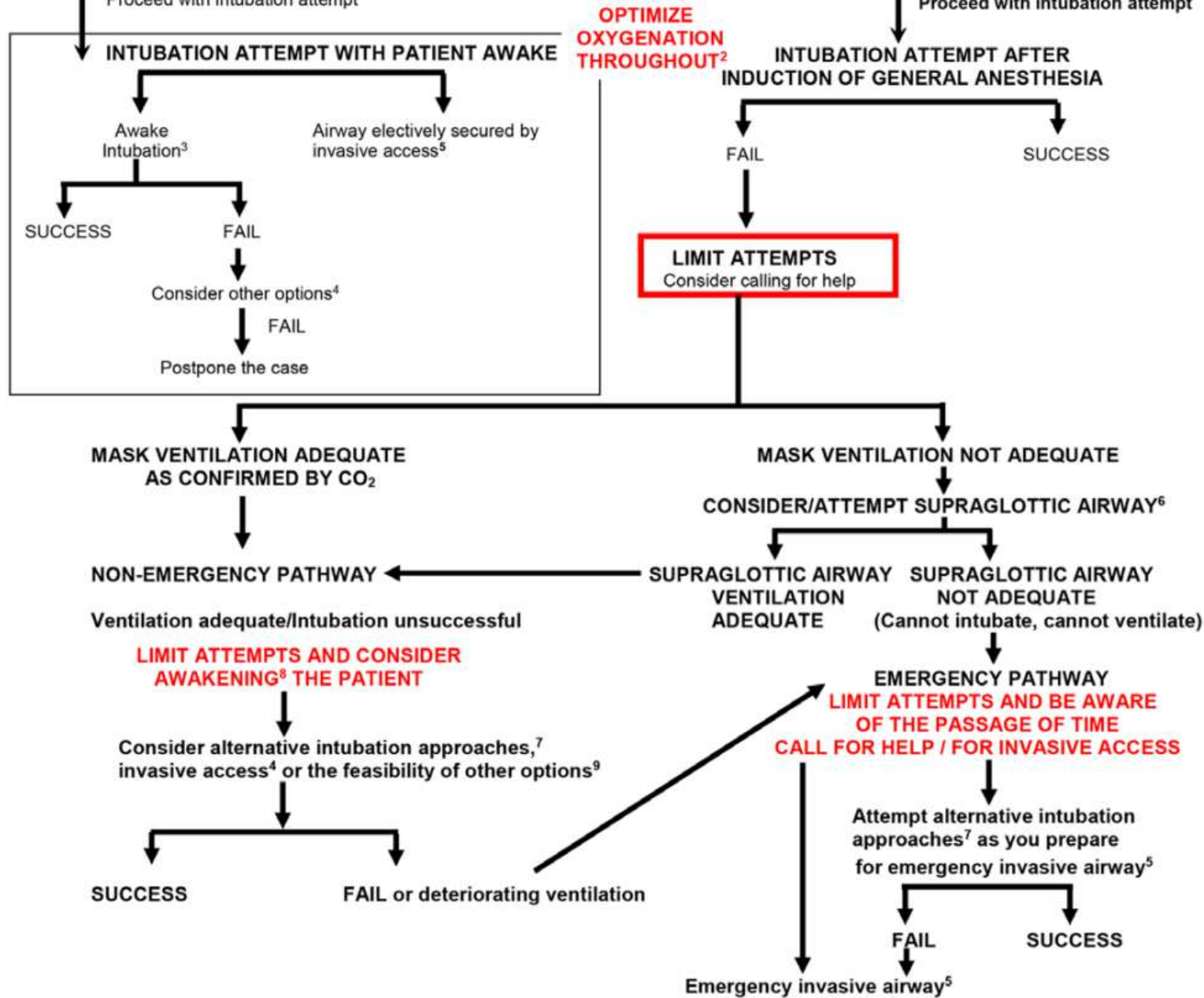


- ❖ Patient is seated or from the side of the bed, the tip of the scope is angulated down at an angle of 45°

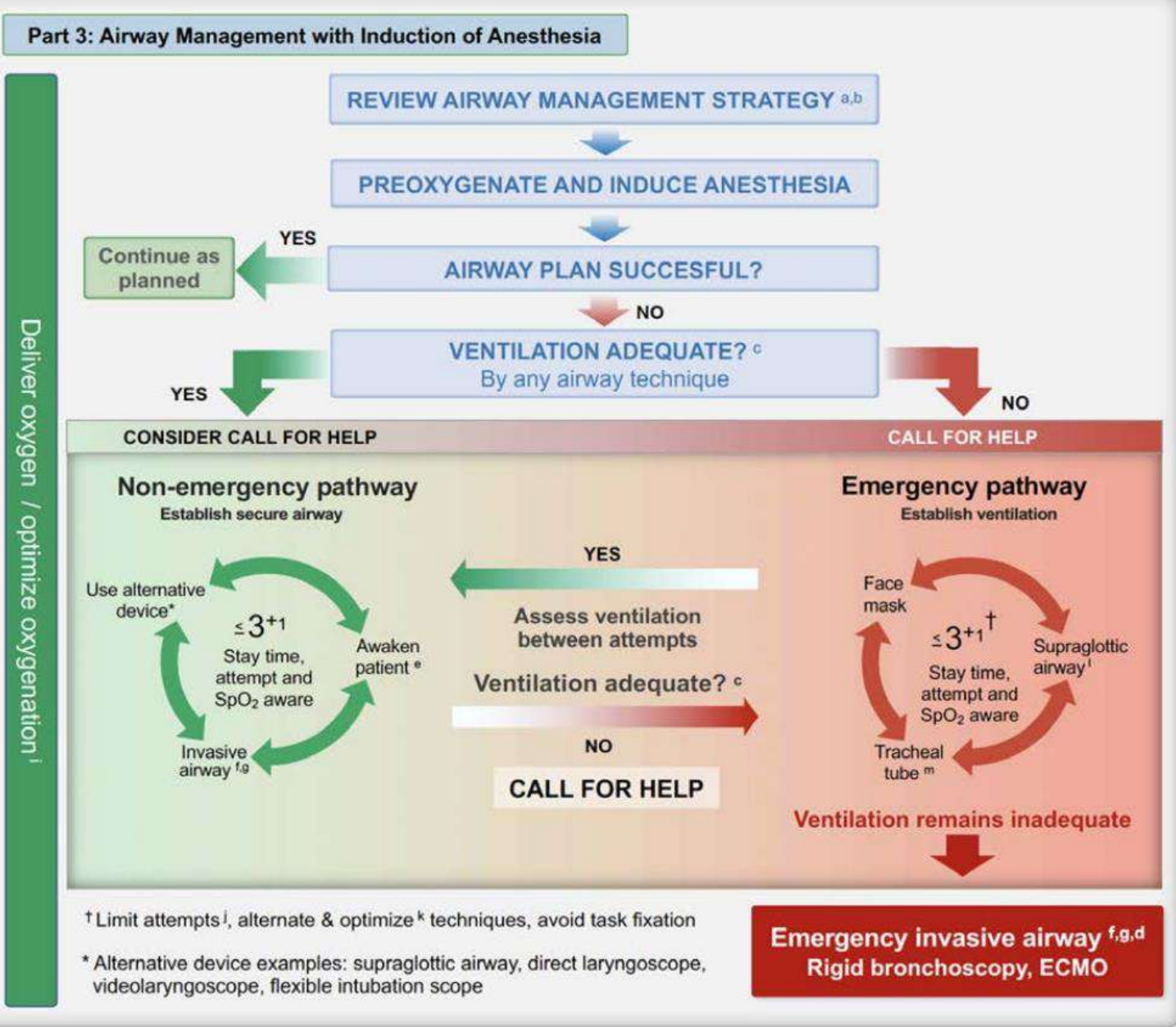
Option 2



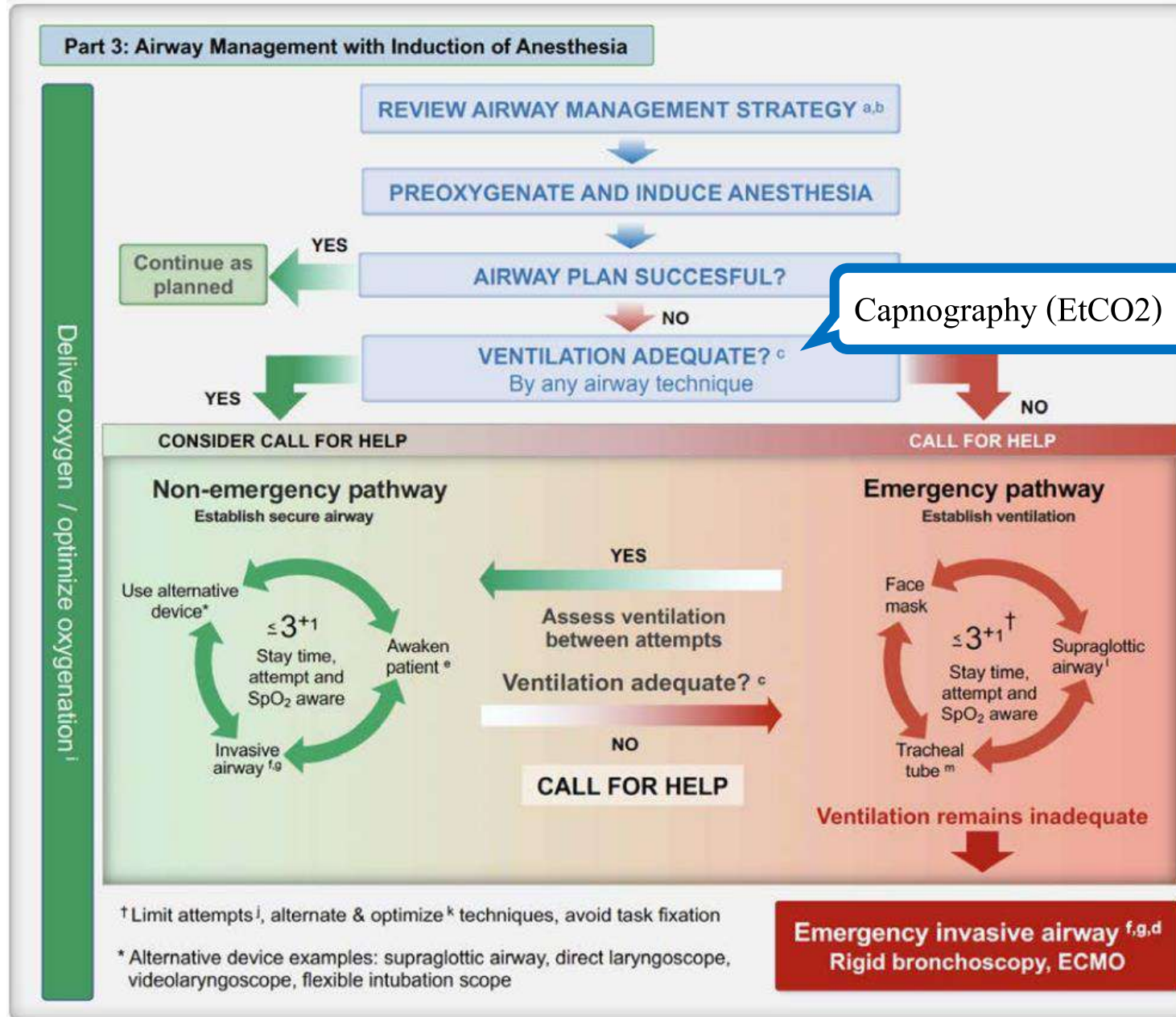
- ❖ Scope from the head of the bed, the tip of the scope is angulated up at an angle of 45°



Guidelines for Management of the Difficult Airway



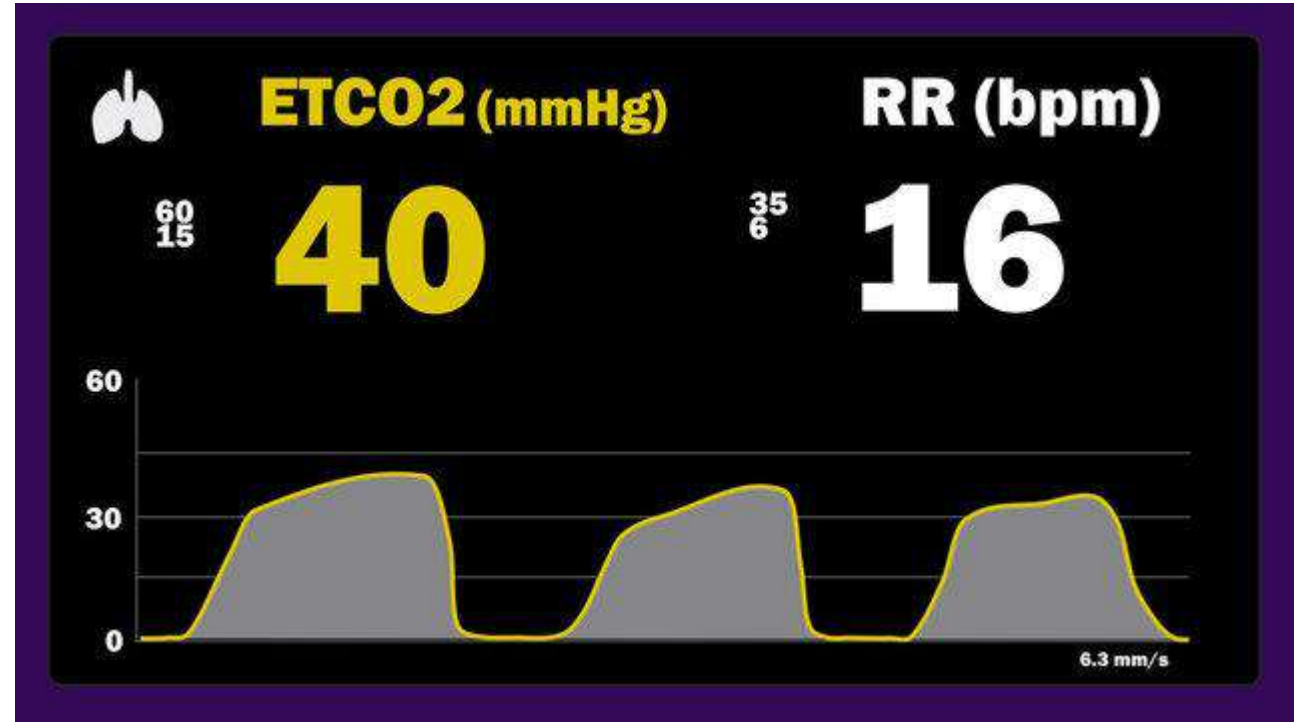
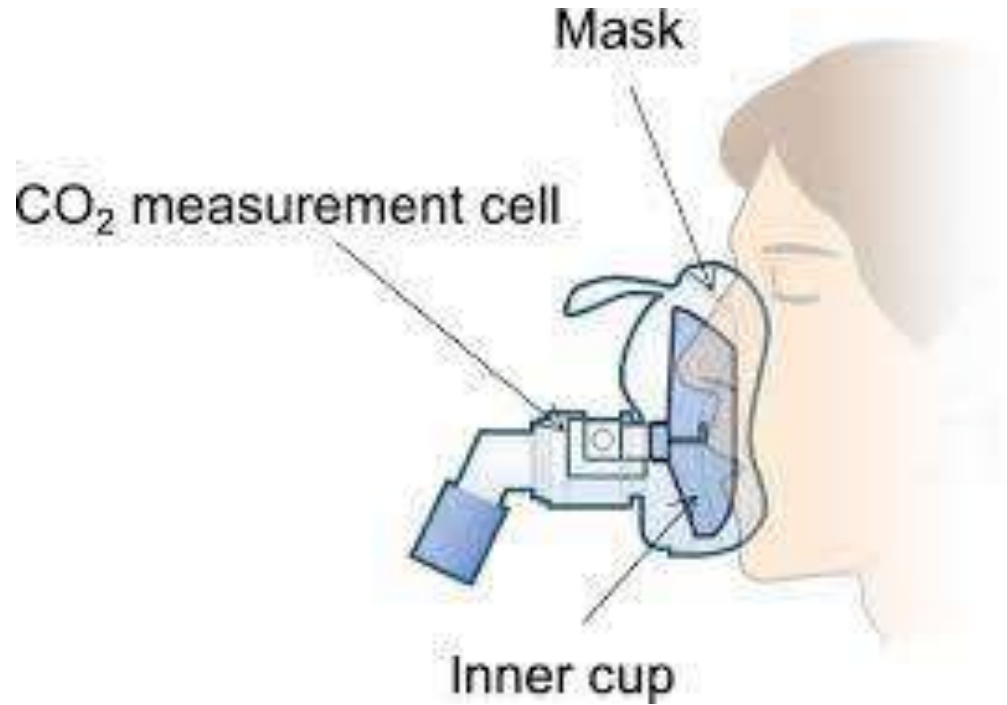
Guidelines for Management of the Difficult Airway



Guidelines for Management of the Difficult Airway

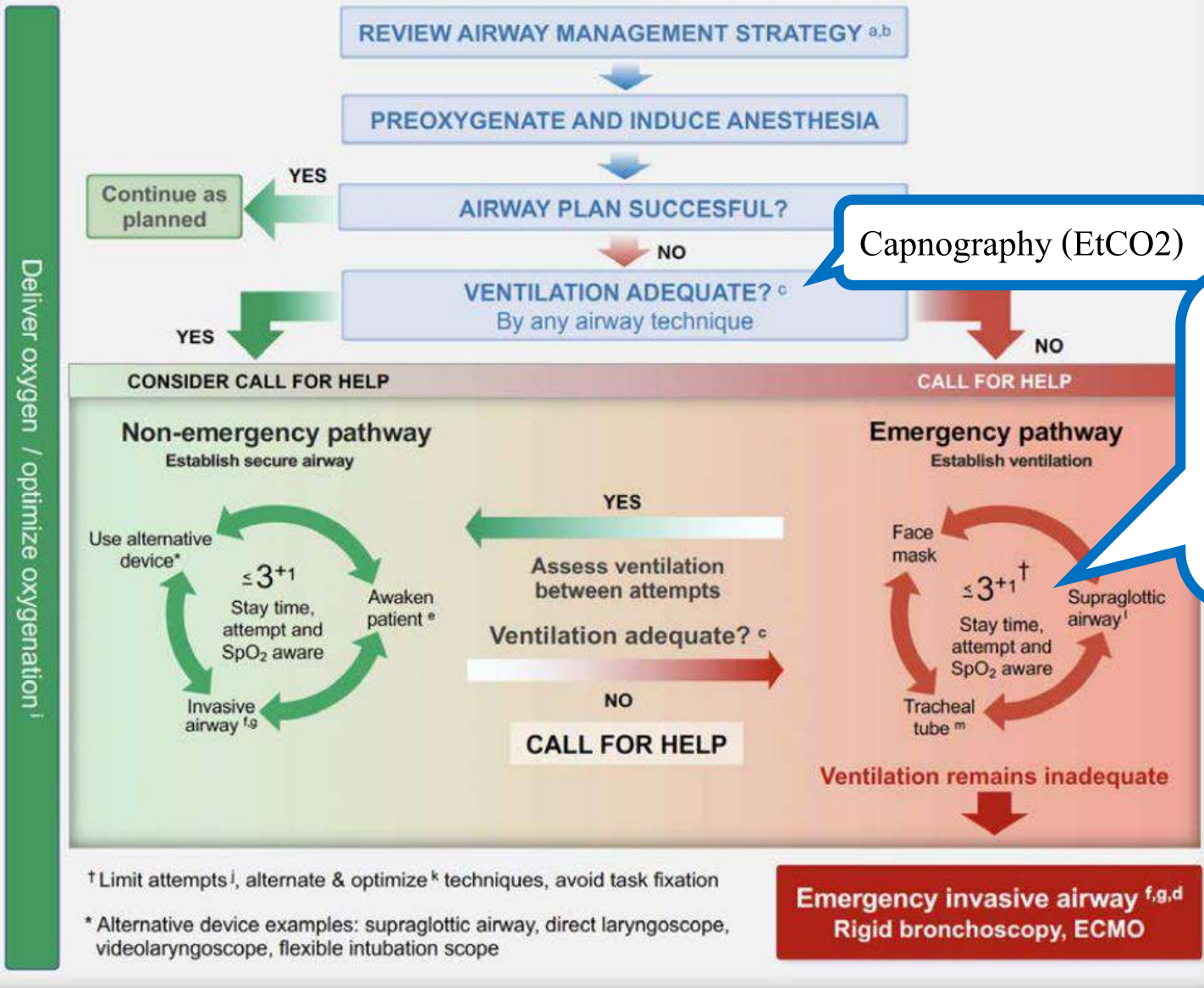


Capnography (EtCO₂)





Part 3: Airway Management with Induction of Anesthesia



Guidelines

for

- Suction
- Repositioning
- Oral/nasal airway
- Two-hand mask grip
- Supraglottic airway

ement

Difficult

Airway

† Limit attemptsⁱ, alternate & optimize^k techniques, avoid task fixation
 * Alternative device examples: supraglottic airway, direct laryngoscope, videolaryngoscope, flexible intubation scope



Establish ventilation

- Suction
- Repositioning
- Oral/nasal airway
- Two-hand mask grip
- Supraglottic airway



Establish ventilation

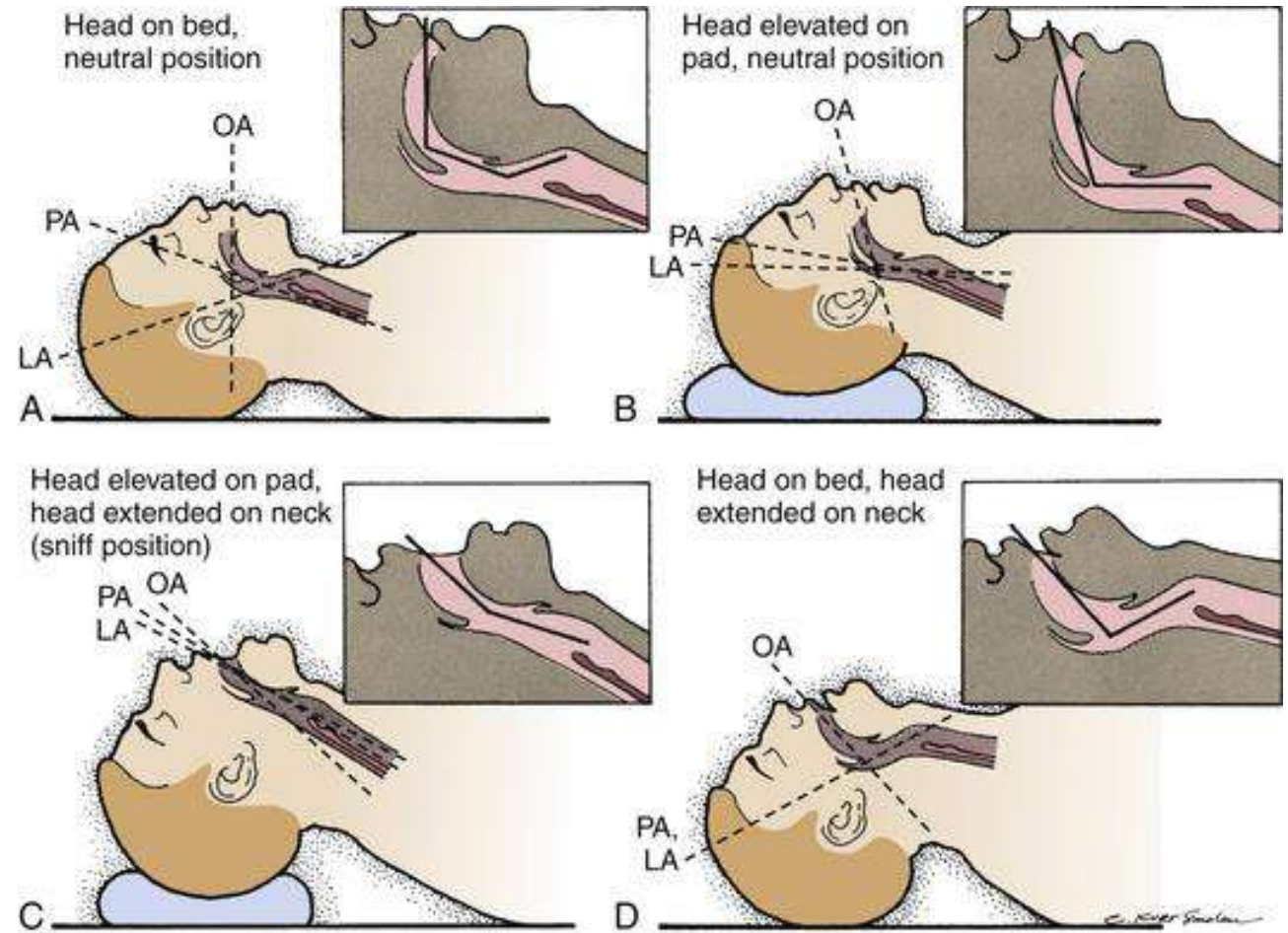
- **Suction**
- Repositioning
- Oral/nasal airway
- Two-hand mask grip
- Supraglottic airway





