

Improve the Quality of Care Amongst Surgical Patients with Obstructive Sleep Apnea

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Background

- Obstructive sleep apnea (OSA) affects about 29 million people in the U.S.
- Up to 80% of individuals with moderate to severe OSA remain undiagnosed
- Patients at risk for OSA are at increased risk for perioperative complications
- Anesthesia providers must identify patients with OSA to optimize and tailor their anesthetic

Purpose

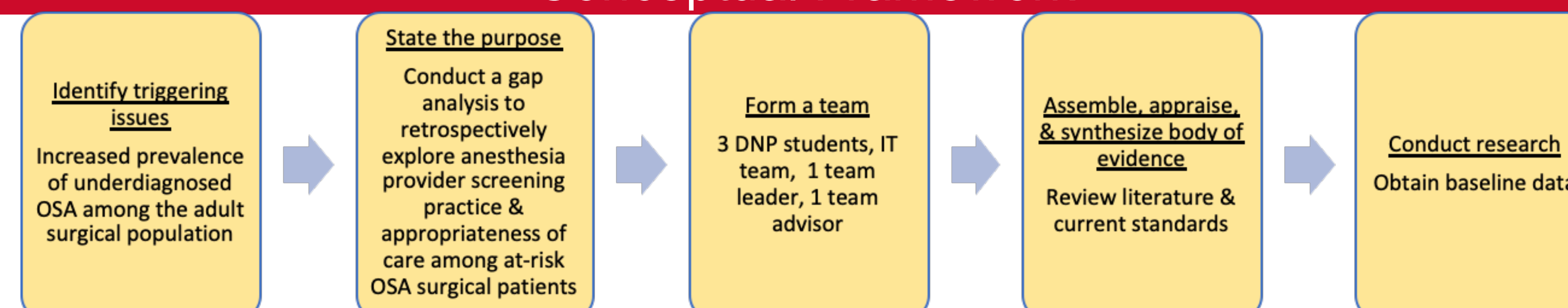
The purpose of this project is to identify whether current management practices amongst the patient population with OSA are in accordance with ASA guidelines for OSA

Current Standards: ASA guidelines for OSA

Clinical Recommendations

- Follow a developed process whereby patients at risk for OSA are evaluated for OSA prior to the day of surgery
- “Cautious” use of benzodiazepines
- Multimodal analgesia approach
- Initial intubation attempt with video laryngoscopy
- Sugammadex vs neostigmine: promising respiratory profile with sugammadex, but no current recommendation

