

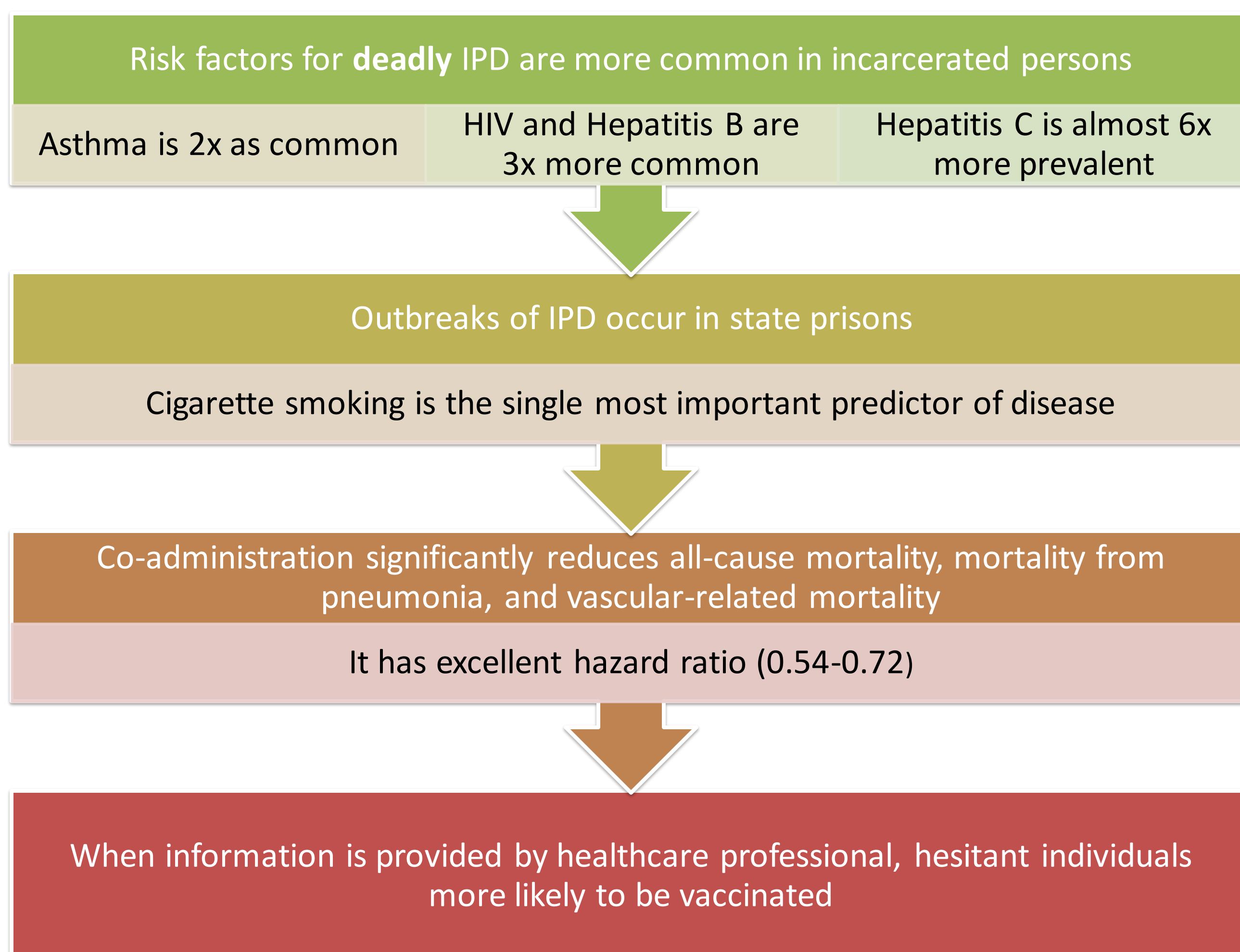
Pneumococcal & Influenza Vaccine Co-Administration in the Incarcerated Population

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Southern Illinois University Edwardsville