Establish ventilation

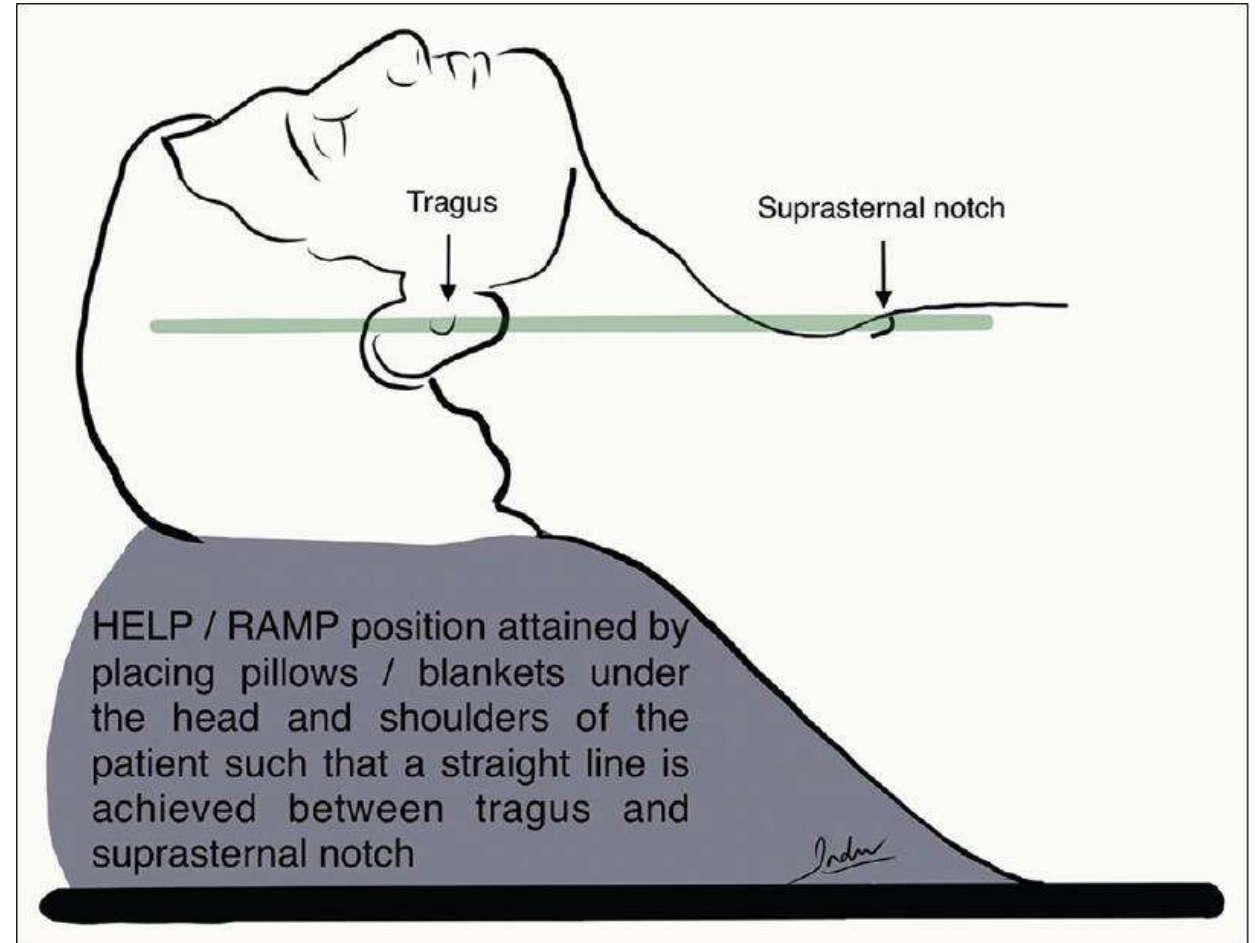
- Suction
- **Repositioning**
- Oral/nasal airway
- Two-hand mask grip
- Supraglottic airway





Establish ventilation

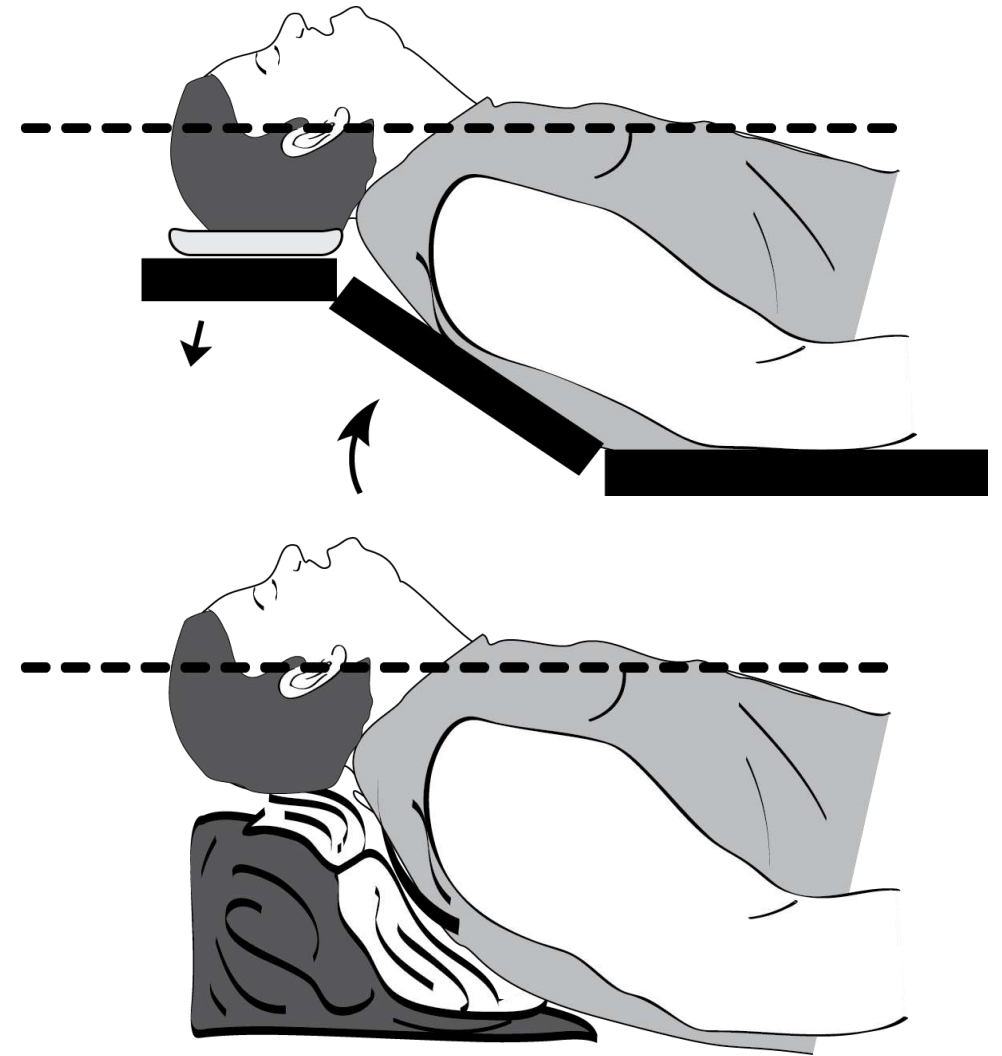
- Suction
- **Repositioning**
- Oral/nasal airway
- Two-hand mask grip
- Supraglottic airway





Establish ventilation

- Suction
- **Repositioning**
- Oral/nasal airway
- Two-hand mask grip
- Supraglottic airway





Establish ventilation

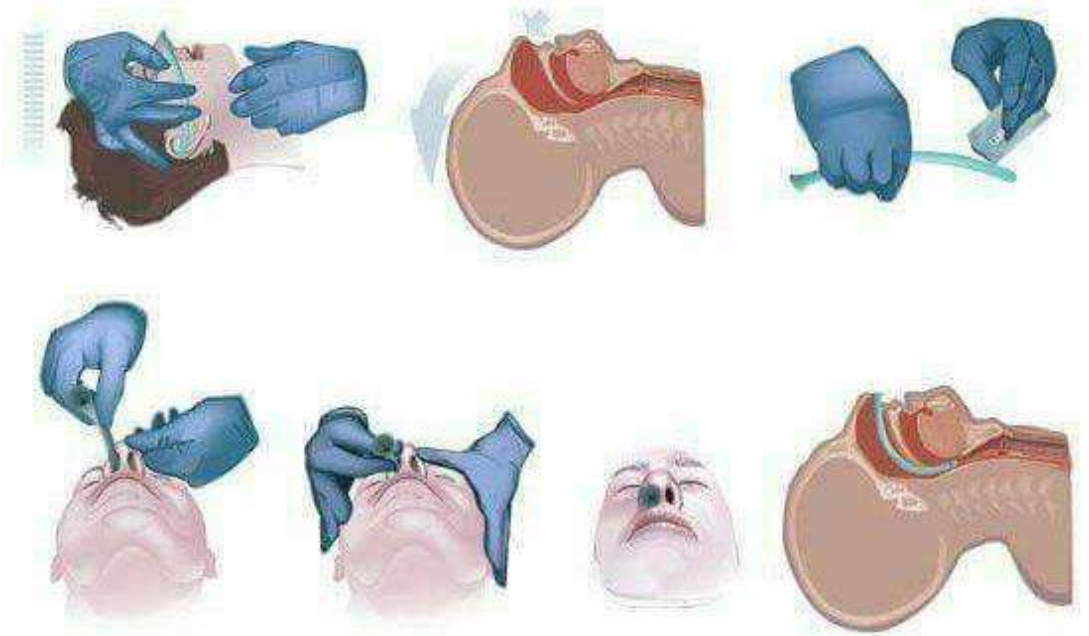
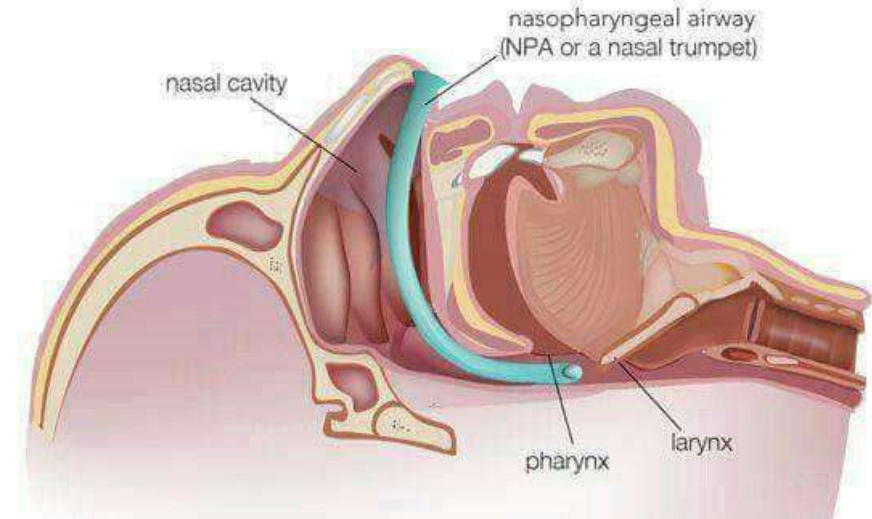
- Suction
- Repositioning
- **Oral/nasal airway**
- Two-hand mask grip
- Supraglottic airway





Establish ventilation

- Suction
- Repositioning
- **Oral/nasal airway**
- Two-hand mask grip
- Supraglottic airway





Establish ventilation

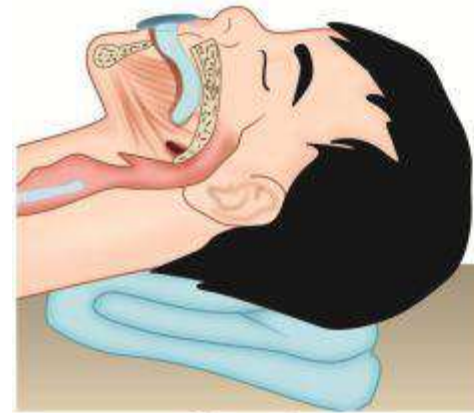
- Suction
- Repositioning
- **Oral/nasal airway**
- Two-hand mask grip
- Supraglottic airway



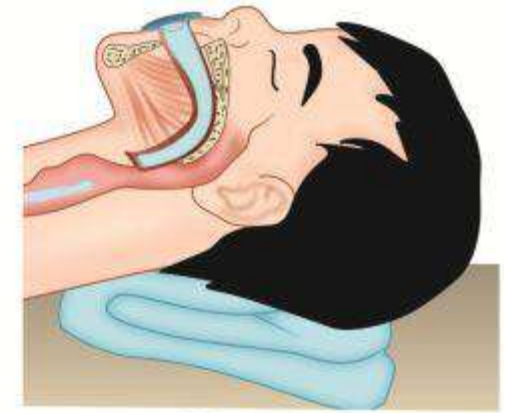


Establish ventilation

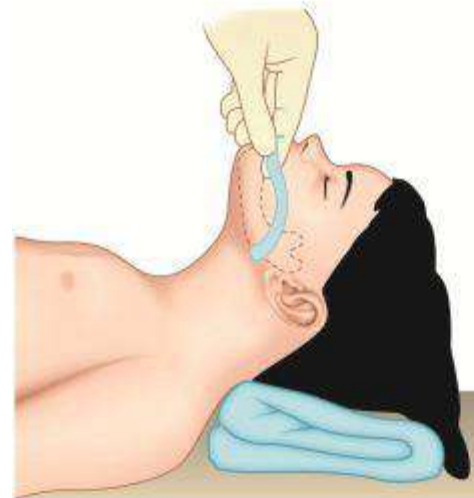
- Suction
- Repositioning
- **Oral/nasal airway**
- Two-hand mask grip
- Supraglottic airway



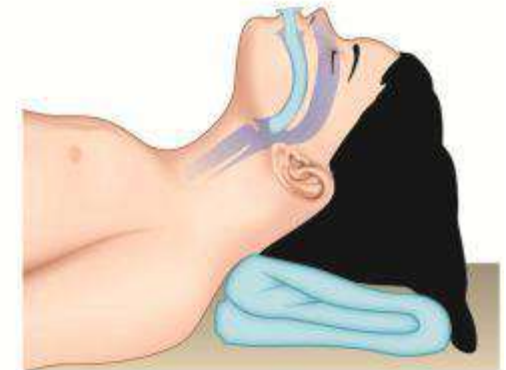
Short size



Bigger size



Proper size nasopharyngeal airways



Proper position



Establish ventilation

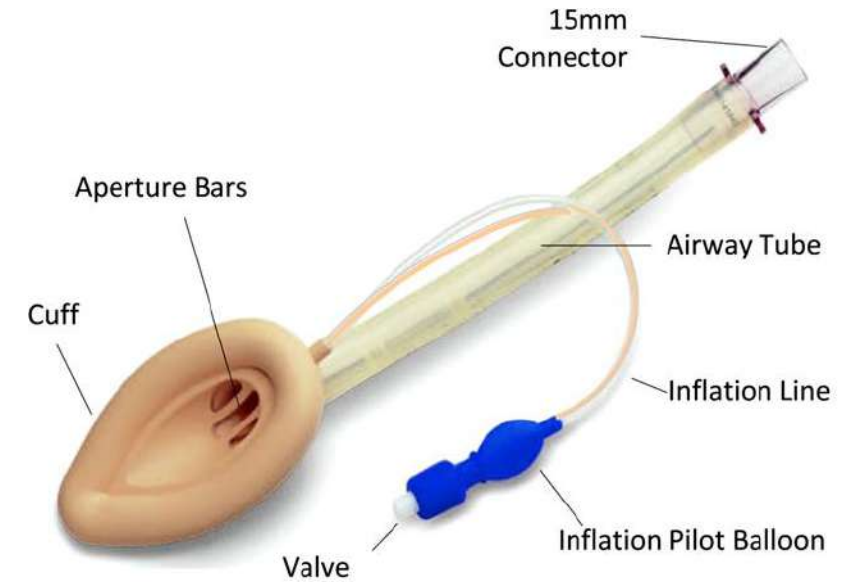
- Suction
- Repositioning
- Oral/nasal airway
- **Two-hand mask grip**
- Supraglottic airway





Establish ventilation

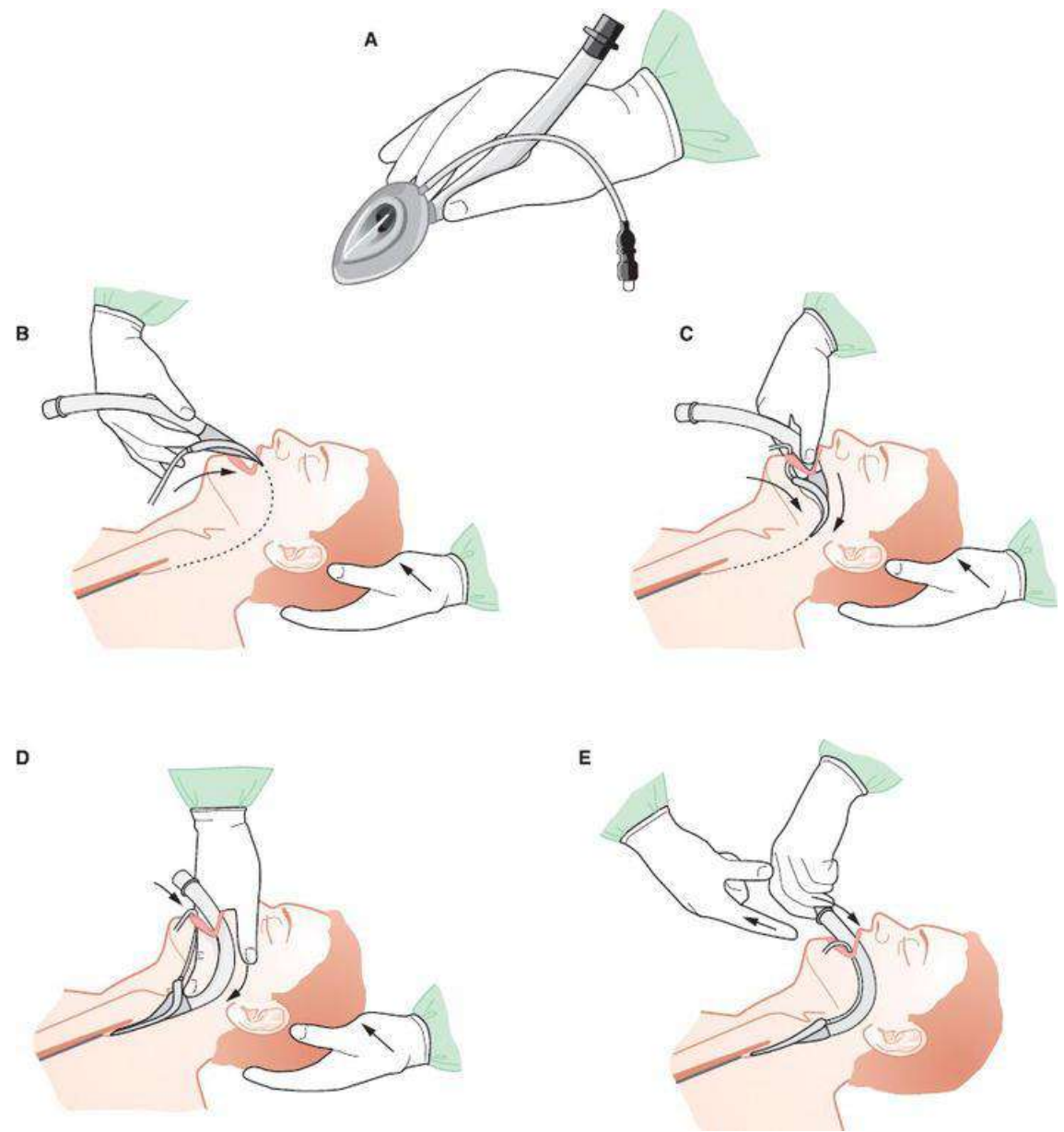
- Suction
- Repositioning
- Oral/nasal airway
- Two-hand mask grip
- **Supraglottic airway**