Conceptual Framework



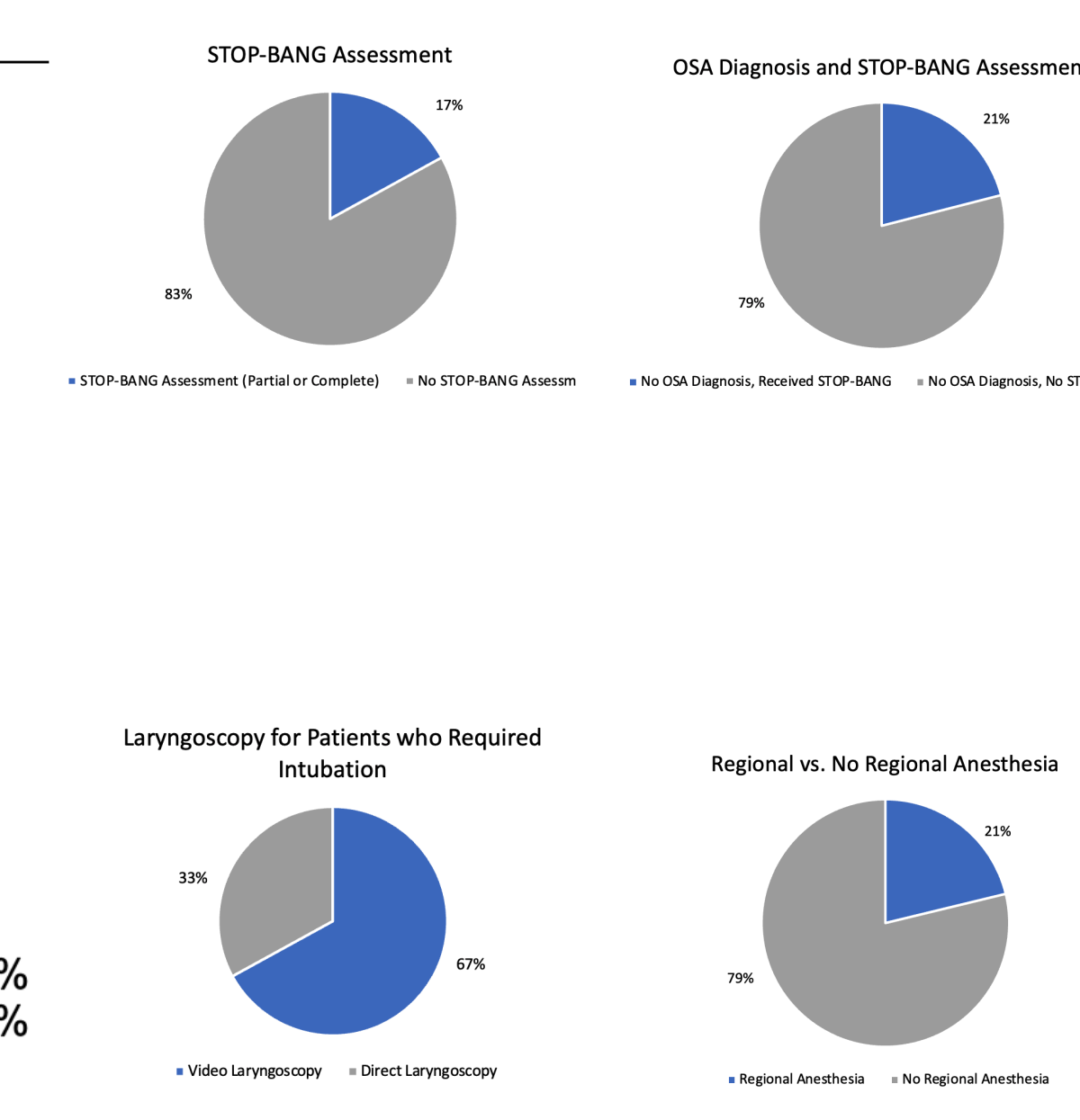
Methods

- **Design:** Retrospective chart review of electronic health records
- **Setting:** Large Integrated Health System
- **Sample:** 100 adult patients at risk for OSA who underwent surgeries in the last quarter of 2019 ($n = 100$)
- **Measures:** Adherence to practice guidelines/current standards in the perioperative period
 1. Percentage of patients who received a STOP-BANG assessment
 2. Percentage of patients without a prior OSA diagnosis and received a STOP-BANG assessment
 3. Percentage of patients intubated with video laryngoscopy
 4. Percentage of patients who received preoperative sedation (midazolam, dexmedetomidine, ketamine)
 5. Percentage of patients discharged with multimodal analgesic medications versus opioids
 6. Percentage of patients who received regional anesthesia/analgesia
 7. Percentage of patients who received reversal agents (neostigmine vs. sugammadex)

Results

Characteristics of Study Subjects

Baseline characteristic	All patients	Baseline characteristic	OSA Diagnosis	No OSA Diagnosis
<i>n</i>	100	<i>n</i>	51	49
ASA class		Location of Procedure		
I	0	Main Operating Room	48	94%
II	13	Ambulatory Surgical Center	3	6%
III	83	STOP-BANG Assessment		
IV	4	No	44	86%
BMI (mean)	43.89	Partial	1	2%
Location of Procedure		Complete	6	12%
Main Operating Room	94	Preoperative Sedation		
Ambulatory Surgical Center	6	Midazolam only	30	30
Pre-existing Diagnosis of OSA		Midazolam dose (mg)	1.33	1.48
Yes	51	Midazolam with Other Sedative	4	4
No	49	Midazolam dose (mg)	1.75	1.75
STOP-BANG Assessment		None	12	24%
No	83	Regional Anesthesia	12	24%
Partial	2	Intubated		
Complete	15	Video Laryngoscopy	31	65%
Preoperative Sedation		Direct Laryngoscopy	17	35%
Midazolam only	60	Reversal Agent, If Given		
Average Midazolam dose (mg)	1.9	Neostigmine	18	50%
Midazolam with Other Sedative	8	Sugammadex	18	50%
Average Midazolam dose (mg)	1.75	Paralysis, No Reversal	14	9
None	22	No Paralysis, No Reversal	1	1
Regional Anesthesia	21			
Intubated	96			
Video Laryngoscopy	65			
Direct Laryngoscopy	31			



- Midazolam only: average dose 1.9mg
- Midazolam with other sedative: average dose of midazolam 1.75mg
- Received no preoperative sedation at all (22%)

Discussion

Major Findings:

- Limited number of STOP-BANG completions for patients not diagnosed with OSA
- Lack of reduction in benzodiazepine use
- One medical service area was an outlier and did not use any benzodiazepines in all patients
- Majority of initial intubation attempts were with video laryngoscopy
- Half of reversals were with sugammadex while half with neostigmine
- Limited amounts of regional anesthesia/analgesia were performed
- Half of the patients were discharged with opioid only analgesic medications
- Minimal difference in care regardless of preexisting diagnosis of OSA

Limitations:

- Retrospective data
- Limited sample size
- Inconsistencies in documentation location
- Human error in the process of data collection

Recommendation

Implementation of a standardized tool to evaluate suspected OSA patients.

Future possibilities of study include strategies to monitor compliance of the tool and dissemination of data regarding guideline adherence after successful tool implementation

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