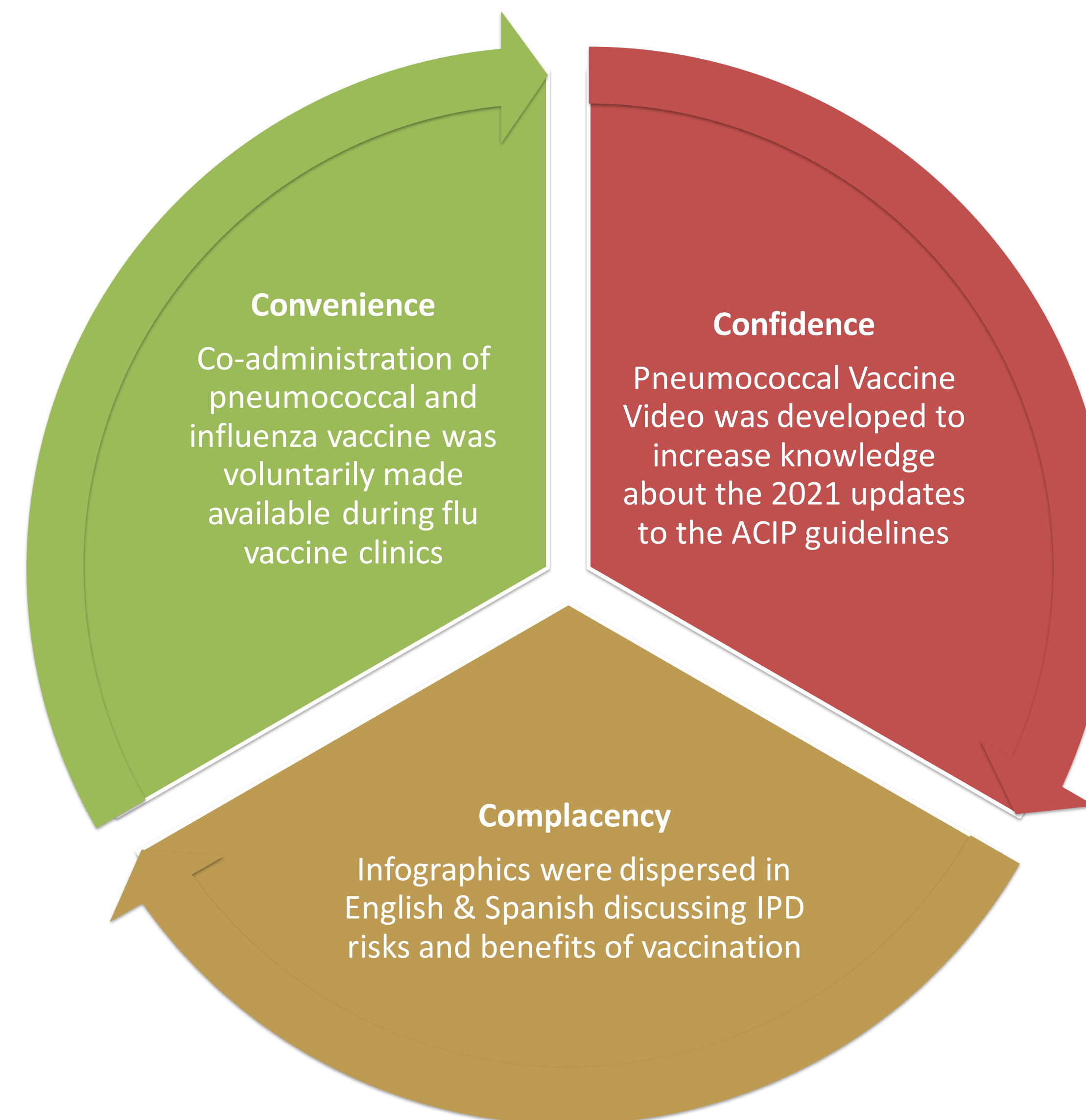
PROBLEM INTRODUCTION

- Streptococcus Pneumoniae causes 20-60% of cases of bacterial pneumonia
- It has a mortality rate of 10-30% depending on risk factors
- Following ACIP Guidelines is 60-70% effective at preventing invasive pneumococcal disease
- Incarcerated individuals have increased risk of contracting pneumococcal disease due to the proximity of living quarters
- Department of Corrections reported 43% of incarcerated individuals <65 & 47% of those >=65 years have received pneumococcal vaccine