Establish ventilation

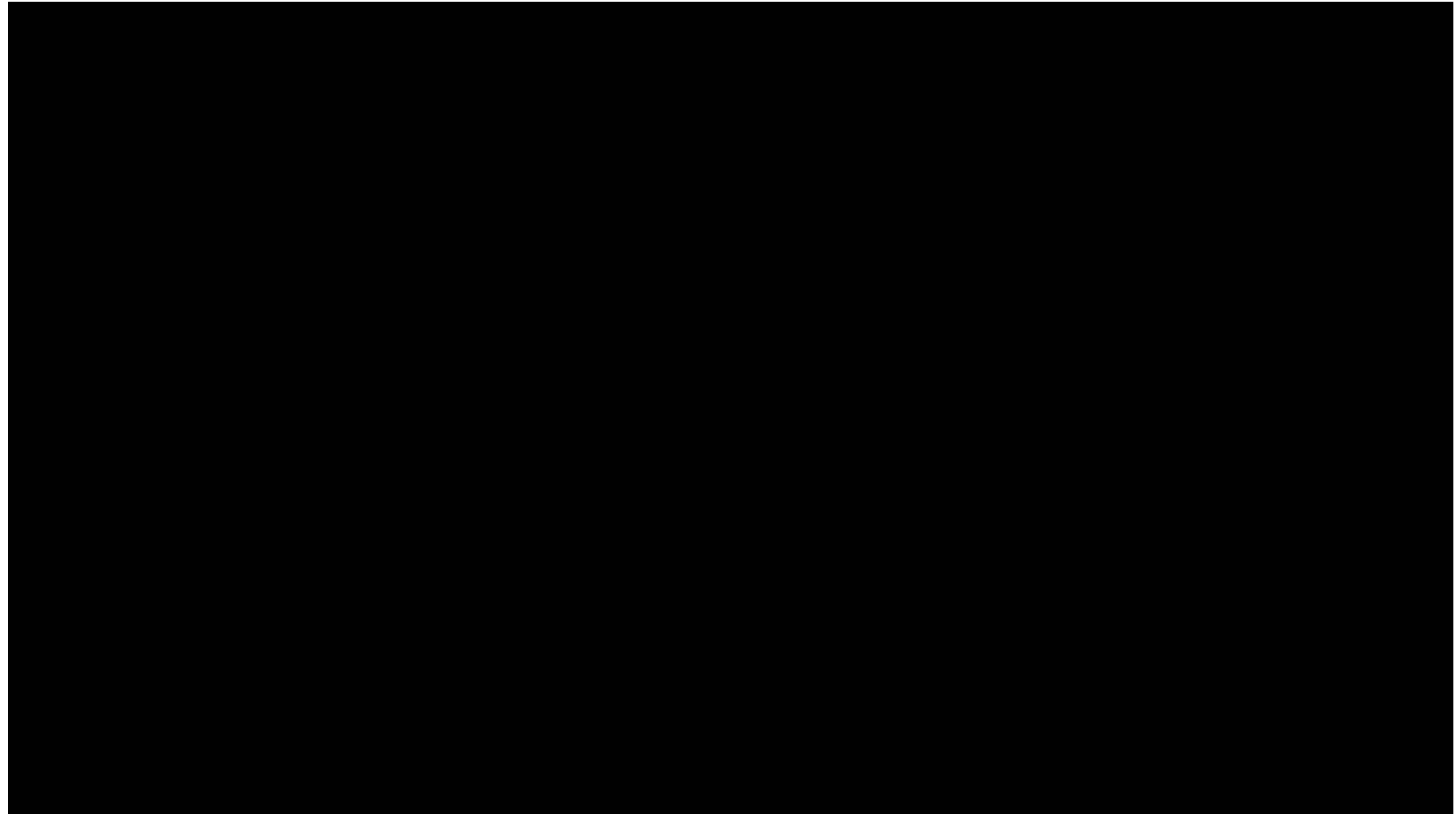
- Suction
- Repositioning
- Oral/nasal airway
- Two-hand mask grip
- **Supraglottic airway**

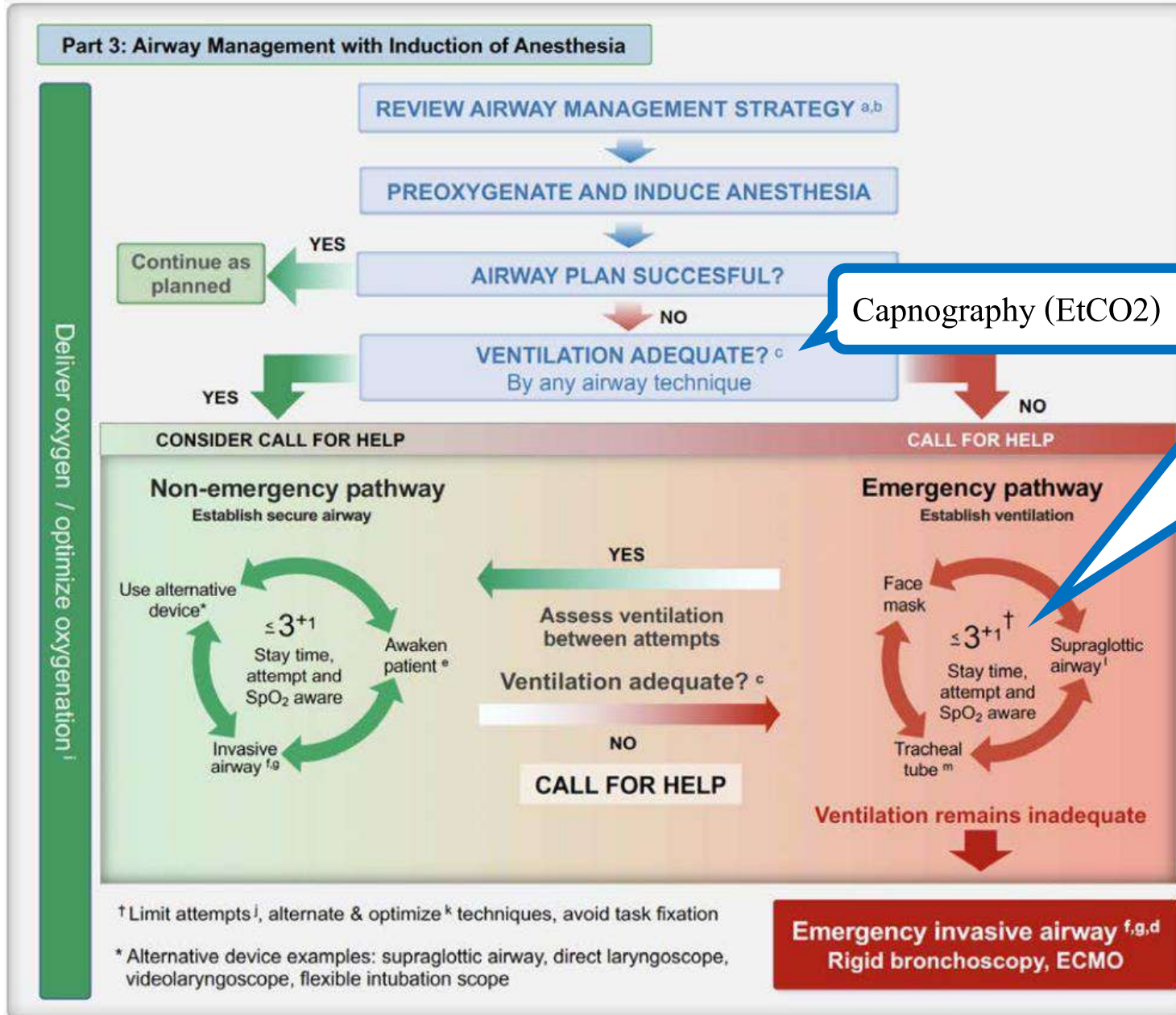




Establish ventilation

- Suction
- Repositioning
- Oral/nasal airway
- Two-hand mask grip
- **Supraglottic airway**

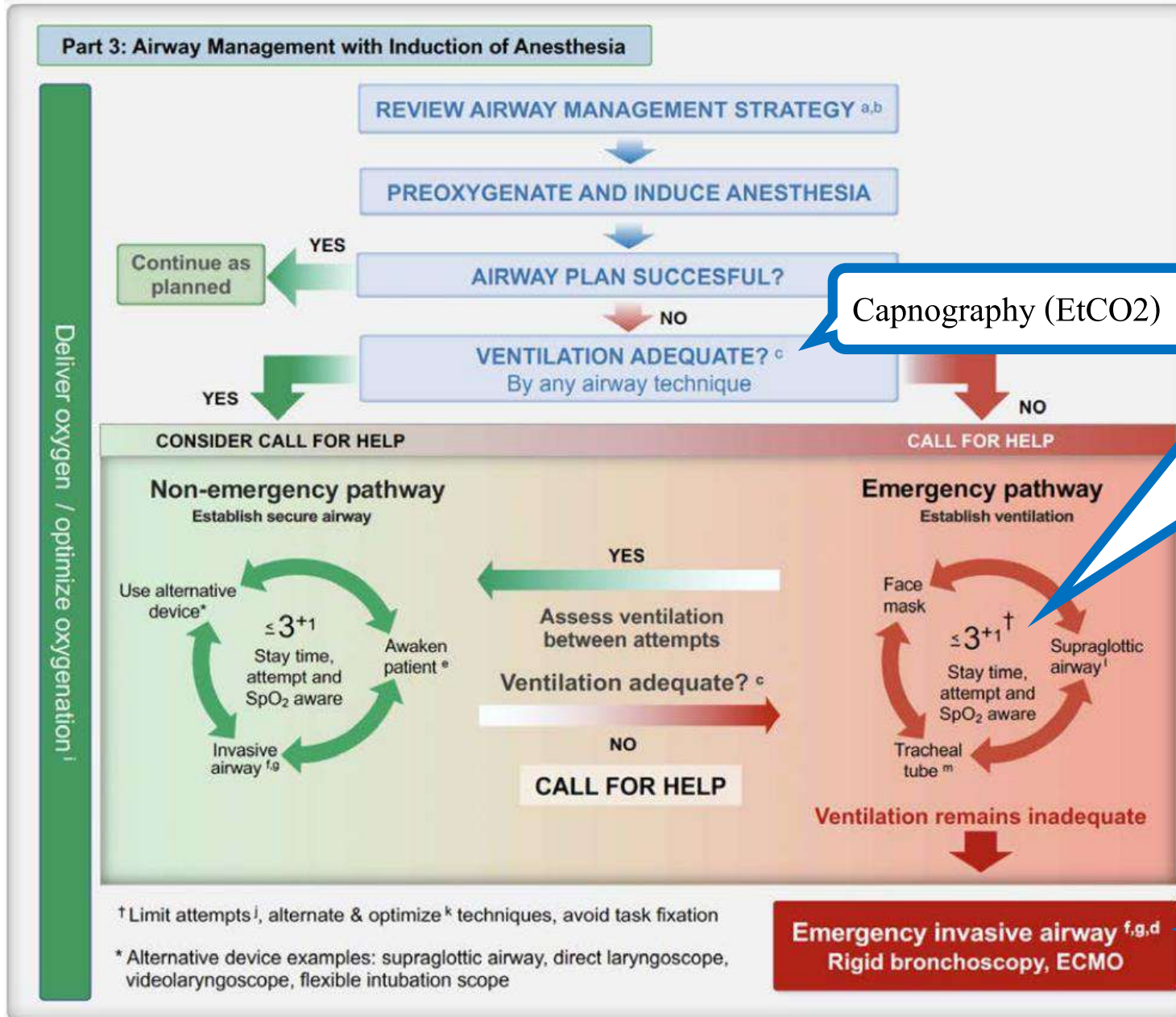




Capnography (EtCO₂)

- Intubating stylets
- External laryngeal manipulation
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- Combined techniques
- Intubating supraglottic airway

Difficult Airway



Capnography (EtCO₂)

- Intubating stylets
- External laryngeal manipulation
- Video-assisted laryngoscopy
- Alternative laryngoscope blades
- Combined techniques
- Intubating supraglottic airway

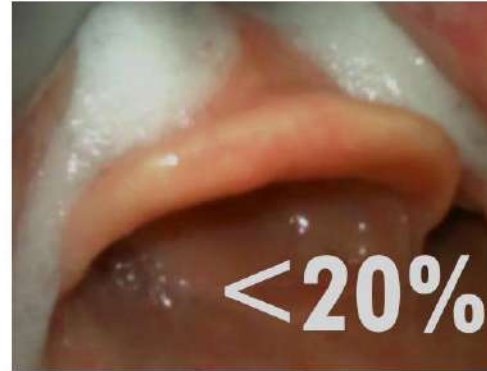
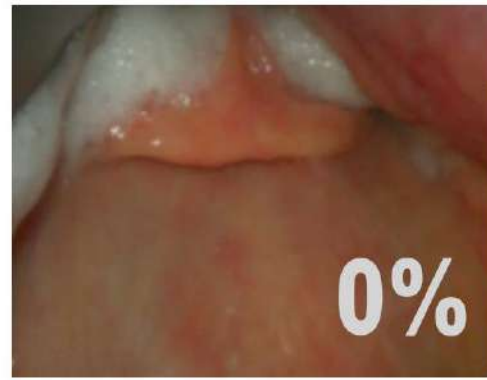
Difficult Airway

- Surgical cricothyrotomy
- Needle cricothyrotomy
- Surgical tracheostomy



What is the problem?

- Uncooperative patient
- Can't see vocal cord
 - Laryngeal view grade $> I$
 - Obscured by Secretion / Blood / Mass
- Seen vocal cord, but can't insert endotracheal tube into vocal cord
 - Can't control tip of ETT to vocal cord
 - Vocal cord edema
- Limited mouth opening or neck movement





What is the problem?

- Uncooperative patient
 - Proper sedation
 - Rapid sequence induction

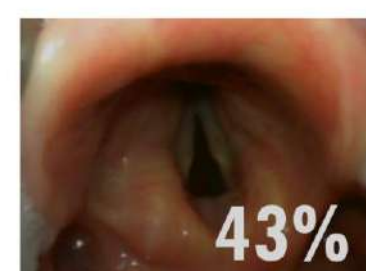
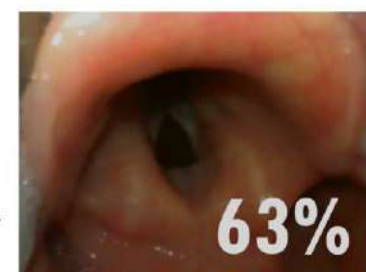
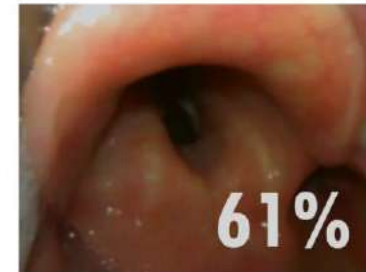


What is the problem?

- Can't see vocal cord **due to poor LV**
 - Appropriated position – Sniff position
 - BURP maneuver
 - Appropriated laryngoscope blades
 - Video laryngoscopy or combine techniques



ELM
→
% IMPVMT





What is the problem?

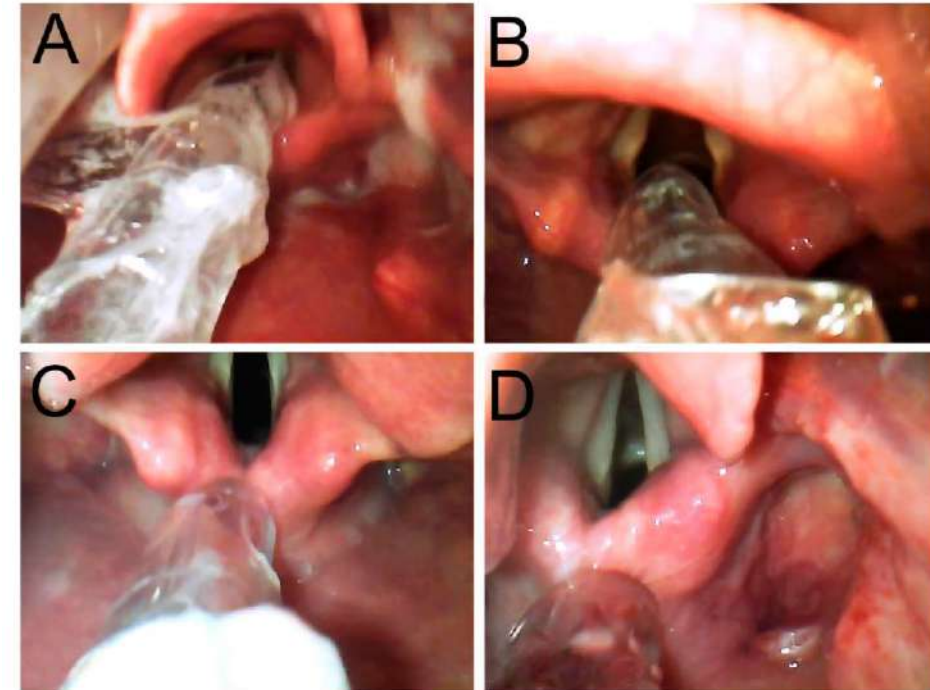


- Can't see vocal cord **due to obscured by Secretion / Blood**
 - Suction
 - Limit attempts
- Can't see vocal cord **due to obscured by large mass e.g. base of tongue tumor**
 - Limit attempts and consider calling for help
 - Consider role of fiberoptic or invasive airway by specialist



What is the problem?

- Seen vocal cord, but can't insert endotracheal tube into vocal cord
 - BURP maneuver
 - Intubating stylets with appropriated curve (correlation with laryngoscope curve blades)
 - Fiberoptic or combine techniques
 - Smaller endotracheal tube size





What is the problem?

- Limited mouth opening or neck movement -> Evaluation cause of limitation
 - Incorporating patient -> Sedation or induction
 - Pain -> Pain controller e.g. Fentanyl, Pethidine



What is the problem?

- Limited mouth opening or neck movement
 - Collar mask
 - **Manual in line stabilization with video laryngoscopy**

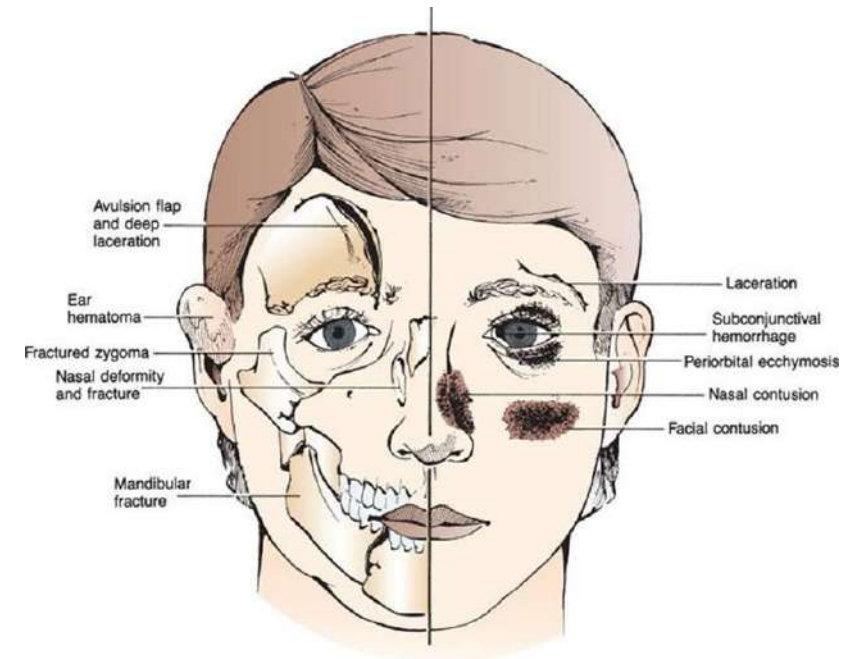


MILS during intubation - standing opposite to intubator



What is the problem?

- Limited mouth opening or neck movement
 - Anatomy defects
 - Limit attempts and consider calling for help
 - Consider role of fiberoptic or invasive airway by specialist





Take home messages

- Face mask ventilation is KEY for survival
- Limit attempts and consider calling for help



ภาควิชากุมารเวชศาสตร์ คณะแพทยศาสตร์มหาวิทยาลัยนเรศวร
Faculty of Medicine Naresuan University



Breaking Bad News

Assoc.Prof.Jiranun Weerakul



What is bad news?

“any information which adversely and seriously effects an individual’s view of his or her future”



ภาควิชากุมารเวชศาสตร์ คณะแพทยศาสตร์มหาวิทยาลัยนเรศวร
Faculty of Medicine Naresuan University



Example of medical bad news





Example of medical bad news

- Informing patients that they have cancer.
- Inform the patient that he/she or her child is HIV positive.
- Inform the first degree relative that the patient has brain death.
- Inform the mother that the child has Down syndrome.
- Inform the patient or relative that he/she need to amputate.



Why is it important?

- A frequent and stressful task
- Breaking bad news can be particularly stressful when the doctor is inexperienced, the patient is young or there are limited prospects for successful treatment



The patients want the truth

- By the late 1970s most physicians were open about telling cancer patient their diagnosis
- In 1982 of 1,251 American indicated that 96% wished to be told if they had diagnosis of cancer
- 85% wished, in case of grave prognosis, to be given a realistic estimate of how long they had to live



Ethical and legal imperatives

- Clear ethical and legal obligations to provide patients with as much information as they desire about their illness and its treatment
- Physicians may not withhold medical information even if they suspect it will have a negative effect on the patient



Clinical outcomes

- How bad news is discussed can effect the patient's comprehension of information, satisfaction with medical care, level of hopefulness, and subsequent psychological adjustment



Barriers to breaking bad news

- Emotional-anxiety
- Burden of responsibility
- Fear of negative evaluation



Principle of breaking bad news

| D | Doctor | Compression. Loving-kindness, empathy |
|---|-------------|---|
| P | Patient | Anxiety, fear, Worries |
| I | Information | Step-by-step approach depending on the patient's capacity to assimilate it |
| H | Hope | Always commit to be on the patient's side, find a way to help especially psychological well-being |



Models of Breaking bad news

- SPIKES model
 - Robert Buckman
 - Professor of oncology-Toronto
 - Trained in Cambridge
 - Used world wide
- KAYES model
- ABCDE model

Buckman R. Breaking bad news: why is it so difficult?. BMJ. 1984;288:1597-9



SPIKES Model

Six steps

- **S-Setting** up the interview
- **P**-assessing the patients **Perception**
- **I**-obtaining the patients **Invitation**
- **K**-giving **Knowledge**
- **E**-addressing **Emotions**
- **S-Strategy** and **Summary**



S-Setting up the interview

- Privacy
- Involve others
- Look attentive and calm
- Listening mode
- Availability



P-Perception

- Ask before you tell
- Find out what the patient know



I-Invitation

- While a majority of patients express a desire for full information about their diagnosis, prognosis, and details of their illness, some patients do not
- How much information would the patient like to know



K-Knowledge

- Warming first
- Mirror language
- Avoid jargon
- Small chunks
- Use of silence
- Allow time for emotions



E-Emotions

- Recognize
- Listen for and identify the emotion
- Identify cause of emotion
- Show the patient you have identified both the emotion and its origin



E-Emotions

- Crying
- Anger
- Denial
- Bargaining
- Shock/silence



S-Strategy and Summary

- Understanding reduces fear
- Summarizes the discussion
- Strategy for future care
- Schedule next meeting
- Allow time for questions
- Leaflets



KAYE's model

- 10 steps
- Logical sequence
- Not based on rigorous research
- Can be used for any serious illness



1. Preparation

- Know all the facts
- Ensure privacy
- Find out who the patient would like present
- Introduce yourself



2. What dose the patient know?

- Open end questions
- Statements may make the best questions
- “How did it all start?”



3. Is more information wanted?

- Not forced on them
- “Would you like me to explain a bit more?”



4. Warning shots

- Not straight out with it
- “I'm afraid it looks rather serious”



5. Allow denial

- Allow the patient to control the amount of information they receive



6. Explain if requested

- Step by step
- Detail will not be remembered but the way you explain it will be



7. Listen to concerns

- “What are your concerns at the moment?”
- Allow time and space for answers



8. Encourage feelings

- Acknowledge the feelings
- Non-judgmental
- Vital step for patient satisfaction



9. Summarizes

- Concerns
- Plans for treatment
- Foster hope
- ? Written information



10.

