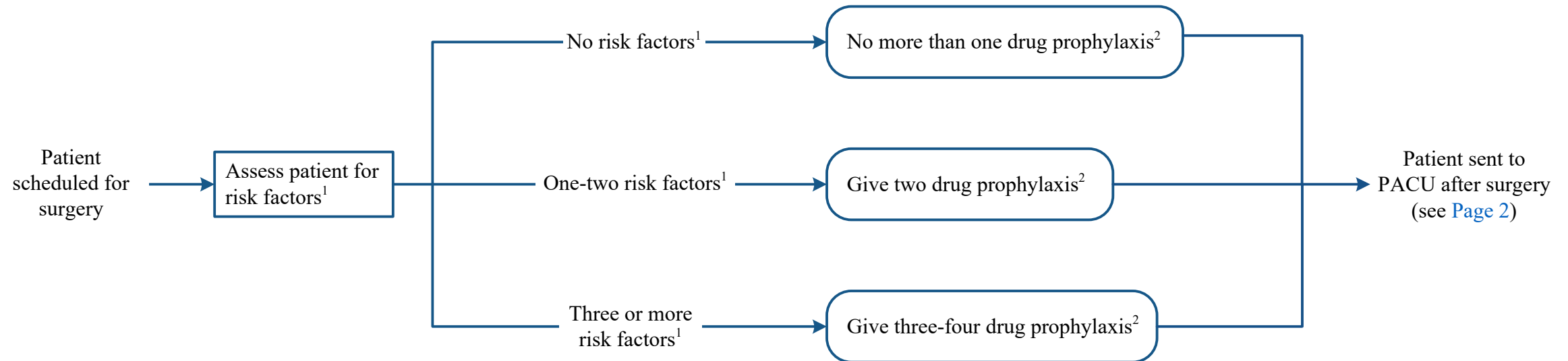


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PATIENT PRESENTATION

PROPHYLAXIS



Note: Strategies to minimize the risk of PONV:

- Minimization of perioperative opioids with the use of multimodal analgesia and regional anesthesia
- Avoidance of volatile anesthetics
- Proper intravascular hydration
- Implementation of total intravenous anesthesia
 - Option to intraoperatively run sub-hypnotic dose of propofol 20 mcg/kg/minute
- Avoid nitrous oxide
- Avoid neostigmine for reversal, use sugammadex