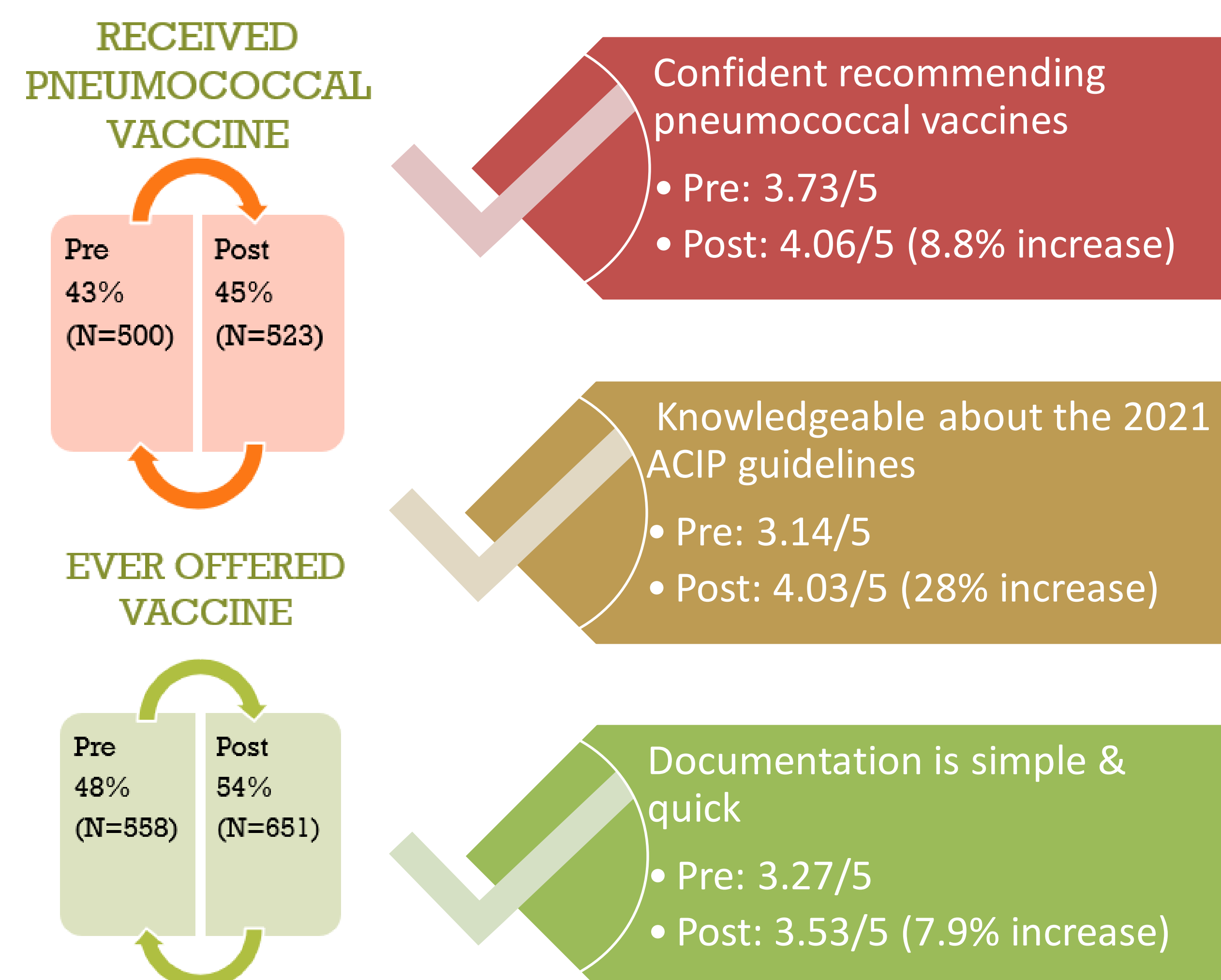
LITERATURE REVIEW



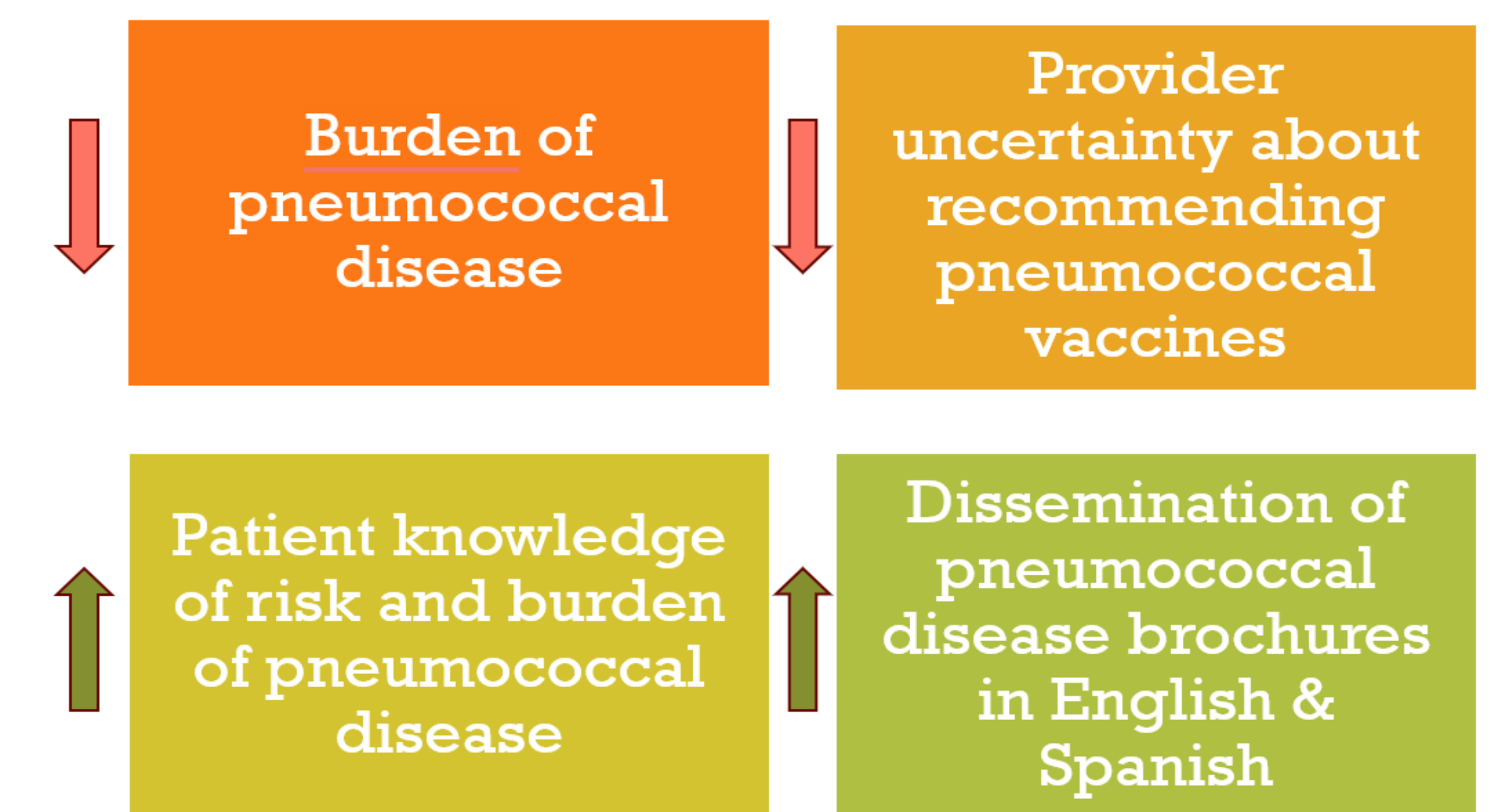
PROJECT METHODS



EVALUATION



IMPACT ON PRACTICE



CONCLUSIONS



PNEUMOCOCCAL PNEUMONIA FACTS

AWARENESS
Streptococcus pneumoniae (the bacteria the pneumonia vaccine prevents) has a mortality of 10-30%. Those with risk factors like smoking, asthma, or HIV are more likely to get the infection and die from it.

THE PNEUMONIA VACCINE IS 60-70% EFFECTIVE AT PREVENTING PNEUMOCOCCAL DISEASE

IDOC
just over 4 out of 10 individuals in custody are vaccinated against pneumococcal disease

REASONS TO GET VACCINATED

- Smoking
- Lung disease
- Diabetes
- Alcoholism
- >64 years old
- Heart disease
- Spleen removal
- Immune deficiency

DID YOU KNOW: YOU CAN GET BOTH THE FLU AND PNEUMOCOCCAL VACCINES ON THE SAME DAY? SCHEDULE YOUR VISIT TODAY

REFERENCES

HECHOS DE NEUMONÍA NEUMOCÓCICA

CONCIENCIA
Streptococcus pneumoniae (la bacteria que previene la vacuna contra la neumonía) tiene una mortalidad del 10-30%. Las personas con factores de riesgo como el tabaquismo, el asma o el VIH tienen más probabilidades de contraer la infección y morir a causa de ella.