- Availability
- Information
- Future needs will change



ABCDE technique

- A-Advance preparation
- B-Build a therapeutic environment or relationship
- C-Communicate well
- D-Deal with patient and family reactions
- E-Encourage and validate emotions



A-Advance preparation

- Arrange for **adequate time, privacy** and no interruptions (turn off or silent mode of mobile phone)
- Review relevant **clinical information**
- **Mental rehearse**, identify words or phrase to use and avoid
- Prepare **yourself emotionally**



B-Build a therapeutic environment or relationship

- Determine **what and how much the patient wants to know**
- Have **family or support persons** present
- **Introduce yourself** to everyone
- **Warm the patient** that bad news is coming
- Use **touch** when appropriate
- Schedule **follow-up** appointments



C-Communicate well

- Ask what the patient or family **already knows**
- Be frank but **compassionate**; avoid euphemisms and medical jargon
- Allow for **silence** and tears; proceed at the patient's pace
- Have the patient **describe his or her understanding** of the news; repeat this information at subsequent visits
- **Allow time to answer** questions; write things down and provide written information
- **Conclude each visit** with a summary and follow-up plan



D-Deal with patient and family reactions

- Assess and **response to the patient and the family's emotional reaction**; repeat at each visit
- Be **empathetic**
- Do **not argue** with or criticize colleagues



E-Encourage and validate emotions

- Explore **what the news means to the patient**
- Offer **realistic hope** according to the patient's goals
- Use **interdisciplinary** resources
- Take care of your own needs; **be attuned to the needs** of involved house staff and officer or hospital personnel



reference

- รัตนา สายพานิชย์. การแจ้งข่าวร้าย. ใน: มาโนช หล่อตระกูล, บรรณาธิการ. คู่มือการดูแลผู้มีปัญหา สุขภาพจิต และจิตเวชสำหรับแพทย์. นนทบุรี: สำนัก พัฒนาสุขภาพจิต กรมสุขภาพจิต; 2544, หน้า 143-56.
- Buckman R. How to break bad news : a guide for health care professionals. Baltimore, Md. : The Johns Hopkins University Press, 1992.
- Sonny Jerome, Breaking bad news.
- Buckman R. Breaking bad news: why is it so difficult?. BMJ. 1984;288:1597-9
- Baile WF, Buckman R, Lenzi R, Gloger G, Beale EA, Kudelka AP. SPIKES-A six-step protocol for delivering bad news: application to the patient with cancer. Oncologist 2000;5:302-11.
- GREGG K. VANDEKIEFT. Breaking Bad News. Fam Physician. 2001;64(12):1975-79



ภาควิชากุมารเวชศาสตร์ คณะแพทยศาสตร์มหาวิทยาลัยนเรศวร
Faculty of Medicine Naresuan University



Case study





เด็กหญิงอายุ 10 ปี ได้รับการวินิจฉัยว่าเป็น
มะเร็งกระดูก (Osteosarcoma) โดยมีก้อนที่
เข้าด้านซ้าย และ มีอาการปวดขาบริเวณก้อน
จนเดินไม่ไหว มา 6 เดือน



- วันนี้ แพทย์นัดมารดามาแจ้งผลการรักษา
หลังจากได้รับยาเคมีบำบัด และ ทำ MRI เพื่อ
ประเมินก่อนการผ่าตัด



SPIKES Model

- **S-Setting** up the interview:
- **P**-assessing the patients **Perception**
- **I**-obtaining the patients **Invitation**
- **K**-giving **Knowledge**
- **E**-addressing **Emotions**
- **S-Strategy** and **Summary**

Eye Problems in General

Phattharaphong Tantichariyangkul, MD, FICO



MED NU 2023: UNEASY SITUATION FOR GENERAL PHYSICIAN



| | SV | PALs | Degressive | Blue/UV coating | Coating | Photochromic | Sunglasses | Reduce Myopic Progression | Techonlogy | Individualised Fitting |
|------------|---|---|---|--|---|--|--|---|---|---|
| ESSILOR | EYEZEN start (antifatique, dualoptim) EYEZEN boost/plus (initial0.4/active0) | Varlux X series (Xclusive4D/Xclusive) Varlux E series (Easy-to-wear) Varlux S series (Nanoptix balance ,Sy) Varlux Physio 3.0 Varlux Comfort Max (Flex Optm) SV Roadpilot II SV Exceptio (-40 to +30 oe beyond) | Varlux Digitime (near80/mid100/room) Varlux Computer 3V (large int) Varlux Computer 2V (int and near) Interview80/130 (large near) | Crizal Sapphire 360 UV Crizal Provencia Crizal Rock Crizal Alize UV Crizal Easy UV | Opifog (antiglare & fog both sides) Antifog AR (antiglare front & antifog b | Transitions Signature GENB Transitions XTRActive Polarized (activ Transitions Vantage (variable polarizat Transitions Drivewear (polarized) | Xperio Sun Xperio Polarised Xperio Mirror - Crizal Sun XProtect (ionic shield strengthen) | Stellest (HALT) MyoPiLux SightGlass (DOT) | WAVE & WAVE 2.0 (wavefront advanc Path Optimizer Binocularbooster Flexoptim Nanoptix (anti off-balance) Synchroneyes (calculate both lenses as a pair for smooth far-near refraction) Xtend (extend sharp vision area within arm-length 16-28" or 60-85% add power with ED DualOptix | Eyecode (with Visiooffice automatic device) Fit (frame) AWA (customise 0.01 D scale) 4D (dominant eye fast saccade) NVB (ipad, near visual behaviour) NVA (near vision add power with ED) DualOptix |
| HOYA | Nulux Identity V+ SYNC III (antifatigue) Nulux Trueform (freeform) HiLux Trueform Nulux HiLux Sportive SV EnRoutePro SV (Driving) Nulux Seamless Aspheric (Aphakia) | Hoyalux ID MySelf Hoyalux ID MyStyle V+ (Trueview I) Hoyalux ID LifeStyle 3/3 Hoyalux ID Balansis (integrated double surface FF) Daynamic (full back surface PAL) Amplitude Trueform Amplitude EnRoutePro Progressive(Driving) Sportive Progressive | Hoyalux ID WorkStyleV+ 200,400 Tact200,400 Addpower80 (60cm) | UV control Blue control UV block385/400 | Hi-Vision LongLife Super Hi-Vision Hi-Vision Aqua Hi-Vision Aqua Concave Glare filter | Sensity Sensity Dark Sensity Shine | | MIYOSMART+ (DIMS) | | Trueview I / Lite automatic device |
| NIKON | SEEMAX Infinite (freeform 8 axis, talk MyopSee (DAS) RELAXEE Neo (antifatigue) Lite AS | SEEMAX Ultimate SEEMAX Master SEEMAX Power Presio Master Presio Power Presio W | SOLTES Wide Neo Home&Office Neo DigiLife | Pure blue UV pro Pure blue UV SEECOAT Next Blue SEECOAT Blue UV SEECOAT plus UV | SEECOAT Bright/Next Bright (Augme SEECOAT Drive SEECOAT Next | Transitions XTRActive | Sunstyle Polarshade (polarize) | | | |
| ZEISS | SV Sph SV AS SV Superb + Asia SV Individual (w Digital Lens (antif DriveSafe Individual DriveSafe SV EnergizeMe SV (CL user, antifatique, EnergizeME Digital (0.65D) | Individual/balan EnergizeMe PAL (+0.75-4.00 D) | | | | | | | | |
| RODENSTOCK | Perfalt (Sph) Cosmolit (AS) Perfalt Mono Plus 2 (antifatigue) Perfalt Mono Sport 2 (high wrap) Multigressiv Mono2 + Plus (Back mult Impression Mono2 + Plus (Back mult Impression Mono Sport2 | Progressiv Life Free2 Progressiv PureLift Free2 B.I.G. Multigressiv MyView Impression2 Impression Freesign3 | Progressiv Ergo2 Book/PC/Room Multigressiv Ergo2 Book/PC/Room Impression Ergo2 Book/PC/Room Impression Ergo2 S2 Myxema (color versions) | | X-tra Clean Solitaire Protect P 2 | ColorMatic IQ2 | | | | Impression ST DrEye |
| TOKAI | | | | | | | | | | |

I HAVE NO CONFLICT OF INTERESTS IN THE PRESENTATION

Ophthalmology Subspecialties

Low vision
& Rehabilitation

Ped-Oph
& Strabismologist

Uveitis

Retina

Ocular
Pathologist

Neuro-Oph

Cornea &
Refractive Sx

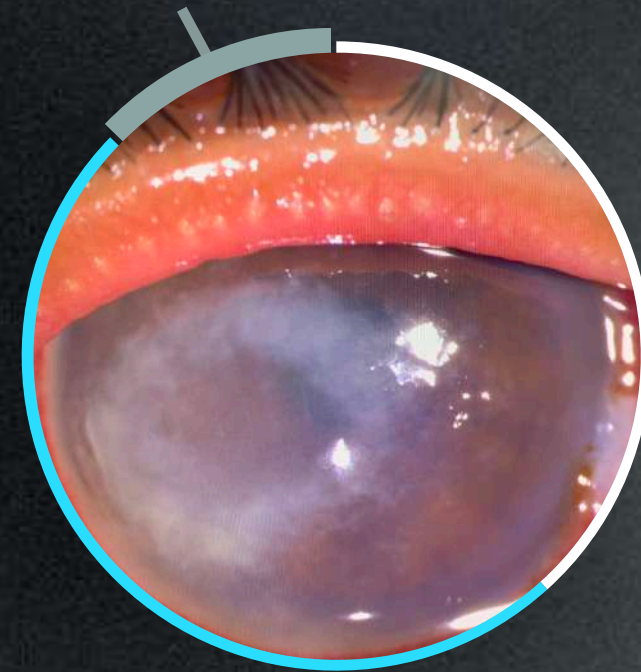
Glaucoma

Oculoplastic

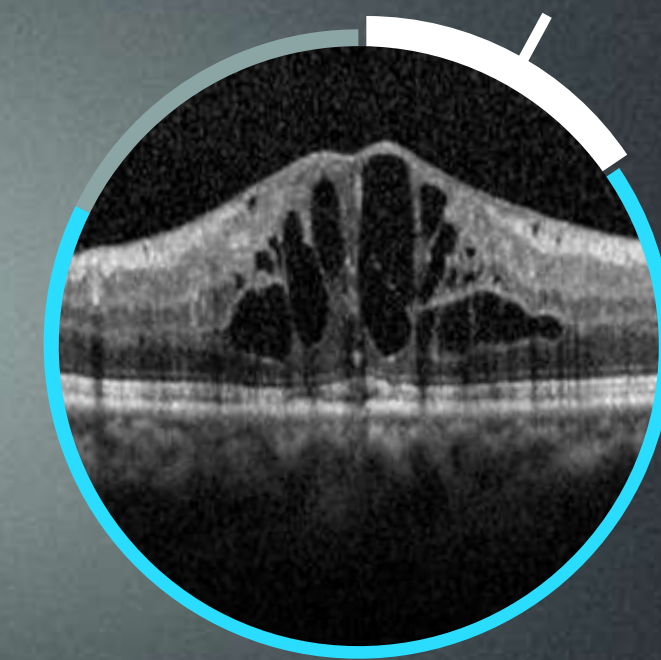


WHAT are common PRESENTATIONS of EYE PROBLEMS?

Ocular Pain & Discomfort



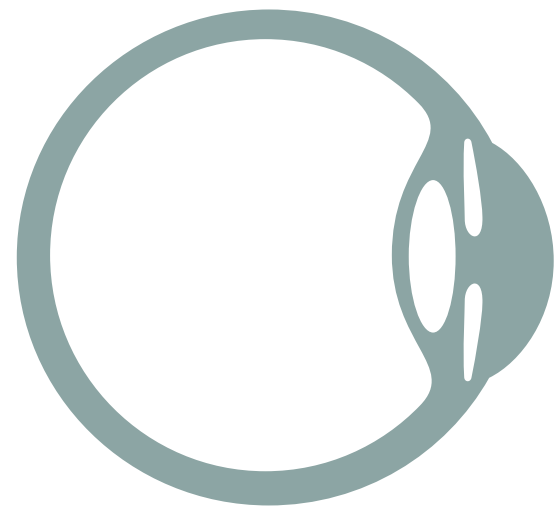
Others: Appearance, Screening



Visual Symptoms



Modalities of Visual Functions



Visual Acuity

Visual Field

Contrast Sensitivity

Color (Ishihara, Fransworth, Red pin)

RAPD (Anterior visual pathway)

Binocular vision (Diplopia, Stereoacuity)

Refractive Error (Defocus)

ERG, VEP, EOG



Image Resolution

Image/Canvas Size

Contrast + Resolution

Hue & Saturation

Exposure/Brightness between 2 images

Merge images > Ghost images, etc

Blur or Sharpen tool

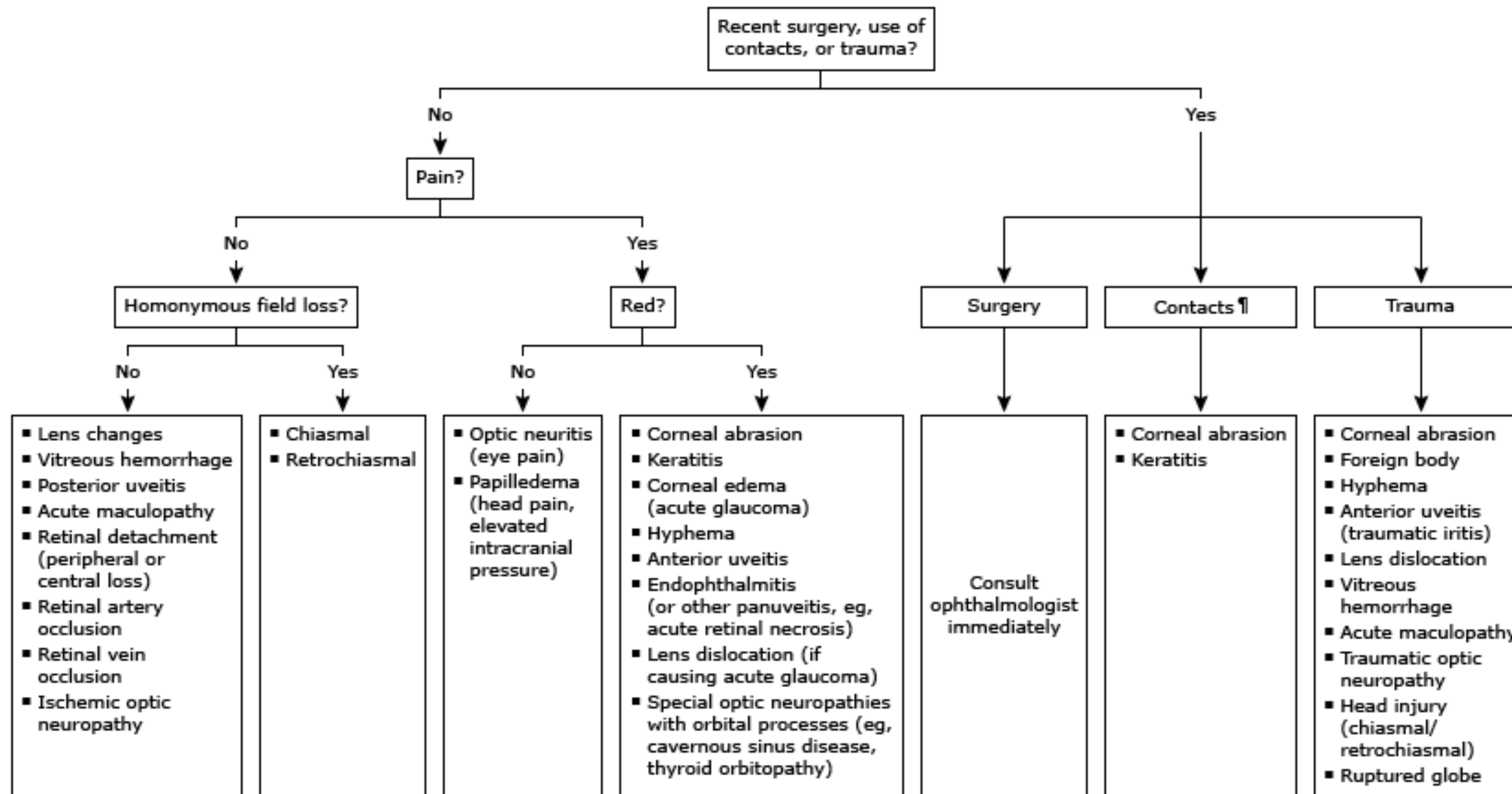
Calibrate for Dead pixel

Approaching Ocular Problems



OUTSIDE IN

Approaching Visual Problems





Approaching Visual Problems

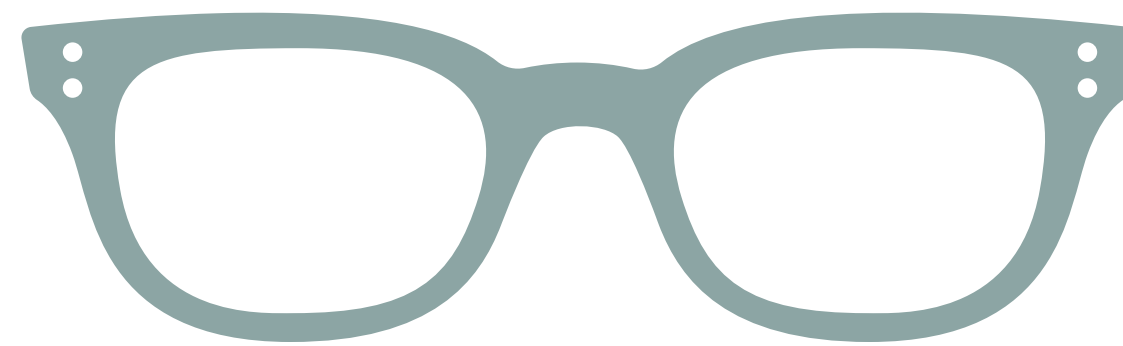
| Refractive Error | Media Opacity | Retina & Choroid | Neural Visual Pathway |
|--|--|--|--|
| Hyperopia | Ptosis | Vasculopathies DR CRAO CRVO Leukemia Hyperviscosity | Pre-chiasmal lesions from ganglion cell layer (GCL) |
| Myopia | Discharge, Tear film instability | Retinal detachment | Optic neuropathies; glaucomatous ischemia, inflammatory, infectious |
| Astigmatism | Cornea; keratitis, scar edema (glaucoma), dystrophy | Maculopathy / Choroidopathy eg. AMD, PCV, CSC, Drug-induced | Disc swelling / Papilledema |
| Others eg. Higher order aberrations (HOA) | Anterior chamber: hyphema (spontaneous or traumatic), uveitis | Retinitis Infection / Inflammation | Chiasmal lesions Pituitary, Craniopharyngioma |
| Ocular misalignment eg. Strabismus | Cataract | Neoplasm | Post-chiasmal lesions Brain lesions eg. stroke, mass, PRES |
| | Vitreous hemorrhage Vitritis, Floaters | Dystrophies eg. Retinitis pigmentosa | Visual Aura, Migraine |

Approaching Visual Problems



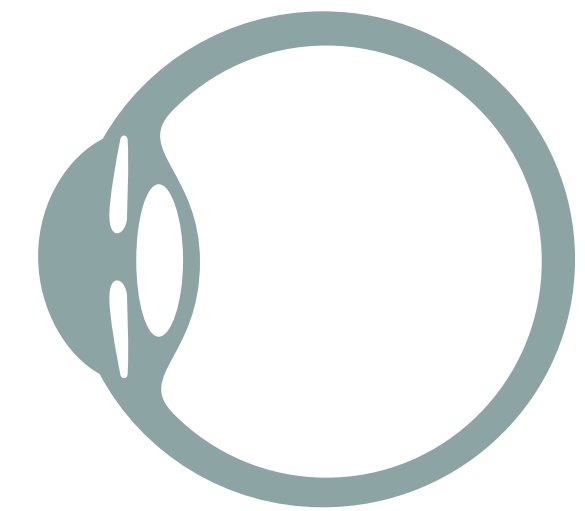
Hx Taking

LODCRAFT
PHx FHx
Trauma/Sx
Prior Tx



Refraction

Pinhole
Spectacles
Contact lens
Day/Night Vision



Eye Exam

Lid, Lash, Conjunctiva,
Tear film, Cornea, A/C, Iris,
Lens, Vitreous, Retina, Optic
nerve, Chiasm, Tract, LGB
to visual cortex, CN III, IV, VI



Approaching Visual Problems

Cataract RE

Hx Taking

70 yr old
Blurry + glare RE
progressed for 6 months
No other symptoms

Refraction

VA RE 20/100 PH 20/70
VA LE 20/40 PH 20/25
Non-significant refractive error

Eye Exam

Conjunctiva, Cornea, A/C WNL
Lens: NS + CC
Fundus: WNL



Approaching Visual Problems

Dry Eyes from Meibomian Gland Dysfunction

Hx Taking

Generalized blurry BE
off and on for years
Worse: outdoor/working
Improve with blink

Refraction

VA - 20/25⁺¹
VA c PH - 20/20⁻¹
Unstable refraction
(astigmatism)

Eye Exam

Lid margin: meibum plugs
Conj: mildly injected
Tear film instability
Cornea: epith. erosions



Approaching Visual Problems

Posterior Vitreous Detachment (PVD) RE

Hx Taking

Floater/flashing RE
for months
Worse: lateral gaze

Refraction

VA 20/20
Emmetropia

Eye Exam

Anterior segment: WNL
Vitreous floaters
Lattice degeneration



Approaching Visual Problems

PVD RE, High Risk Retinal Break > Laser Retinopexy

Hx Taking

Floater/flashing RE
for months
Hx of laser LE
Hx of head trauma

Refraction

VA 20/20
High Myopia

Eye Exam

Anterior segment: WNL
Vitreous floaters
Lattice degeneration



Approaching Visual Problems

VH RE, Suspicious of PVD or Retinal Break

Hx Taking

Floater/flashing RE
for months
Full of floaters today
Last checkup: no U/D

Refraction

VA 20/20⁻²
Emmetropia

Eye Exam

Anterior segment: WNL
Vitreous hemorrhage gr I-II
Normal background fundus



Approaching Visual Problems

RRD, Macula on > Emergency retina repair Sx

Hx Taking

Floaters/flashing RE
for months
Blurry inferior (1 dPTA)
to central vision today

Refraction

VA 20/100 PH 20/70⁻²
VA LE 20/20
Hx prior RE vision = LE

Eye Exam

Anterior segment: WNL
Superior RRD, macula on
(involve superior macula)

Approaching Visual Problems

VH RE, Suspicious of PDR > Consult

Hx Taking

Blurry RE for years
Floaters for days
Can't see anything today
Last visit: 1st Dx DM

Refraction

VA RE HM, LE 20/40
Auto-Refracton: RE
unable, LE emmetropia

Eye Exam

Anterior segment: WNL
Except pupil SRTL BE
RE Vitreous hemorrhage gr IV
(Cannot evaluate fundus)



Approaching Visual Problems

Migraine with Visual aura (Scintillating scotoma)

Hx Taking

Headache after seeing zigzag dancing light followed by blurry image, expand to 50% of visual field in both eyes, lasting 30 min

Refraction

VA 20/20
(no symptom now)
Emmetropia

Eye Exam

WNL



Approaching Visual Problems

Acute Angle Closure RE > IOP lowering agent & LPI

Hx Taking

At 4 AM

Headache N/V 4 hrs

RE blurry & see halo

around lightings

Refraction

VA 20/70

Auto-Refracton: unable

Eye Exam

Tense globe

Ciliary injection

Slightly cloudy cornea

Mid-dilated fixed pupil



Approaching Visual Problems

Optic Neuritis RE

Hx Taking

Headache for 4 day
Blurry RE with
Color desaturation

Refraction

VA RE 20/100 PH NI
VA LE 20/20

Eye Exam

Anterior segment: WNL
Except RAPD RE
no disc swelling RE
Ishihara RE 6/24, LE 24/24