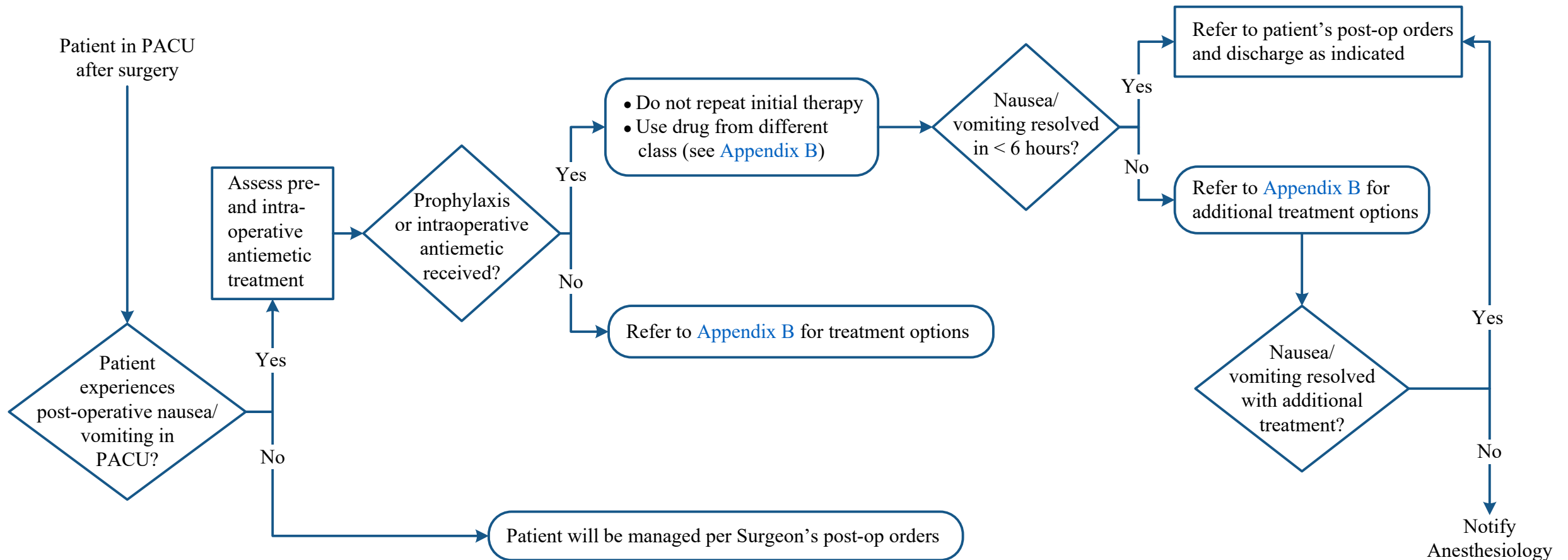
¹ MD Anderson risk factors

- **Patient specific risk factors:**
 - Female gender
 - Non-smoking status
 - History of post-operative nausea/vomiting (PONV) or motion sickness
 - Age < 50 years
- **Anesthetic risk factors:**
 - Use of volatile anesthetics
 - Post-operative opioids
- **Surgical risk factors:**
 - Duration of anesthesia > 3 hours
 - Type of surgery (abdominal, gynecologic, breast, head & neck surgery)

² See [Appendix A](#) – Antiemetic Medication Options for Prophylaxis

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TREATMENT



Nausea/Vomiting Associated with Surgery - Adult

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APPENDIX A: Antiemetic Medication Options for Prophylaxis

Drug	Dosage	Timing	Comments
Anticholinergics Scopolamine Patch (Transderm Scop [®])	1.5 mg disc placed behind ear	At least 2 - 4 hours before surgery	<ul style="list-style-type: none"> • Caution in patients > 60 years old • Patch may be applied the night prior to surgery • If not discontinued prior to hospital discharge, patients should be instructed in the safe removal and disposal of the patch
Benzodiazepines Midazolam (Versed [®])	35 - 75 mcg/kg IV	May be given pre-operatively or intra-operatively	
Butyrophenones Droperidol (Inapsine [®]) ¹	0.625 mg IV	Most effective if given at the end of surgery	<ul style="list-style-type: none"> • Requires 2 – 3 hours of EKG monitoring • Known risk² of TdP
Haloperidol (Haldol [®])	1 mg IV	Give at the end of surgery	<ul style="list-style-type: none"> • Alternative to droperidol • Known risk² of TdP and precludes its use as a first-line agent
Corticosteroids Dexamethasone	4-8 mg IV	Give shortly after induction	Avoid in labile diabetic patients
Dopamine Antagonist Amisulpride (Barhemsys [®])	5 mg IV	Give at the time of induction	<ul style="list-style-type: none"> • Less likely to cause adverse reactions such as extrapyramidal symptoms • Conditional risk² of TdP

TdP = torsades de pointes

¹ Availability varies based on supply

² The Arizona Center for Education and Research on Therapeutics (AZCERT)'s Adverse Drug Event Causality Analysis (ADECA) Risk Categories

- Known risk: Drugs in this category prolong the QT interval and are clearly associated with a risk of TdP, even when taken as recommended
- Possible risk: Drugs in this category can cause QT prolongation but currently lack compelling evidence for a risk of TdP when the drug is taken as recommended
- Conditional risk: Drugs in this category have evidence of TdP but only under certain conditions of their use (e.g., excessive dose, in patients with conditions such as hypokalemia or when they are taken with interacting drugs) or by creating conditions that facilitate or induce TdP (e.g., by inhibiting metabolism of a QT-prolonging drug or by causing an electrolyte disturbance that induces and/or facilitates TdP)

Continued on next page

Nausea/Vomiting Associated with Surgery - Adult

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APPENDIX A: Antiemetic Medication Options for Prophylaxis - continued

Drug	Dosage	Timing	Comments
Neurokinin-1 Receptor Antagonists Aprepitant (Emend [®])	40 mg PO	Give within 3 hours before the induction of anesthesia	
Phenothiazines Promethazine (Phenergan [®])	6.25 mg IV	Give shortly after induction	<ul style="list-style-type: none"> • 6.25 mg dose may require a second dose after 15 minutes; may repeat up to 3 times for a maximum dose of 25 mg • Should not be used in children ≤ 2 years old • Possible risk¹ of TdP
Prochlorperazine (Compazine [®])	5 - 10 mg IV	Give at the end of surgery	Risk of QTc prolongation did not reach the level to be placed in any of the TdP risk categories ¹ ; however, other tertiary drug information references ² and the product information ³ indicate ECG abnormalities (Q and T wave distortions) at an undefined frequency
Serotonin Antagonists Ondansetron (Zofran [®])	4 mg IV	Give at the end of surgery	Known risk ¹ of TdP
Granisetron	0.35 - 3 mg IV	Give at the end of surgery	<ul style="list-style-type: none"> • For patients with history of delayed (post-discharge) post-operative nausea and vomiting • Possible risk¹ of TdP