LA VACUNA TIENE UNA EFICACIA DEL 60-70% EN LA PREVENCIÓN DE LA ENFERMEDAD NEUMOCÓCICA

POCO MÁS DE 4 DE CADA 10 PERSONAS DETENIDAS ESTÁN VACUNADAS CONTRA LA ENFERMEDAD NEUMOCÓCICA

RAZONES PARA VACUNARSE

- Tabaquismo
- Enfermedad Pulmonar
- Diabetes
- Alcoholismo
- >64 años
- Cardiopatía
- Extirpación del Bazo
- Inmunodeficiencia del Bazo

¿SABÍA USTED QUE PUEDE RECIBIR LAS VACUNAS CONTRA LA GRIPE Y EL NEUMOCOCCO EL MISMO DÍA? AGENDA TU VISITA HOY

REFERENCIAS

Recommendations for Sugammadex Administration in Standard and Special Populations

Kristen Mattson, BSN, SRNA and Alexa Brummund, BSN, SRNA
Southern Illinois University Edwardsville

PROBLEM

- Lack of standardization for use of sugammadex in standard and special populations including renal failure, breast feeding, pregnancy, and pediatrics at Memorial Hospital Belleville.
- Concerns about lack of evidence-based information about the use of sugammadex, access to quick references, and understanding of the cost analysis about the drug.
- This project will bridge the knowledge gap, provide evidence-based references, and explore causes for barriers to use of the medication.



PROJECT METHODS

IRB approval was obtained from SIUE and Memorial Hospital Belleville.

Non-experimental single group design using a convenience sample of approximately 25 anesthesia providers.

Guidelines were created based on the current literature regarding sugammadex in standard and special patient populations.

Current guidelines presented to the anesthesia staff. A quick reference card was distributed to all providers for ease of access.

A post-implementation survey assessed current use of sugammadex, barriers to use, and understanding of the use in special populations.

DOSING

≥ 2 twitches = 2 mg/kg
 < 2 twitches or only post-tetanic twitches = 4 mg/kg
 Cannot intubate / cannot ventilate = 16 mg/kg
 Dose should be calculated on **ACTUAL BODY WEIGHT.**

Side Effects: Most common include rash, bradycardia, hypotension, nausea and vomiting, and prolonged clotting times. More commonly seen with large doses (16mg/kg) (Merck & Co, 2022).