Approaching Visual Problems

Bilat. Optic Neuritis, Suspicious of NMOSD/AntiMOG

Hx Taking

Headache for 1 day
Blurry RE > LE with
Color desaturation

Refraction

VA RE 20/400 PH NI
VA LE 20/50 PH NI

Eye Exam

Anterior segment: WNL
Except RAPD RE
Mild disc swelling RE
Ishihara RE 0/24, LE 0/24



Approaching Visual Problems

Advance glaucoma RE > LE

Hx Taking

70 yr old
Blurry + dark vision RE
progressed for 6 months
No other symptoms

Refraction

VA RE 20/20
VA LE 20/20
Emmetropia

Eye Exam

IOP RE 28 LE 25
Anterior segment: WNL
Except RAPD RE
C:D RE 0.9 LE 0.7



Approaching Visual Problems

Rt Post-chiasmatic lesion, Suspicious of Stroke

Hx Taking

75 yr old, Sudden
painless blurry vision BE
U/D Old CVA, HT, DM
Hx Cataract Sx BE 10 yr

Refraction

VA RE 20/25 PH 20/20
VA LE 20/25 PH 20/20
Nearly emmetropia

Eye Exam

Anterior segment: WNL
RAPD negative
Normal fundus
Confrontation: Lt hemianopia



Approaching Visual Problems

Asthenopia, Presbyopia

Hx Taking

37 yr old
Intermittent blurry vision
periocular dull-aching pain
worsen during E-sport

Refraction

VA RE 20/20
VA LE 20/20
Hyperopia +1 BE

Eye Exam

WNL



Approaching Visual Problems

Monocular Diplopia from Refractive Error

Hx Taking

80 yr old
Diplopia 2 months
Cannot drive his car
U/D: HT DM DLP

Refraction

VA cc RE 20/50 PH 20/25
VA cc LE 20/70 PH 20/30
Hyperopic astigmatism

Eye Exam

Anterior segment WNL
Normal fundus examination
EOM full auctions & versions
APCT orthophoria



Approaching Visual Problems

Bilateral Sixth Nerve Palsies > Workup

Hx Taking

80 yr old
Diplopia 1 day
Cannot drive his car
U/D: HT DM DLP

Refraction

VA (BE open) 20/20
without diplopia
both Snellen & near chart
Emmetropia

Eye Exam

Anterior segment WNL
Normal fundus examination
EOM limit abduction 80 PD BE
APCT ET 30 PD on side gazes



Approaching Visual Problems

Divergence Insufficiency > Prism spectacles

Hx Taking

80 yr old
Diplopia 1 day
Cannot drive his car
Last checkup: no U/D

Refraction

VA (BE open) 20/20
without diplopia
both Snellen & near chart
Emmetropia

Eye Exam

Anterior segment WNL
Normal fundus examination
EOM full ductions & versions
APCT: ET 10 PD, ortho at near

Approaching Red Eye & Discomfort



OUTSIDE IN

Approaching Red Eye & Discomfort

Lid & Lashes

Cellulitis / Panophthalmitis

Hordeolum / Chalazion / Neoplasm

Meibomian Gland Dysfunction

Thyroid orbitopathy

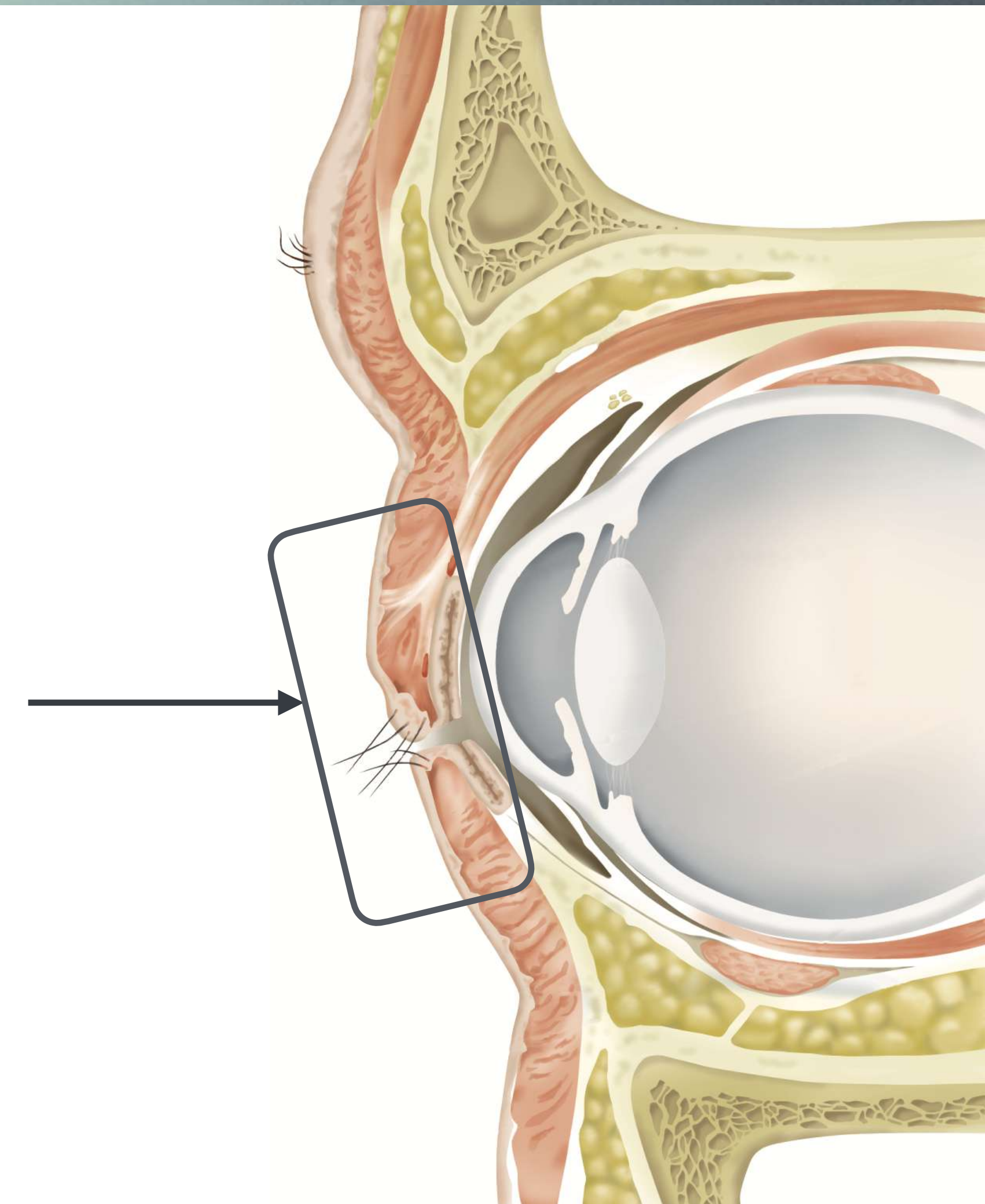
Herpes Simplex / Zoster Blepharitis

Trichiasis / Distichiasis

Ectropion / Entropion / Epiblepharon

Blepharospasm / Hemifacial spasm

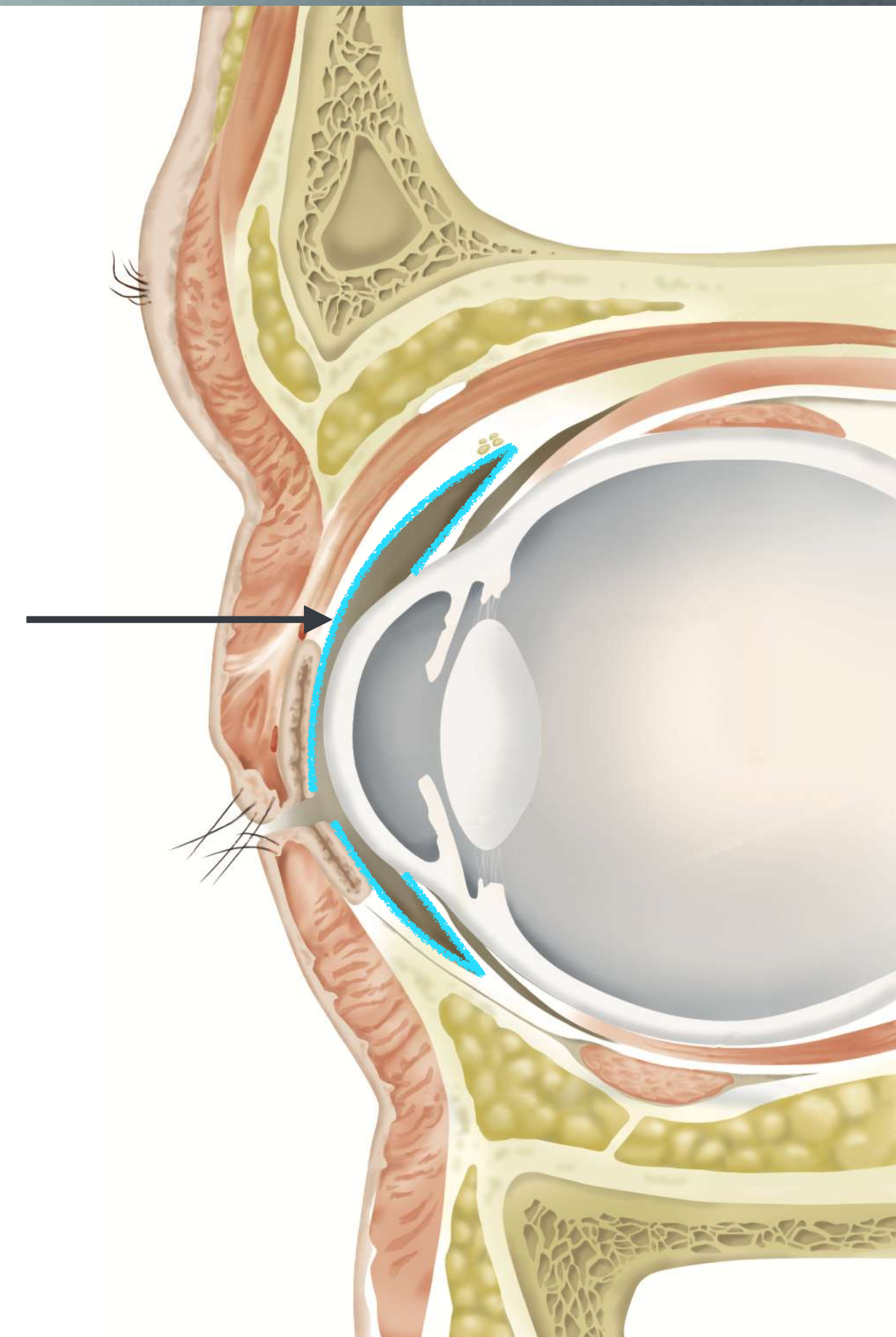
Dacryocystitis / Canaliculitis



Approaching Red Eye & Discomfort

Conjunctiva

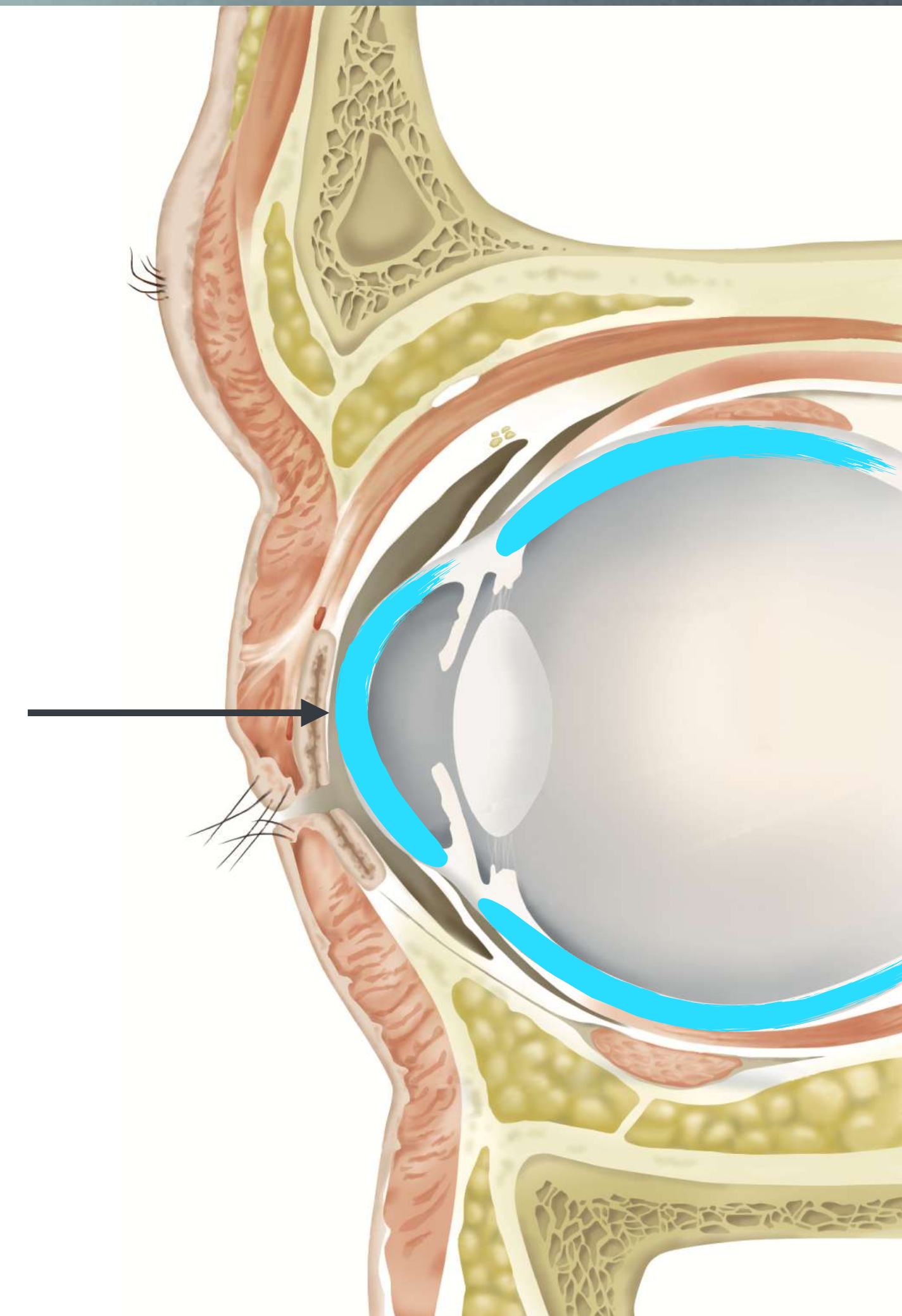
Conjunctivitis / Episcleritis
Infection: Viral, Bacterial, Parasitic
Inflammation / SJS / Allergic
Masquerade / Neoplasm
Thyroid orbitopathy
Subconjunctival Hemorrhage
Foreign body / Lithiasis



Approaching Red Eye & Discomfort

Tear film Cornea & Sclera

- Dry Eyes
- Epithelial defect / Abrasion
- Contact lens overwear
- Scleritis / Keratitis
- Infection / Inflammation
- Foreign body / Neoplasm



Approaching Red Eye & Discomfort

A/C - Iris - Lens Ciliary body

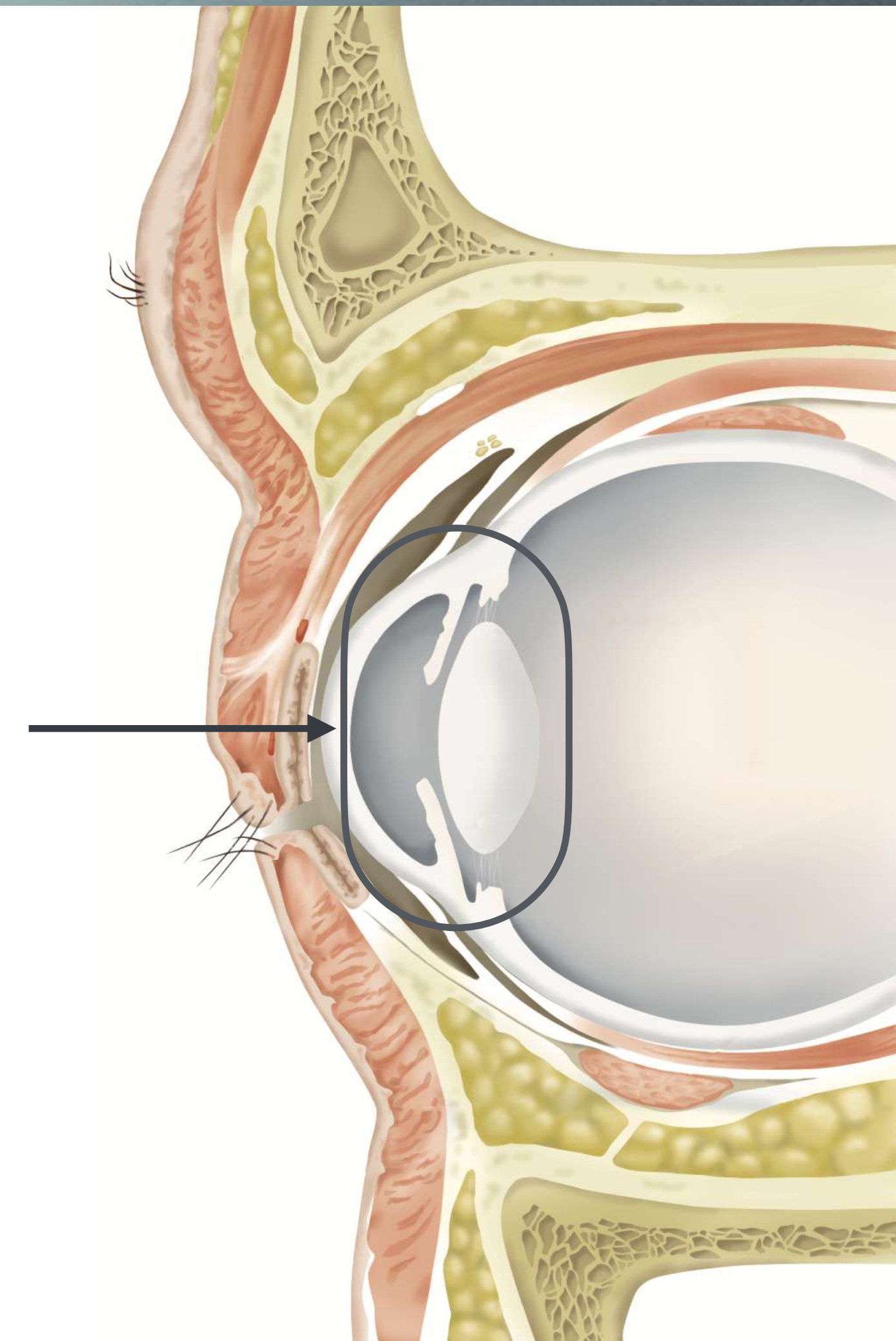
Anterior uveitis (Iritis, Iridocyclitis, Cyclitis)

Angle closure (Primary / Lens-induced)