¹ The Arizona Center for Education and Research on Therapeutics (AZCERT)'s Adverse Drug Event Causality Analysis (ADECA) Risk Categories

- Known risk: Drugs in this category prolong the QT interval and are clearly associated with a risk of TdP, even when taken as recommended
- Possible risk: Drugs in this category can cause QT prolongation but currently lack compelling evidence for a risk of TdP when the drug is taken as recommended
- Conditional risk: Drugs in this category have evidence of TdP but only under certain conditions of their use (e.g., excessive dose, in patients with conditions such as hypokalemia or when they are taken with interacting drugs) or by creating conditions that facilitate or induce TdP (e.g., by inhibiting metabolism of a QT-prolonging drug or by causing an electrolyte disturbance that induces and/or facilitates TdP)

² See [Lexicomp](#)

³ See [prochlorperazine product information](#)

Nausea/Vomiting Associated with Surgery - Adult

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APPENDIX B: Antiemetic Medication Options for Treatment or Rescue

Drug	Dosage	Comments
Serotonin Antagonists Ondansetron (Zofran [®])	First Line Agent 4 mg IV	Known risk ¹ of TdP
Phenothiazines Promethazine (Phenergan [®])	Second Line Agents 6.25 mg IV	<ul style="list-style-type: none"> • 6.25 mg dose may require a second dose after 15 minutes; may repeat up to 3 times for a maximum dose of 25 mg • Possible risk¹ of TdP
Prochlorperazine (Compazine [®])	5 - 10 mg IV	Risk of QTc prolongation did not reach the level to be placed in any of the TdP risk categories ¹ ; however, other tertiary drug information references ² and the product information ³ indicate ECG abnormalities (Q and T wave distortions) at an undefined frequency
Butyrophenones Droperidol (Inapsine [®]) ⁴	Third Line Agents 0.625 mg IV	<ul style="list-style-type: none"> • Requires 2 - 3 hours of EKG monitoring • Known risk¹ of TdP
Haloperidol (Haldol [®])	1 mg IV	<ul style="list-style-type: none"> • Known risk¹ of TdP and precludes its use as a first-line agent • Alternative to droperidol
Prokinetic Metoclopramide (Reglan [®])	Rescue 10 mg IV	Conditional risk ¹ of TdP
Dopamine Antagonist Amisulpride (Barhemsys [®])	Rescue 10 mg IV	<ul style="list-style-type: none"> • Less likely to cause adverse reactions such as extrapyramidal symptoms • Conditional risk¹ of TdP

Notes:

- When nausea and vomiting occur post-operatively, treatment should be administered with an antiemetic from a DIFFERENT pharmacologic class than the drug given for prophylaxis initially
- Re-dosing should only occur if ≥ 6 hours has elapsed since the last dose from that class was given

TdP = torsades de pointes

¹ The Arizona Center for Education and Research on Therapeutics (AZCERT)'s Adverse Drug Event Causality Analysis (ADECA) Risk Categories

- Known risk: Drugs in this category prolong the QT interval and are clearly associated with a risk of TdP, even when taken as recommended
- Possible risk: Drugs in this category can cause QT prolongation but currently lack compelling evidence for a risk of TdP when the drug is taken as recommended
- Conditional risk: Drugs in this category have evidence of TdP but only under certain conditions of their use (e.g., excessive dose, in patients with conditions such as hypokalemia or when they are taken with interacting drugs) or by creating conditions that facilitate or induce TdP (e.g., by inhibiting metabolism of a QT-prolonging drug or by causing an electrolyte disturbance that induces and/or facilitates TdP)

² See [Lexicomp](#)

³ See [prochlorperazine product information](#)

⁴ Availability varies based on supply

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DEVELOPMENT CREDITS

This practice consensus statement is based on majority expert opinion of the Nausea and Vomiting experts at the University of Texas MD Anderson Cancer Center for the patient population. These experts included:

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