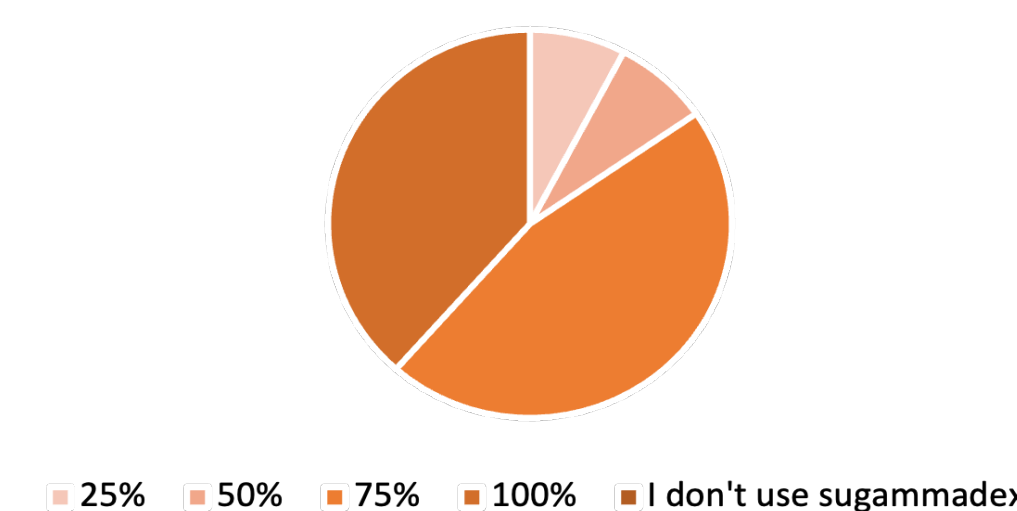
LITERATURE REVIEW

Clinical Relevance	<ul style="list-style-type: none"> Inadequate reversal of chemically induced paralysis can lead to increased morbidity & mortality including hypoxia, respiratory failure & increased length of hospital stay (Ayad et al., 2019).
Renal Impairment	<ul style="list-style-type: none"> Should not be used if creatinine clearance < 30ml/min No dose adjustments required. Slightly prolonged onset High flux dialysis within 24-48 hours of administration (Paredes et al., 2020).
Breast Feeding	<ul style="list-style-type: none"> Avoid in the first 10 days postpartum. The large molecule can pass through maternal lactating ducts (Willett et al., 2019). Weigh risks versus benefits; effects on lactation are unknown (Willett et al., 2019).
Pregnancy	<ul style="list-style-type: none"> Avoid in 1st trimester Safe to use near term (37 weeks) (Willett et al., 2019)
Birth Control	<ul style="list-style-type: none"> Utilize nonhormonal birth control for 7 days after administration Hormonal birth control includes pills, IUDs, vaginal rings & implants (Willett et al., 2019).
Pediatrics	<ul style="list-style-type: none"> Not FDA approved in children < 2 years (Merck & Co, 2022). Safe in children > 2 years; same dosing as adults
Cost Analysis	<ul style="list-style-type: none"> Sugammadex resulted in fewer minutes in the OR and PACU when compared to neostigmine (Moss et al., 2022). Although sugammadex is more costly than neostigmine, saving OR time results in decreased overall costs (Childers & Maggard-Gibbons, 2018).

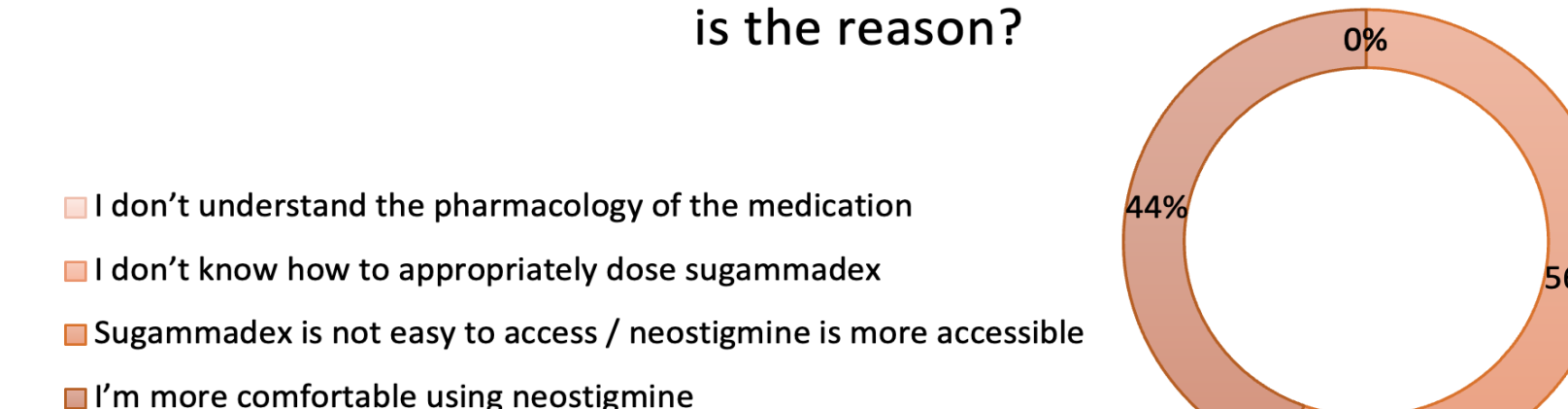
EVALUATION

- Multiple choice & Likert-style questions utilized in survey
- 13 total participants included in analysis
- 84.6% of participants indicated an increase in knowledge of sugammadex in standard and special patient populations and increased confidence in using sugammadex after implementation.
- Majority used sugammadex due to its reliable course of reversal and shorter reversal time.
- 100% said they would increase their use of sugammadex if it was accessible in a Pyxis in each OR.