HypHEMA / Hypopyon

eg. Endophthalmitis

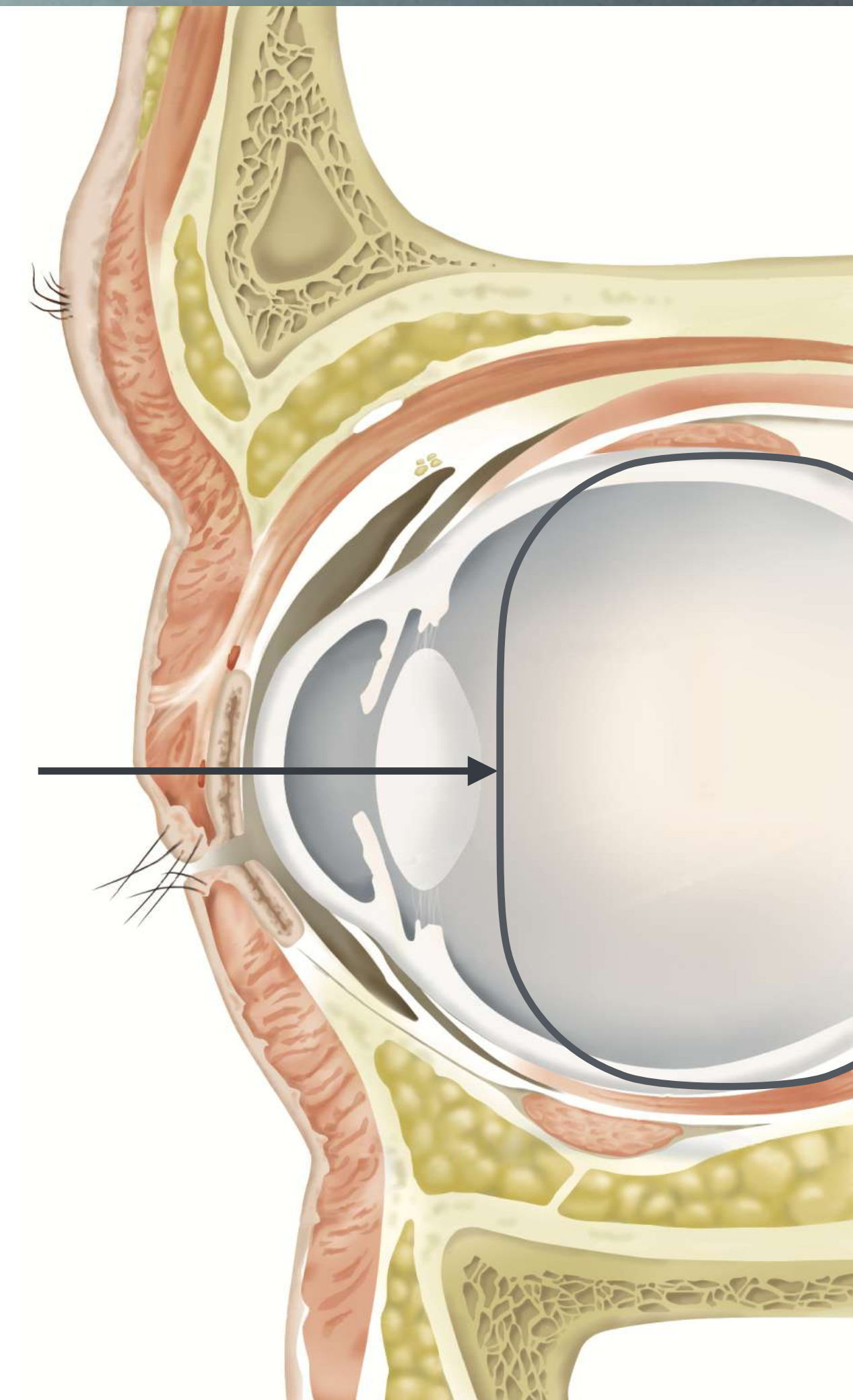
Masquerade



Approaching Red Eye & Discomfort

Posterior Segment

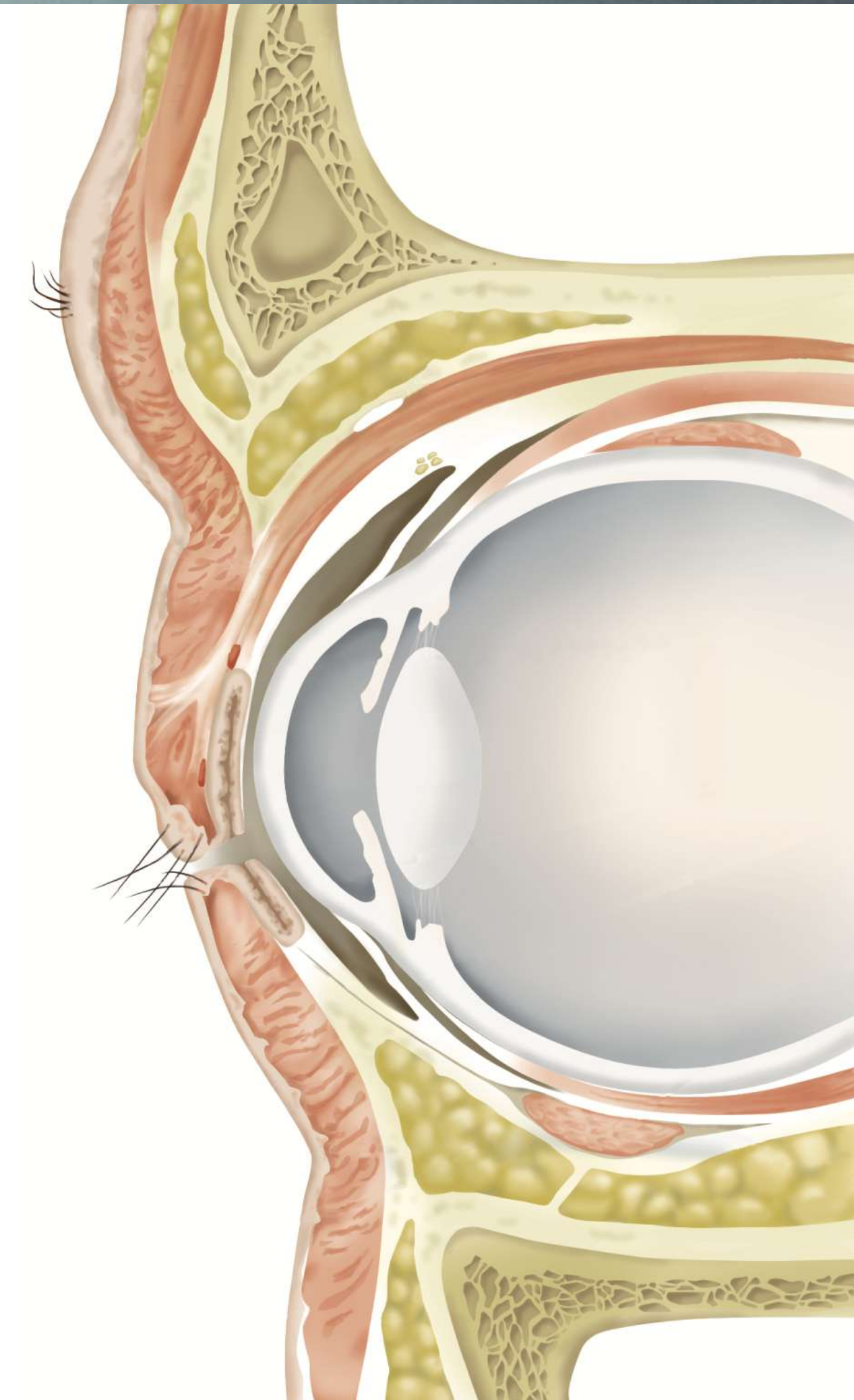
- Vitritis / Endophthalmitis
- Panophthalmitis
- Posterior Scleritis
- Retrobulbar Optic Neuritis



Approaching Red Eye & Discomfort

Asthenopia (Fatigue)

- Excessive near work
- Hyperopia
- Presbyopia
- Overcorrection of Myopia
- Spasm of near reflex
- Strabismus eg. X(T) maintain fusion



Eye Screening

Non-specific

For general patients

Walk in / screening package

Visual functions (VA, Refraction, Colour), IOP
Comprehensive eye exam (may not dilate pupil)

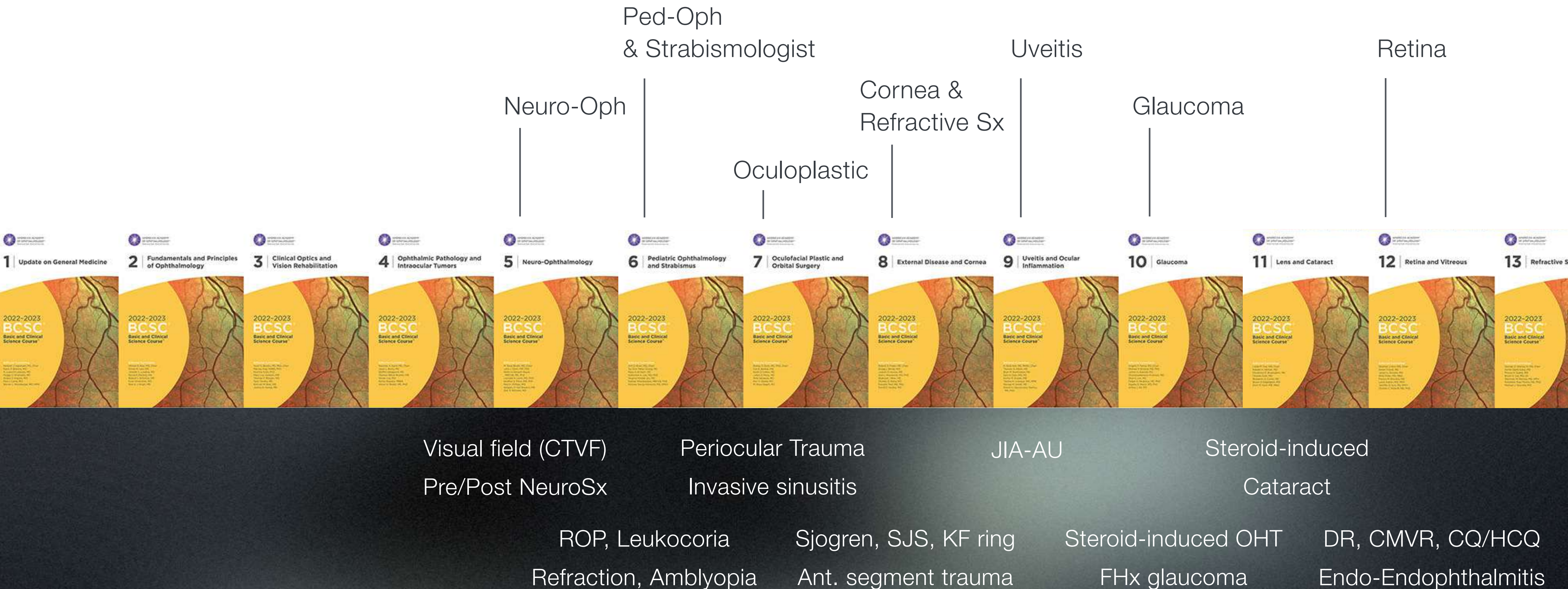
Specific

For increased risk patients

By consultation

Eg. DR, CMVR, HCQ/CQ Maculopathy,
Endogenous endophthalmitis, ROP

Common Specific Screening





TAKE HOME MESSAGE

Hx Taking: LODCRAFT, PH, FHx, Trauma/Sx, Prior Tx

Outside-in Approach: VA cc/pinhole, ant -> post -> visual pathway

Photograph: obtained with patient consent, very useful



THANKS FOR
YOUR ATTENTION



Update management of diabetes mellitus

Preaw Suwannasrisuk, M.D.
Division of Endocrine and metabolism
Department of medicine Naresuan Hospital



Update guideline

- Standard of Care in Diabetes: ADA 2023
- A consensus report by ADA and EASD, Sep 2022
- Thai guideline 2020



Outlines

- Screening and diagnosis
- Lifestyle modification
- Pharmacological management
- Summary

Screening

1. Age \geq 35 years
2. Hx of GDM (testing at least every 3 years)
3. Hx of prediabetes (A1C $>5.7\%$, IGT, IFG) (testing at least yearly)
4. Overweight or obesity (BMI ≥ 23 kg/m²) with one or more risk factor*

Screening

Risk factor*

- First degree relative with diabetes
- History of CVD
- Hypertension
- HDL < 35 mg/dL and/or triglyceride level >250 mg/dL
- Hx PCOS
- Physical inactivity
- Clinical of insulin resistance eg. acanthosis nigrican

Diagnosis

- **FPG \geq 126 mg/dL** . Fasting is defined as no caloric intake for at least 8 h. * or
- **2-h PG \geq 200 mg/dL** during 75 gm-OGTT.*or
- **A1C \geq 6.5%** (A1C method that is NGSP certified and standardized to the DCCT assay.)* or
- **Random plasma glucose \geq 200 mg/dL** + classic symptoms or hyperglycemia crisis

* Diagnosis requires 2 abnormal test results from the same sample or in 2 separate test samples

Assessment and treatment plan

| | |
|--------------------------------------|--|
| Assess risk of diabetes complication | <ul style="list-style-type: none">• ASCVD and heart failure history• ASCVD risk factors and 10-yr ASCVD risk assessment• Staging of CKD• Hypoglycemia risk• Retinopathy and neuropathy |
| Goal setting | <ul style="list-style-type: none">• Set A1C target• Blood pressure target• Diabetes self-management goals |
| Treatment plan | <ul style="list-style-type: none">• Lifestyle management• Pharmacological management (glucose lowering)• Pharmacological management (CVD risk factors and renal) |

ASCVD : atherosclerotic cardiovascular disease



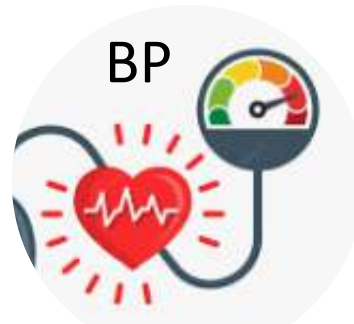
A1C

ADA : A1C < 7%

Premeal CPG 80-130 mg/dL

Peak post prandial < 180 mg/dL

KDIGO: target of <6.5% to <8.0%
(individualized A1C target)



BP

ADA:

- BP < 130/80 mmHg
- BP < 140/90 mmHg if older adult and multiple comorbidities

- **KDIGO2021**: SBP < 120 mmHg

ACEI or ARB : 1 st line treatment



Thai guideline 2017: target LDL

LDL < 100 mg/dL for 1° prevention

LDL < 70 mg/dL for 2° prevention

ADA

Moderate potency statin for 1° prevention

High potency statin for 2° prevention

Kidney Int.2022Sep27:S0085-2538(22)00634-2.

Diabetes care 2022;45(suppl.1):S1895-S207

Thai guideline2017

Treatment goals for glycemia and blood pressure in older adults with diabetes

| Patient character/ health status | A1C goal | Fasting glucose mg/dL | Bedtime glucose mg/dL | Blood pressure mmHg |
|-------------------------------------|---|--------------------------|--------------------------|------------------------|
| Healthy | < 7-7.5 % | 80-130 | 80-180 | <130/80 |
| Complex/ intermediate* | < 8 % | 90-150 | 100-180 | <130/80 |
| Very complex/ poor health** | A1C base on avoid hypoglycemia and symptomatic hyperglycemia | 100-180 | 110-200 | <140/80 |

*Multiple coexisting chronic illness or mild to moderate cognitive impairment or ≥ 2 instrument activity daily living impairments

**End stage chronic illness or moderate to severe cognitive impairment or ≥ 2 activity daily living impairment

Treatment plan

- Lifestyle management
- Pharmacological management (glucose lowering)
- Pharmacological management (CVD risk factors and renal)

Lifestyle modification

| | Adiposity-related diabetes | Diabetes with cardiovascular disease | Isolated hyperglycaemia |
|---|--|--|--------------------------------------|
| Primary pathophysiological driver | Insulin resistance | Atherosclerosis, inflammation | β -cell dysfunction |
| Approximate prevalence* | 40–70% | 20–40% | 10–20% |
| Primary morbidity | Obesity | Cardiovascular disease | Hyperglycaemia |
| Foundational diabetes treatment target | Weight-centric | Cardiocentric | Glucocentric |
| Target | >15% bodyweight loss | Use of proven cardio-protective agents | HbA _{1c} <7% |
| Examples of foundational diabetes treatment | Anti-obesity agents or intervention, GLP1R agonist, SGLT2 inhibitor, metformin | SGLT2 inhibitor, GLP1R agonist (thiazolidinediones) | Sulfonylurea, insulin, GLP1R agonist |
| Secondary treatment targets | Glucose, blood pressure, lipids | Weight, glucose, blood pressure, lipids, coagulation | NA |

HbA_{1c}=glycated haemoglobin. NA=not applicable. *Prevalence varies by definition and population.

Table 3: Proposed primary and secondary treatment goals for type 2 diabetes by prevailing disease phenotype

Weight-centric approach

Cardiocentric approach

Glucocentric approach

Treatment effectiveness of weight loss

| % weight loss | Procedure |
|---------------|---|
| 5-7% | <ul style="list-style-type: none">• Self monitor diet• Intensive lifestyle program |
| 8-15% | <ul style="list-style-type: none">• Meal replacement• Very low calories diet |
| 9-15% | <ul style="list-style-type: none">• Weight-loss medications• GI procedure |
| 20-30 % | <ul style="list-style-type: none">• SLEEVE gastrectomy• RYGB |

Lifestyle modification

- $\geq 5\%$ weight loss are recommended for most people with type 2 diabetes and overweight or obesity.
- Method of 3-5% weight loss:
 - 500–750 kcal/day energy deficit or
 - calorie restriction : 1,200 –1,500 kcal/day for women
 - : 1,500 –1,800 kcal/day for men

History taking

- Medication
- Hypoglycemia
- Meal: frequency, CHO count
- Simple CHO : sweetener drinks, bakery, Thai dessert





บวมจีน 1 จับ หรือ 1/2 ถ้วยตวง



ข้าวเหนียว 1/2 ถ้วยตวง หรือ 1/4 ถ้วยตวง



ข้าวขาว 1 ถ้วยตวง หรือ 1/3 ถ้วยตวง

| | | |
|--------------|---------------|------------------|
| คาร์โบไฮเดรต | 18 กรัม | 1 ส่วน = 1 คาร์บ |
| โปรตีน | 2 กรัม | |
| พลังงาน | 80 กิโลแคลอรี | |

หมวดข้าวแป้ง และผลิตภัณฑ์

คาร์โบไฮเดรต

18 กรัม

โปรตีน

2 กรัม

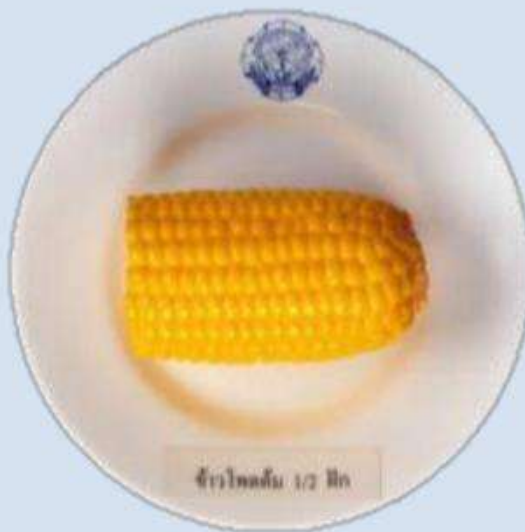
พลังงาน

80 กิโลแคลอรี

1 ส่วน = 1 คาร์บ



ขนมปัง
1 แผ่น



ข้าวโพดต้ม
1/2 ฝัก



ขนมปังแครกเกอร์
4 - 6 แผ่น

หมวดผัก ก.
(ผักที่ไม่มีแป้ง)

ไม่คิดพลังงาน อุดมด้วยแร่ธาตุ วิตามิน
และใยอาหาร



ผักกาดขาว



แตงกวา



ตำลึง



ผักบุ้งจีน



กะหล่ำปลี



ผักกวางตุ้ง

หมวดผัก ข. (ผักที่มีแป้ง)

| | |
|--------------|---------------|
| คาร์โบไฮเดรต | 5 กรัม |
| โปรตีน | 2 กรัม |
| พลังงาน | 28 กิโลแคลอรี |

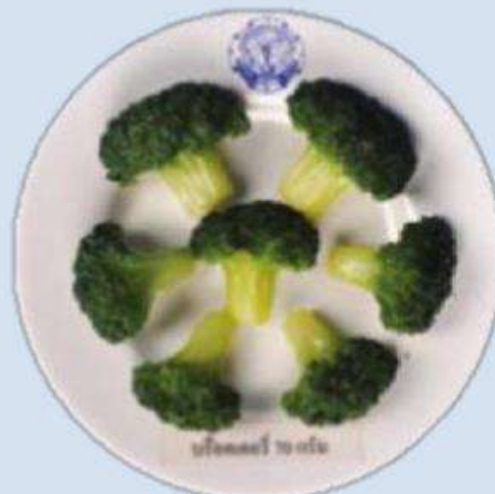
1 ส่วน = 0.3 คาร์บ



แครอท



ผักคะน้า



บร็อคเคอรี่



ฟักทอง



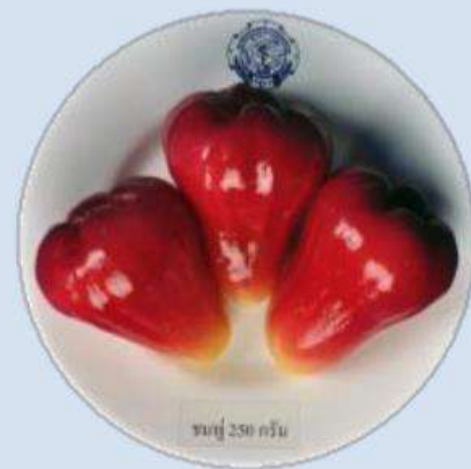
ฝรั่ง



แอปเปิ้ล



มะละกอ



ชมพู่

ผลไม้ วันละ 3-4 ส่วน

Plate model

- มืออาหารหลักของฉัน -



กินให้ถูกส่วนใน 1 มือ

2 : 1 : 1
(ผัก) (ข้าว/แป้ง) (เนื้อสัตว์)



Case scenario

- Goal of treatment
- Diabetes management

Case 1: 60-year-old woman

Underlying diseases: T₂DM for 2 years

Physical examination

- V/S: BP 140/90 mmHg, HR 80 bpm, BMI 25 kg/m²
- Acanthosis nigrican at neck
- RS and CVS: unremarkable
- Ext: no pitting edema

Case1 : 60-year-old woman: T2DM for 2 years

Current medication

Metformin 2,000mg/day

Physical examination

- V/S: BP 120/80 mmHg
- BMI 30 kg/m²

Laboratory

HbA1c=8.5%, FPG=180mg%, LDL=80mg/dL
Cr=0.8mg/dL
Urine microalbumin=20mg/gm.cr

Management ?

Update on Thai DM CPG August 2563

2. การเริ่มต้นให้การรักษารักษาขึ้นอยู่กับ

2.1 ระดับน้ำตาลในเลือด และ A_{1c} (ถ้ามีผลการตรวจ)

2.2 อาการหรือความรุนแรงของโรค (อาการแสดงของโรคเบาหวานและโรคแทรกซ้อน)

2.3 สภาพร่างกายของผู้ป่วย ได้แก่ โรคอ้วน โรคอื่น ๆ ที่อาจมีส่วนร่วมด้วย การทำงานของตับและไต

2.4 โรคร่วมของผู้ป่วย ได้แก่ โรคหัวใจและหลอดเลือด และ/หรือ โรคไตเรื้อรัง (GFR < 60 มล/นาที

และ/หรืออัลบูมินในปัสสาวะ ≥ 300 มก/ก)



Case 1: 60-year-old woman: T2DM for 2 years

Current medication

Metformin 2,000 mg/day

Physical examination

- V/S: BP 140/90 mmHg
- BMI 30 kg/m²

Laboratory

HbA1c 8.5%, FPG 180 mg%, LDL 80 mg/dL
Cr=0.8, Urine microalbumin 20mg/gm.cr

Add glipizide or
pioglitazone

Case 2: 60-year-old woman

Underlying diseases: T₂DM for 20 years, hypertension, DLD

Physical examination

- V/S: BP 140/90 mmHg, HR 80 bpm, BMI 30 kg/m²
- Acanthosis nigrican at neck
- RS and CVS: unremarkable
- Ext: pitting edema¹⁺

Case 2: 60-year-old woman

Underlying disease

T₂DM for 20 years, hypertension, DLD

Physical examination

- V/S: BP_{140/90} mmHg

- BMI 30 kg/m²

Laboratory

HbA_{1c} 8.5%, FPG 180 mg%, LDL 120 mg/dL

Cr=1.8, **eGFR=40** ml/min/1.73m³ (persistent)

Urine microalbumin 2,000 mg/gm.cr

Current medication

- Metformin 2,000 mg/day
- Glipizide 10 mg/day
- Pioglitazone 30 mg/day
- Sitagliptin 100 mg/day
- Amlodipine 10 mg/day
- Simvastatin 10 mg/day

F 60 yr

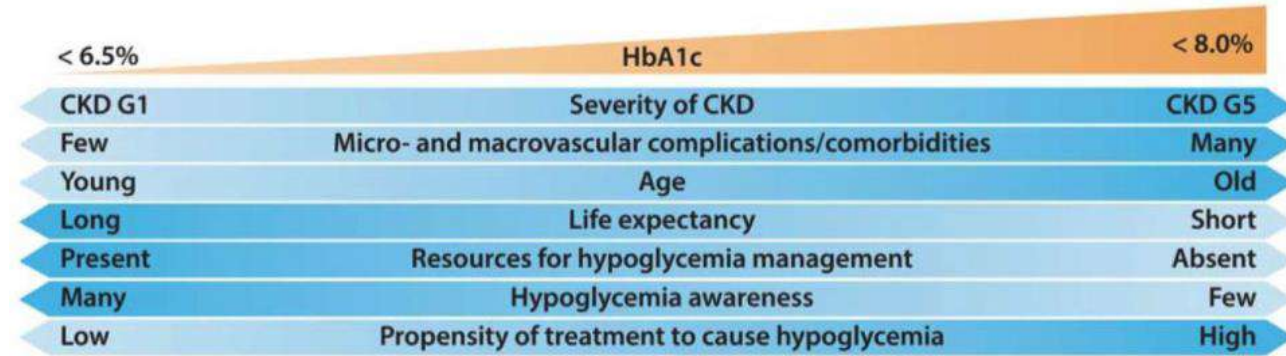
:T₂DM for 20 years, hypertension, DLD

Physical examination

- V/S: BP 140/90 mmHg
- BMI 30 kg/m²

Laboratory

HbA_{1c} 8.5% , FPG 180 mg%,
LDL 120 mg/dL
Cr=1.8, eGFR 40 ml/min/1.73m³
Urine microalbumin 2,000 mg/gm.cr



ADA : target of HbA_{1c} <7%

KDIGO : target of <6.5% to <8.0%
(individualized HbA_{1c} target)