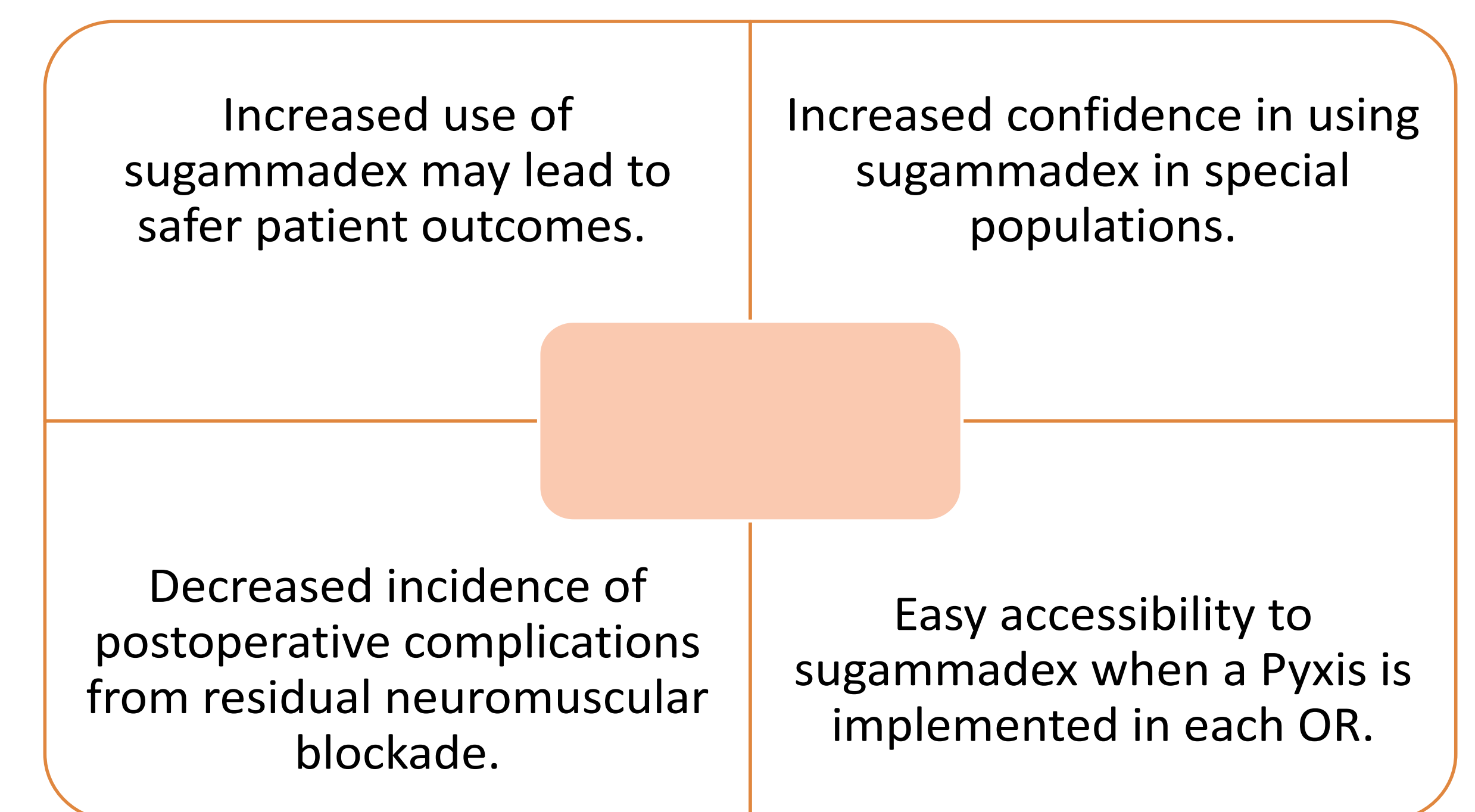
Approximately how often do you use sugammadex?



In the cases in which you do not use sugammadex, what is the reason?



IMPACT ON PRACTICE



CONCLUSIONS

Overall, current literature shows that sugammadex is superior to neostigmine in the reversal of steroidal NDMRs. Most of the data for the use of sugammadex in special populations shows that additional research is needed. Sugammadex has shown to decrease time in the OR, potentially further reducing healthcare costs. Further conclusions after implementation showed an increase in provider knowledge about the use of sugammadex in standard and special populations, with additional education and quick reference card. Barriers included lack of accessibility and more comfort with other reversal agents. 100% of participants found the quick reference card user friendly along with an increase in confidence in using sugammadex.