Opinion: target of HbA_{1c} 7-8 %

F 60 yr

:T₂DM for 20 years, hypertension, DLD

Physical examination

- V/S: BP 140/90 mmHg
- BMI 30 kg/m²

Laboratory

LDL 120 mg/dL, cholesterol 250 mg/dl,
HDL 40 mg/dl
Cr=1.8, **eGFR=40** ml/min/1.73m³
Urine microalbumin 2,000 mg/gm.cr



ACEI or ARB : 1 st line treatment

ADA:

- BP < 140/90 mmHg if 10-yr ASCVD risk <15%
- BP <130/80 mmHg if 10-yr ASCVD risk >15%

KDIGO 2021: SBP<120 mmHg

Opinion <130/80 mmHg (Thai guideline2017)

F 60 yr

:T2DM for 20 years, hypertension, DLD

Physical examination

- V/S: BP140/90 mmHg
- BMI 30 kg/m²

Laboratory

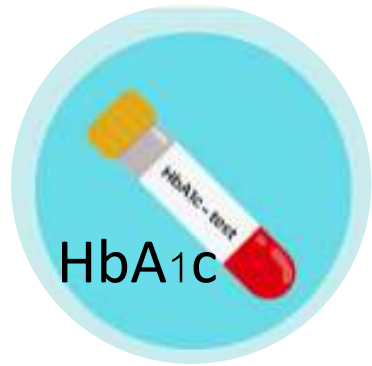
LDL 120 mg/dL , cholesterol 250 mg/dl,
HDL 40 mg/dl
Cr=1.8, **eGFR=40** ml/min/1.73m³
Urine microalbumin 2,000 mg/gm.cr



Target : LDL < 100 mg/dl

- Primary prevention: moderate intensity statin
- Secondary prevention: high intensity statin

| | High Intensity | Moderate Intensity | Low Intensity |
|-----------------|---|---|--|
| LDL-C lowering† | ≥50% | 30%-49% | <30% |
| Statins | Atorvastatin (40 mg‡) 80 mg Rosuvastatin 20 mg (40 mg) | Atorvastatin 10 mg (20 mg) Rosuvastatin (5 mg) 10 mg Simvastatin 20-40 mg§ | Simvastatin 10 mg |
| ... | ... | Pravastatin 40 mg (80 mg) Lovastatin 40 mg (80 mg) Fluvastatin XL 80 mg Fluvastatin 40 mg BID Pitavastatin 1-4 mg | Pravastatin 10-20 mg Lovastatin 20 mg Fluvastatin 20-40 mg |



ADA : target of HbA1c <7%

KDIGO : target of <6.5% to<8.0
(individualized HbA1c target)



ADA:

- BP < 140/90 mmHg
if 10-yr ASCVD risk < 15%
- BP < 130/80 mmHg
if 10-yr ASCVD risk > 15%

KDIGO 2021: SBP < 120 mmHg

ACEI or ARB : 1 st line treatment



Thai guideline 2017: Target LDL
LDL < 100 mg/dl for 1° prevention
LDL < 70 mg/dl for 2° prevention

ADA

1° prevention : moderate potency statin
2° prevention : high potency statin

Diabetes and CKD management



| | ADA 2022 | KDIGO 2022 |
|-------------------|---|--|
| Protein | | 0.8 gm protein/kg/day |
| Sodium | <2,300 mg/d | <2,000 mg/d |
| Physical activity | Moderate intensity (≥ 150 min/wk) Vigorous intensity (≥ 75 min/wk) | Moderate intensity (at least 150 min/wk) |
| Weight | At least 5% weight loss | Advice weight loss in CKD with obese (eGFR ≥ 30) Consider GLP-1 agonist for promote weight loss |
| Alcohol | ≤ 1 drink in women ≤ 2 drinks in men | No recommendation on alcohol intake |

Diabetes and CKD management

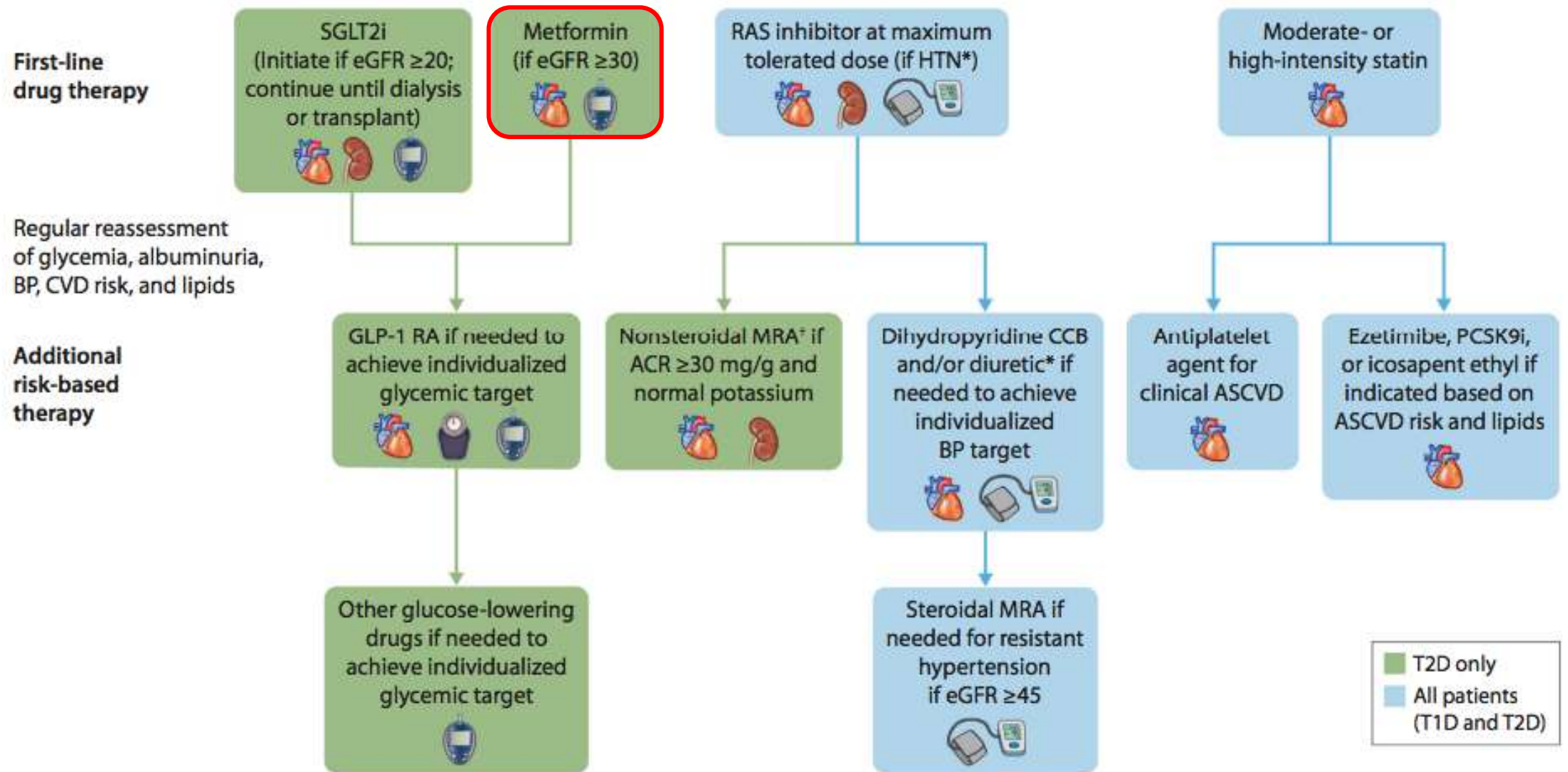


Table 2 | Considerations for selecting glucose-lowering agents in patients with T2D and CKD^{2,17}

| | Progression of CKD | ASCVD | Heart failure | Glucose-lowering efficacy | Hypoglycemia risk | Weight effects | Cost |
|--------------------------------|----------------------|----------------------------------|---|---------------------------|-------------------|----------------|------------------|
| Metformin | Neutral | Potential benefit | Potential benefit | High | Low | Neutral | Low |
| SGLT2 inhibitors | Benefit ^a | Benefit ^c | Benefit | Intermediate | Low | Loss | High |
| GLP-1 receptor agonists | Benefit ^b | Benefit ^c | Potential benefit | High | Low | Loss | High |
| DPP-4 inhibitors | Neutral | Neutral | Potential risk ^c (saxagliptin) | Intermediate | Low | Neutral | High |
| Insulin | Neutral | Neutral | Neutral | Highest | High | Gain | High (analogues) |
| | | | | | | | Low (human) |
| Sulfonylureas | Neutral | Neutral | Neutral | High | High | Gain | Low |
| Thiazolidinediones | Neutral | Potential benefit (pioglitazone) | Increased risk | High | Low | Gain | Low |
| α-Glucosidase inhibitors | Neutral | Neutral | Neutral | Intermediate | Low | Neutral | Low |

Neutral

Potential benefit or intermediate glucose-lowering efficacy

Benefit (organ protection, high efficacy, low hypoglycemia risk, weight loss, or low cost)

Potential risk or high cost to patient

Increased risk for adverse effects

Metformin for T₂DM with CKD

- eGFR 45-59 ml/min/1.73m³ : a reduction should be considered
- eGFR 30-44 ml/min/1.73m³: max dose 1,000 mg/day
- eGFR <30 ml/min/1.73m³ : contraindication

- Sick day protocol: **holding metformin** doses during acute illness.

SGLT2 inhibitors (SGLT2i)

| Benefit | Side effect |
|--|---|
| Decrease A1C 0.5-0.9 % Low risk hypoglycemia Weight loss 2 kg SBP lowering 2.5-5 mmHg DBP lowering 1-2 mmHg Cardiovascular and renal protection | Volume depletion from polyuria Fungal genital infection (F 10%, M 2-3%) DKA (<0.1%) |

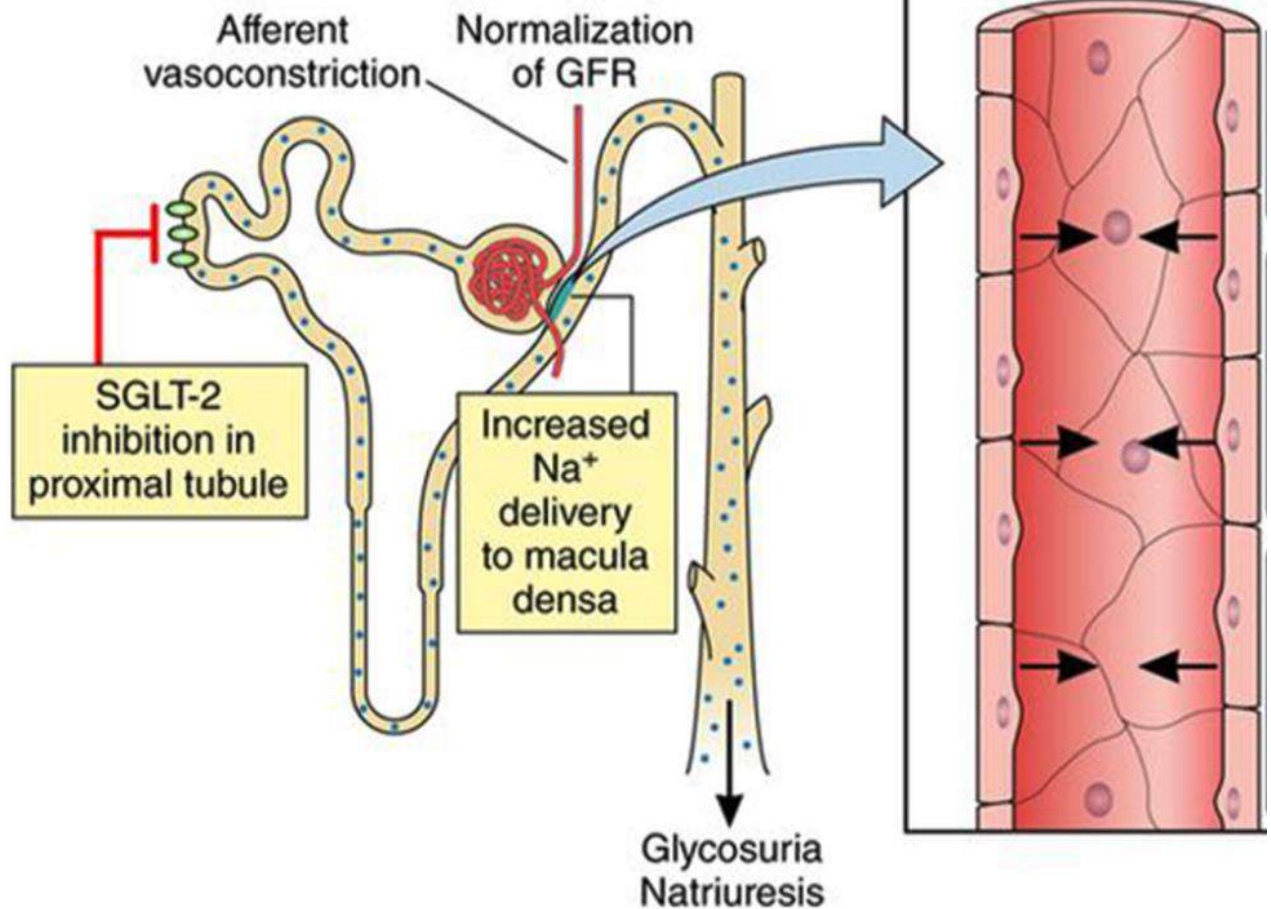
Table 4—Dose adjustments for eGFR <45 mL/min/1.73 m² (information presented reflects the package inserts rather than guidance from this consensus report)

| | Stage 3b (eGFR 30–44 mL/min/1.73 m ²) | Stage 4 (eGFR 15–29 mL/min/1.73 m ²) | Stage 5 (eGFR <15 mL/min/1.73 m ²) |
|---------------|--|---|--|
| Canagliflozin | Maximum 100 mg daily | Initiation not recommended; may continue 100 mg daily if tolerated for kidney and CV benefit until dialysis | |
| Dapagliflozin | 10 mg daily [†] | | Initiation not recommended with eGFR <25 mL/min/1.73 m ² ; may continue if tolerated for kidney and CV benefit until dialysis |
| Empagliflozin | 10 mg daily [†] | | Initiation not recommended with eGFR <20 mL/min/1.73 m ² ; may continue if tolerated for kidney and CV benefit until dialysis |

| | CREDENCE(N=4401) | DAPA-CKD (N=4094) | EMPA-KIDNEY(N=6609) |
|--|--|---|--|
| Product | Canagliflozin | Dapagliflozin | Empagliflozin |
| Patient population | CKD+T2DM | CKD+T2DM CKD without T2DM | CKD+T2DM CKD without T2DM |
| Required eGFR(ml/min/1.73m ³) and UACR(mg/d) for enrolment | eGFR ≥ 30 -<90 UACR:>300 - ≤ 500 | eGFR ≥ 25 -<75 UACR: ≥ 200 - ≤ 5000 | eGFR >45-<90 UACR:>200 or eGFR ≥ 20 - <45 |
| Primary endpoint | Composite of ESKD, doubling of serum Cr, renal or CV death | Composite of $\geq 50\%$ susptanin decline in eGFR, ESKD, renal or CV death | Kidney disease progression (ESKD, susptanin decline in eGFR to < 10 ml/min/1.73m ³ , renal death or sustain decline of $\geq 40\%$ in eGFR) or CV death |
| Outcome | HR 0.7(0.59-0.82), P=0.00001 | HR 0.61(0.51-0.72), P < 0.001 | HR 0.72(0.64-0.82), P<0.000001 |

N Engl J Med.2019 Jun13;380(24):2295-2306
 N Engl J Med. 2020 Oct 8;383(15):1436-1446.
 N Engl J Med.2022 Nov 4. doi: 10.1056/NEJMoa2204233.

C SGLT-2 inhibition reduces hyperfiltration via TGF



Tubuloglomerular feedback is mediated by the juxtaglomerular apparatus, which contains the macula densa; a specialised group of cells that detect sodium ion (Na^+) concentration within the tubule, signalling to the glomerulus to regulate the filtration rate and avoid dehydration via a feedback loop.

Improved glomerular haemodynamics

- decreased proximal tubular sodium resorption
- glomerular afferent arteriolar vasoconstriction (in response to raised adenosine levels, driven by increased membrane Na^+/K^+ ATPase activity)

Outcome:

intraglomerular pressure and reduces the amount of protein filtered through the glomerulus (albuminuria).

GLP-1 receptor agonists

| Benefit | Side effect |
|---|--|
| Decrease A1C 0.9-2.2% Low risk hypoglycemia Weight loss 1.3-8.7 kg BP lowering 2-3 mmHg Cardiovascular and renal protection | GI side effect Nausea 25-60% Vomiting 5-15% (risk of cholestasis, pancreatitis) |



Liraglutide (once daily)



Dulaglutide (weekly)

Table 4—Dose adjustments for eGFR <45 mL/min/1.73 m² (information presented reflects the package inserts rather than guidance from this consensus report)

| | Stage 3b (eGFR 30–44 mL/min/1.73 m ²) | Stage 4 (eGFR 15–29 mL/min/1.73 m ²) | Stage 5 (eGFR <15 mL/min/1.73 m ²) |
|--|--|---|---|
| GLP-1 receptor agonists[§] | | | |
| Exenatide | Caution initiating or increasing dose; avoid once-weekly formulation | Use not recommended | |
| Dulaglutide | No dose adjustment required | | |
| Liraglutide | No dose adjustment required | | |
| Lixisenatide | No dose adjustment required | | Use not recommended |
| Semaglutide | No dose adjustment required | | |

Case 1: 60-year-old woman

Dx

1. T2DM with poor control with DN
2. CKD G3b A3
3. HT, DLD, obesity

Underlying disease

- T2DM for 20 years, hypertension, DLD
- V/S: BP=140/90 mmHg
 - BMI 30 kg/m², pitting edema 1+

Previous lab (last month)

HbA1c 8.5%, FPG 180 mg%, LDL 120 mg/dL
Cr=1.8, eGFR 40 ml/min/1.73m² (persistent)
Urine microalbumin 2,000 mg/gm.cr

คำแนะนำในการปรับยา

1. ลดขนาด meformin 1,000 mg/day (eGFR 30-44 ml/min/1.73m²)
2. Glipizide คงขนาดเดิมได้ถ้าไม่มีอาการ hypoglycemia
3. ควร off Pioglitazone เนื่องจากเริ่มบวม
4. Sitagliptin ควรลดขนาดยาเป็น 50 mg/day (eGFR 30-44 ml/min/1.73m²)
5. ยาลดความดันควรเพิ่มยาในกลุ่ม ACEI หรือ ARB (low dose)
ติดตาม home BP และค่า Cr, K
6. ควรเพิ่มขนาดยา simvastatin 20 mg/day
หรือปรับเป็น atorvastatin 10-20 mg/day
7. หากต้องเพิ่มยาลดน้ำตาล โดยไม่มีข้อจำกัดเรื่องค่าใช้จ่าย แนะนำ
SGLT2 inh หรือ
GLP1 agonist (หากเริ่มยาในกลุ่มนี้ต้องงดยา DPP4-inh; sitagliptin)



Anti diabetic agent dose adjustment for CKD

| Medication | eGFR | Dose |
|--------------------------------------|--------------------|--------------------|
| Sulfonylureas(2nd generation) | | |
| Glimepiride | Stage 3b-5 | 1-8 mg/d |
| Glipizide | Stage 3b-5 | 2.5 -20 mg/d |
| Thiazolidinediones | | |
| Pioglitazone | No dose adjustment | |
| Alpha-Glucosidase inhibitors | | |
| Acarbose | Stage 3b (30-44) | No dose adjustment |
| | Stage 4-5 | Use not recommend |

SGLT2i inhibitors

| SGLT2i In Thailand: | (T2DM indication) eGFR (ml/min/1.73m ²) | Dose (mg/day) |
|---------------------|--|---------------|
| Canagliflozin | <ul style="list-style-type: none"> ○ EGFR ≥ 30 with UACR > 300 mg/g ○ eGFR >45 | ○ 100 |
| | ○ eGFR >60 | ○ 300 |
| Dapagliflozin | ○ eGFR ≥ 45 | ○ 10 |
| Empagliflozin | ○ eGFR ≥ 30 | ○ 10-25 |
| Lusegliflozin | ○ eGFR ≥ 60 | ○ 5 |

| SGLT2i In Thailand: | (Heart failure indication) eGFR (ml/min/1.73m ²) | Dose(mg/day) |
|---------------------|---|--------------|
| Canagliflozin | ○ Not approved | |
| Dapagliflozin | ○ eGFR ≥ 25 | ○ 10 |
| Empagliflozin | ○ eGFR ≥ 20 | ○ 10 |

| SGLT2i In Thailand: | (CKD: Indication) eGFR(ml/min/1.73m ²) | Dose (mg/day) |
|---------------------|--|---------------|
| Canagliflozin | <ul style="list-style-type: none"> ○ EGFR ≥ 30 ○ with UACR > 300 mg/g | ○ 100 |
| Dapagliflozin | ○ eGFR ≥ 25 | ○ 10 |
| Empagliflozin | ○ eGFR ≥ 30 | ○ 10 |

Kidney Int2022. Sep 27:S0085-2538(22)00634-2.

DPP4 inhibitors

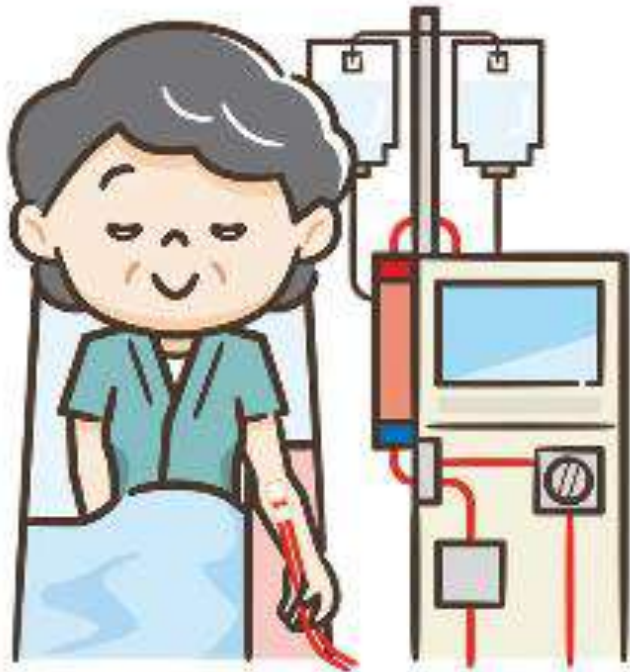
| Medication | eGFR | Dose |
|-------------|--------------------------|------------|
| Sitagliptin | Stage 1-3a (≥ 45) | 100 mg/d |
| | Stage 3b (30-44) | 50 mg/d |
| | Stage 4-5 | 25 mg/d |
| Gemigliptin | No dose adjustment | |
| Linagliptin | No dose adjustment | |
| Saxagliptin | Stage 1-3a (≥ 45) | 2.5-5 mg/d |
| | Stage 3b-5 | 2.5 mg/d |

GLP1 agonist

| Medication | eGFR | Dose | Thai FDA |
|-------------|--------------------|-----------------|-----------|
| Liraglutide | No dose adjustment | 0.6-1.8mg sc OD | eGFR >15 |
| Dulaglutide | No dose adjustment | 1.5mg sc weekly | eGFR > 15 |

Case 3:
60-year-old woman

- T2DM for 20 years, hypertension, DLD
- **Symptomatic hyperglycemia**
- BW 60 kg



Previous Lab

HbA1c=**13%**, FPG 240 mg%, LDL 80 mg/dL

Cr=1.8 , eGFR 40 ml/min/1.73m² (persistent)

UACR 2,000 mg/gm

Medication: glipizide 10 mg/day

Add basal insulin³

Choice of basal insulin should be based on person-specific considerations, including cost. Refer to Table 9.4 for insulin cost information. Consider prescription of glucagon for emergent hypoglycemia.

Add basal analog or bedtime NPH insulin⁴

INITIATION: Start 10 units per day OR 0.1–0.2 units/kg per day

TITRATION:

- Set FPG target (see Section 6, “Glycemic Targets”)
- Choose evidence-based titration algorithm, e.g., increase 2 units every 3 days to reach FPG target without hypoglycemia
- For hypoglycemia determine cause, if no clear reason lower dose by 10–20%

Assess adequacy of basal insulin dose

Consider clinical signals to evaluate for overbasalization and need to consider adjunctive therapies (e.g., basal dose more than ~0.5 units/kg/day, elevated bedtime–morning and/or post–preprandial differential, hypoglycemia [aware or unaware], high variability)

Neutral Protamine Hagedorn (NPH)



| ชนิดยา (ชื่อยา) | เวลาที่เริ่มออกฤทธิ์ | เวลาที่มีฤทธิ์สูงสุด | ระยะเวลาการออกฤทธิ์ |
|---|----------------------|----------------------|---------------------|
| อินซูลินออกฤทธิ์ปานกลาง (Insulin Isophane Suspension, NPH) (Insulatard HM [®] , Humulin N [®] , Gensulin N [®] , Insugen N [®] , Insuman basal [®] , Winsulin N [®]) | 2-4 ชั่วโมง | 4-8 ชั่วโมง | 10-16 ชั่วโมง |



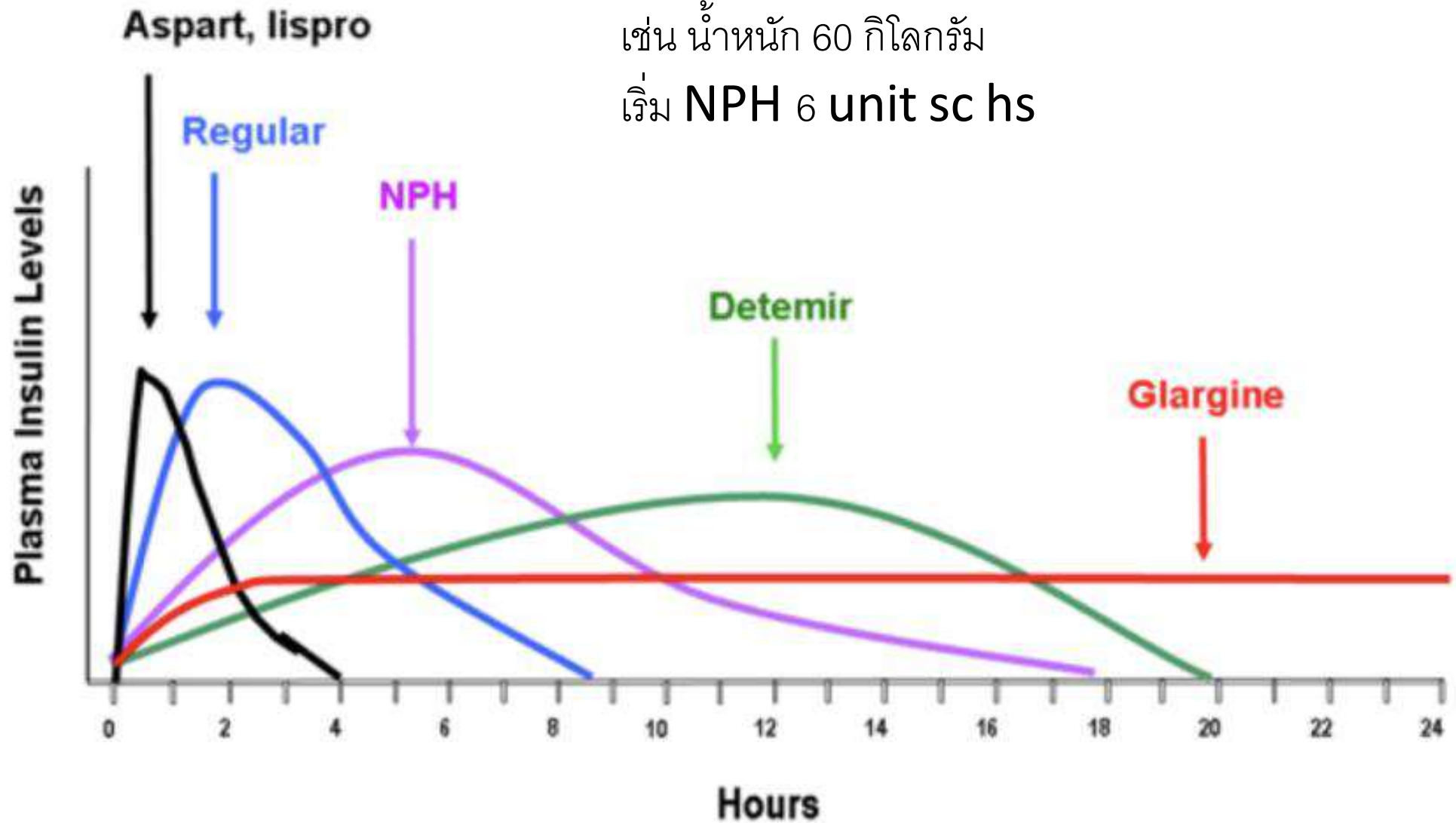
ตารางที่ 2. แสดงยาฉีดอินซูลินชนิดต่างๆ ที่มีในประเทศไทย และเวลาการออกฤทธิ์

| ชนิดยา (ชื่อยา) | เวลาที่เริ่มออกฤทธิ์ | เวลาที่มีฤทธิ์สูงสุด | ระยะเวลาการออกฤทธิ์ |
|--|----------------------|----------------------|---------------------|
| อินซูลินอะนาล็อกออกฤทธิ์เร็ว | | | |
| - Insulin lispro (Humalog [®]) | 5-15 นาที | 1-2 ชั่วโมง | 3-4 ชั่วโมง |
| - Insulin aspart (NovoRapid [®]) | 10-20 นาที | 1-2 ชั่วโมง | 3-4 ชั่วโมง |
| - Insulin glulisine (Aprida [®]) | 10-20 นาที | 1-2 ชั่วโมง | 3-4 ชั่วโมง |
| อินซูลินอะนาล็อกออกฤทธิ์ยาว | | | |
| - Insulin glargine (Lantus [®] , Basalin [®] , Glaritus [®]) | 2 ชั่วโมง | ไม่มี | 24 ชั่วโมง |
| - Insulin detemir (Levemir [®]) | 2 ชั่วโมง | ไม่มี | 18-24 ชั่วโมง |
| - Insulin degludec (Tresiba [®]) | 6 ชั่วโมง | ไม่มี | 24-36 ชั่วโมง |

Start basal insulin 0.1 unit /kg/day

เช่น น้ำหนัก 60 กิโลกรัม

เริ่ม NPH 6 unit sc hs



Case 3: BW 60 kg

60-year-old woman, last A1c=13%, glipizide 10 mg/day

| DTX เช้า | DTX เย็น | |
|-----------------|-----------------|------------------|
| 240 | 280 | NPH 6 unit sc hs |
| 204 | 275 | NPH 6 unit sc hs |
| 184 | 267 | NPH 6 unit sc hs |
| 189 | 289 | NPH 8 unit sc hs |
| 156 | 300 | NPH 8 unit sc hs |

If above A1C target

Basal plus:

Basal + bolus RI เฉพาะมื้อใหญ่

Multiple insulin injection

**ควร off glipizide

Premix bid

(Total 0.4-0.8 unit/kg/day)

** ควร off glipizide

Case 3: BW 60 kg

60-year-old woman, last A1c=13%, off glipizide

Start premix 0.5 unit/kg/day

| DTX เช้า | DTX เย็น | |
|-----------------|-----------------|--|
| 155 | 200 | Mixtard 20 unit sc ac เช้า Mixtard 10 unit sc ac เย็น |
| 136 | 174 | ฉีดเท่าเดิม |
| 124 | 155 | ฉีดเท่าเดิม |
| 132 | 185 | ฉีดเท่าเดิม |
| 125 | 177 | Mixtard 22 unit sc ac เช้า Mixtard 10 unit sc ac เย็น |

Premix insulin (30%RI+70%NPH)



| ชนิดยา (ชื่อยา) | เวลาที่เริ่มออกฤทธิ์ | เวลาที่มีฤทธิ์สูงสุด | ระยะเวลาการออกฤทธิ์ |
|--|----------------------|----------------------|---------------------|
| <p>ฮิวแมนอินซูลินผสมสำเร็จรูป</p> <ul style="list-style-type: none"> - Premixed 30% RI + 70% NPH (Mixtard 30 HM[®], Humulin 70/30[®], Gensulin M30[®], Insugen 30/70[®], Insuman combo30[®], Winsulin 30/70[®]) - Premixed 50% RI + 50% NPH (Gensulin M50[®]) | 30-60 นาที | 2 และ 8 ชั่วโมง | 12-20 ชั่วโมง |
| | 30-60 นาที | 2 และ 8 ชั่วโมง | 12-20 ชั่วโมง |



Novomix: 70% protamine aspart+30%Aspart



| Preparation | Trade name | Timing of Action | | |
|--------------------------------------|----------------------------|------------------|------|---------------|
| | | Onset | Peak | Duration |
| Pre-mixed insulin | | | | |
| 70%NPH / 30%Regular | Humulin 70/30 Mixtard30 | 30-60 นาที | Dual | 10-16 ชั่วโมง |
| 75% protaminated Lispro / 25% Lispro | Humalog mix 25 | 15-30 นาที | Dual | 10-16 ชั่วโมง |
| 70% Protaminated aspart / 30% aspart | Novomix 30 | 15-30 นาที | Dual | 10-16 ชั่วโมง |

Insulin in CKD

Table 4: Insulin preparations: Considerations in hemodialysis patients.

| INSULIN PREPARATION | ONSET OF ACTION | PEAK ACTION | EFFECTIVE DURATION |
|----------------------------------|-----------------|-------------|--------------------|
| Rapid-acting | | | |
| Regular | 30-60 min | 2-3 hr | 8-10 hr |
| Lispro (Humalog) | 5-15 min | 30-90 min | 4-6 hr |
| Aspart (NovoLog) | 5-15 min | 30-90 min | 4-6 hr |
| Long-acting | | | |
| Neutral protamine Hagedorn (NPH) | 2-4 hr | 4-10 hr | 12-18 hr |
| Glargine (Lantus) | 2-4 hr | None | 20-24 hr |
| Detemir (Levemir) | 3-4 hr | 3-14 hr | 6-23 (19.9) hr |
| Premixed | | | |
| 70/30 human mix | 30-60 min | 3-12 hr | 12-18 hr |
| 70/30 aspart mix | 5-15 min | 30-90 min | 12-18 hr |
| 75/25 lispro mix | 5-15 min | 30-90 min | 12-18 hr |

| EGFR (ml/min/1.73m ²) | Decrease doses of insulin |
|-----------------------------------|---------------------------|
| 10-50 | 25% |
| <10 | 50% |



How to start insulin in a patient with CKD

- Total daily dose (TDD) for insulin : **0.1 to 0.3 units/kg** (depend on nutritional status or frailty of the patient. (Obese patient : 1.2-1.5 units/kg)
- Regimens: multiple doses of insulin (MDI) or basal bolus regimen > premix
- The rule of thumb to prevent nocturnal hypoglycemia is-
“Bedtime glucose should always be higher than before dinner glucose by at least 40mg/dl



Conclusion

- Lifestyle modification is important management.
- Pharmacological management: risk specific selection:
 - SGLT2 inhibitor (ASCVD, Hx HF, CKD)
 - GLP1 agonist (ASCVD, CKD, weight management goal)

Thank you

For attention



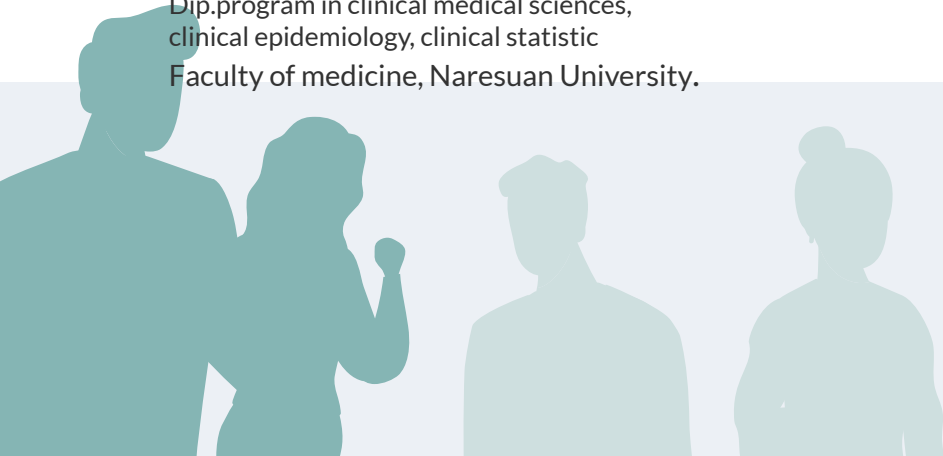
แนวทางเบื้องต้นในการจัดการ ผู้ป่วยก้าวร้าว

การใช้ยาทางจิตเวชเบื้องต้น



Dr. Fasinee Arunrodpanya

M.D., Dip.Thai Board of psychiatry,
Dip.program in clinical medical sciences,
clinical epidemiology, clinical statistic
Faculty of medicine, Naresuan University.

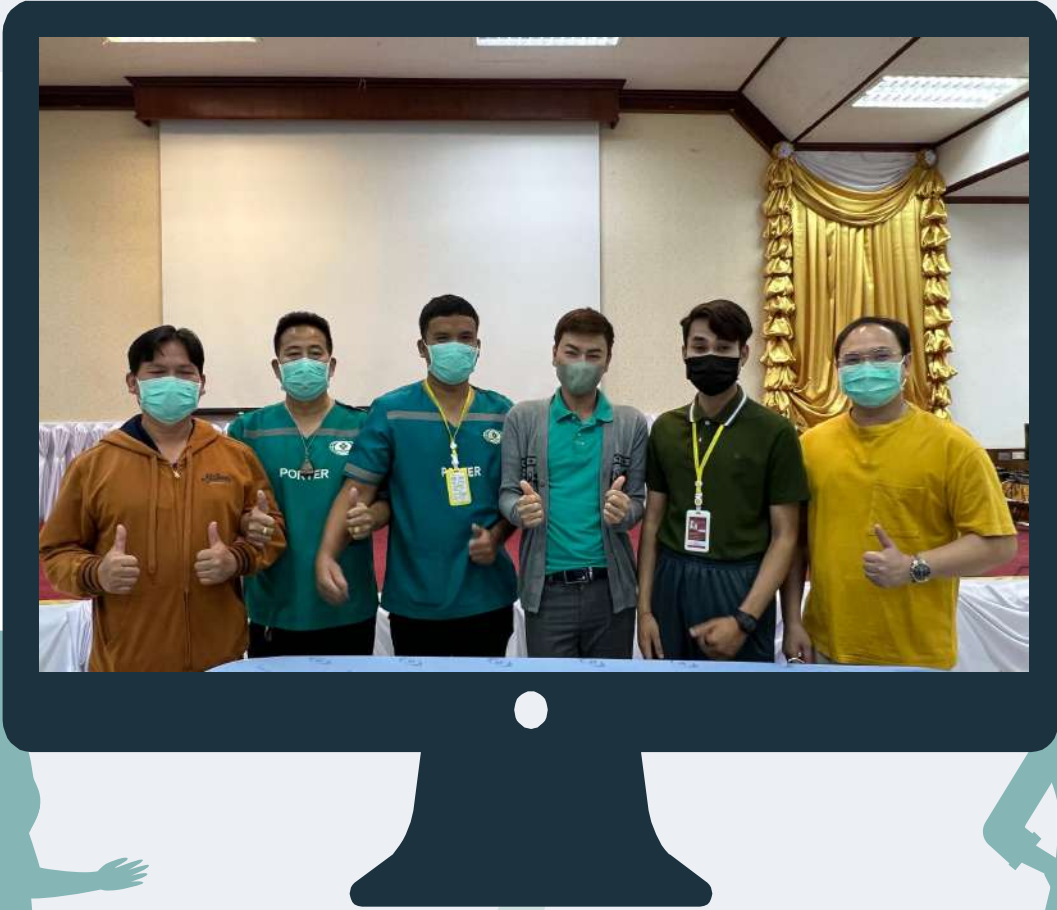


Special guest

**Assit.Prof.Dr.Jatuwit
Howannapakorn,MD.**

Forensic Medicine Department,
Naresuan University Hospital





พฤติกรรมก้าวร้าว ?

- ผู้ป่วยที่เคยมีประวัติพฤติกรรมรุนแรง
- ผู้ที่มีอาการแสดงที่เป็นสัญญาณพฤติกรรมรุนแรง



พฤติกรรมก้าวร้าว ?

- ผู้ที่มีอาการแสดงที่เป็นสัญญาณพฤติกรรมรุนแรง
 - **การแสดงออกทางสีหน้า/ท่าทาง**
 - สีหน้าบึ้งตึง โกรธ แหวงตาไม่เป็นมิตร
 - กัดกรามแน่น ดวงตาเปิดกว้าง
 - ท่าทางเกรี้ยว ดื้อไม่ผ่อนคลาย



พฤติกรรมก้าวร้าว ?

- ผู้ที่มีอาการแสดงที่เป็นสัญญาณพฤติกรรมรุนแรง

➤ การเคลื่อนไหว การกระทำ

- กระวนกระวาย อยู่ไม่นิ่ง เดินไปมา
- ตัวเกร็ง กำมือแน่น กำหมัด
- กระทบหรือ กระทำด้วยความรุนแรง
- หยุดการกระทำที่ทำอยู่ทันทีทันใด



พฤติกรรมก้าวร้าว ?

- ผู้ที่มีอาการแสดงที่เป็นสัญญาณพฤติกรรมรุนแรง
 - **การแสดงออกทางคำพูด**
 - เจ็บแค้นผิดปกติ
 - โต้ตอบด้วยน้ำเสียงห้วน
 - พูดก้าวร้าว วิพากวิจารณ์ ต่ำหนิติเตียน ต่ำว่าคำหยาบ สาบแซง
 - **การเปลี่ยนแปลงความรู้สึกอย่างทันทีทันใด**



การพูดคุยกับผู้ป่วย

- ทำที่เป็นมิตร สงบ จริงใจ
- พูดคุยด้วยน้ำเสียงนุ่มนวล
- ให้ผู้ป่วยได้ระบายความคิด
- แยกผู้ป่วยออกจากสถานการณ์
ลดสิ่งแวดล้อมกระตุ้นผู้ป่วย



การเตรียมอุปกรณ์การผูกยึด

- ผ้าผูกยึด ควรเป็นผ้าที่แข็งแรง เหนียว แต่นุ่ม และมีลักษณะพร้อมใช้
- ตำแหน่งข้อมือข้อเท้า และลำตัว
- เตียงที่สามารถใช้ผ้าผูกยึดได้



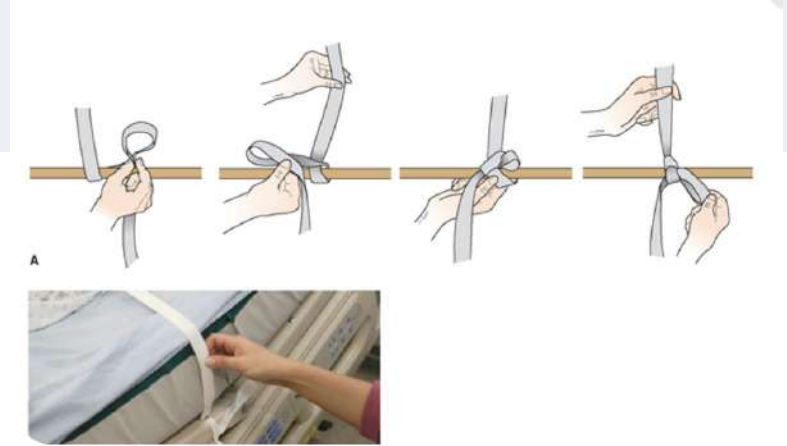


https://richardswsmith.files.wordpress.com/2020/11/437bc5547cb05ce157c78439f200feeb_large.jpg



Copyright © 2008 Lippincott Williams & Wilkins.

<https://o.quizlet.com/aBW7-FPuZvMipzm3xPE5SA.jpg>



<https://www.pinterest.com/pin/583497695449310990/>

ทีมที่ดูแลรักษา

- ควรมีอย่างน้อย 5-6 คน มีการตกลงร่วมกัน ปฏิบัติไปใน**แนวทางเดียวกัน**
- แพทย์ หรือ พยาบาล หรือสมาชิกที่สามารถพูด**เบี่ยงเบน**ความสนใจ
- สมาชิกคนที่**เข้าจับ**ผู้ป่วย เมื่อผู้ป่วยเผลอ
- สมาชิกที่**ช่วยจับ**แขนขาเพื่อการผูกยึด



ข้อควรปฏิบัติ

- การเข้าจับควรทำขณะผู้ป่วย**เพลอ**
- ผู้นำทีมต้องให้**สัญญาณ**ในการเข้าจับผู้ป่วย
- การเข้าจับควรยึดจับบริเวณ**ข้อพับใหญ่ๆ**
ข้อศอก เข่า เป็นต้น
- ควรผูกยึดผู้ป่วยในท่านอนหงาย



ข้อควรปฏิบัติ

- การผูกยึดที่ดีควรใช้นิ้วมือ **2 นิ้ว** สอดใต้ผ้าผูกยึดได้
- ตรวจเยี่ยมประเมินทุก **15-30 นาที** การเปลี่ยนท่า การรับประทานน้ำ อาหาร การขับถ่าย
- เมื่อสงบลง พุดคุยรู้เรื่องถึงยุติการผูกยึด



ยาทางจิตเวชเบื้องต้นสำหรับสถานการณ์นี้



First choice; antipsychotic

ระวัง alcohol intoxication

| | |
|---|---------------------------------|
| Haloperidol 5 mg IM q 30 minutes | EPS; dystonia, akathisia |
| Diazepam 10 mg IV, IM | Respiratory repression |
| Risperidone solution 2-4 cc oral | Sedative |
| | |

ยาทางจิตเวชเบื้องต้น



บุคคลากรผู้เหนือยล้า และหัวร้อน หงุดหงิดมานานเหลือเกิน
ควร work up **baseline** metabolic profile

| | |
|--|--------------------------------------|
| Sertraline (50) 1 tab oral OD pc เย็น | Loss appetite, N/V, palpitate |
| Lorazepam 0.5-1 mg oral hs | Sedative , daytime somnolence |
| | |
| | |

Thank you

For your attention