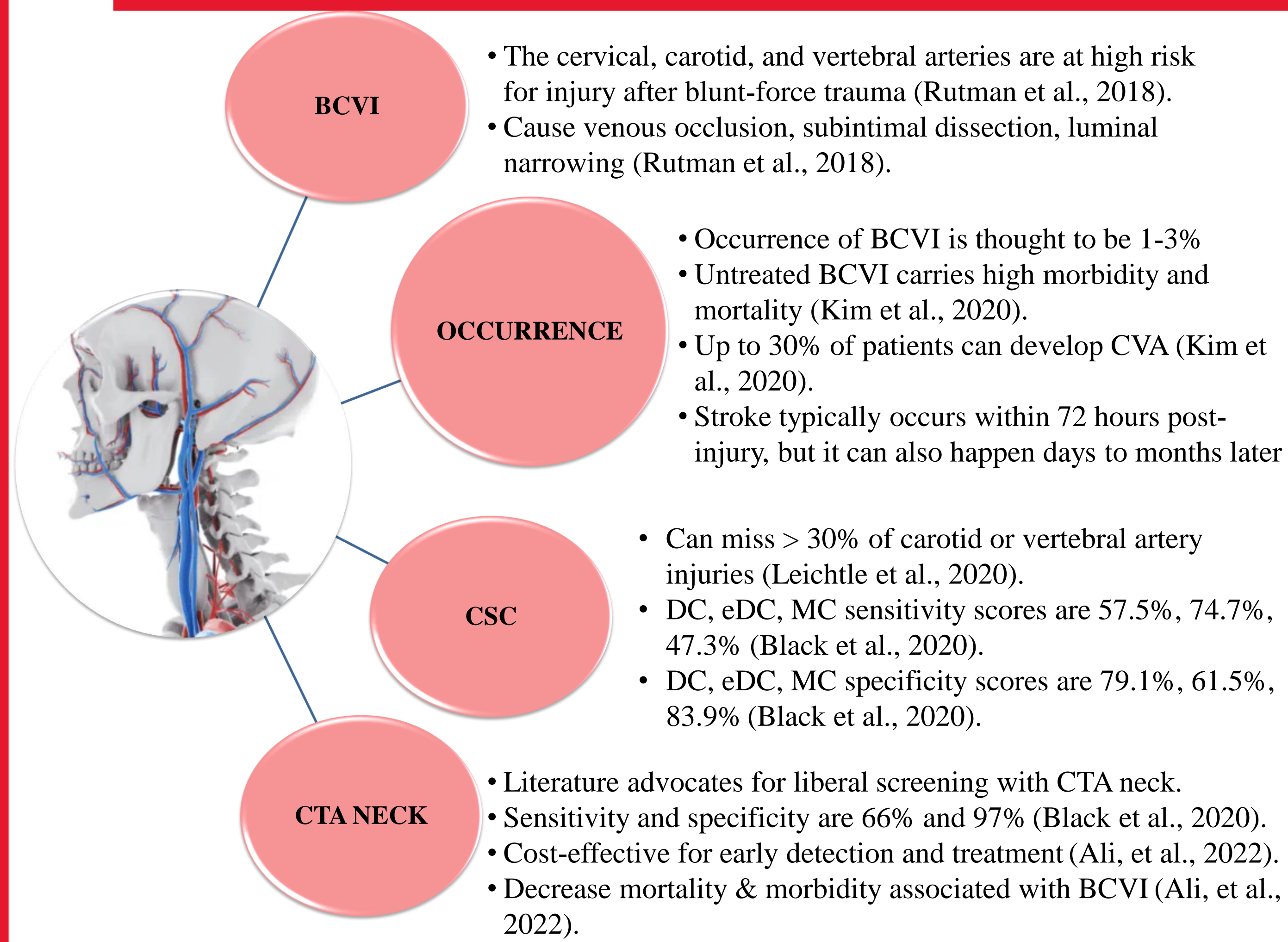
Blunt Cerebrovascular Injury (BCVI): Universal CTA Neck Screening at Level 2 Trauma Center

Tamine Gogel, MSN, APRN, Student DNP/MBA
Southern Illinois University Edwardsville

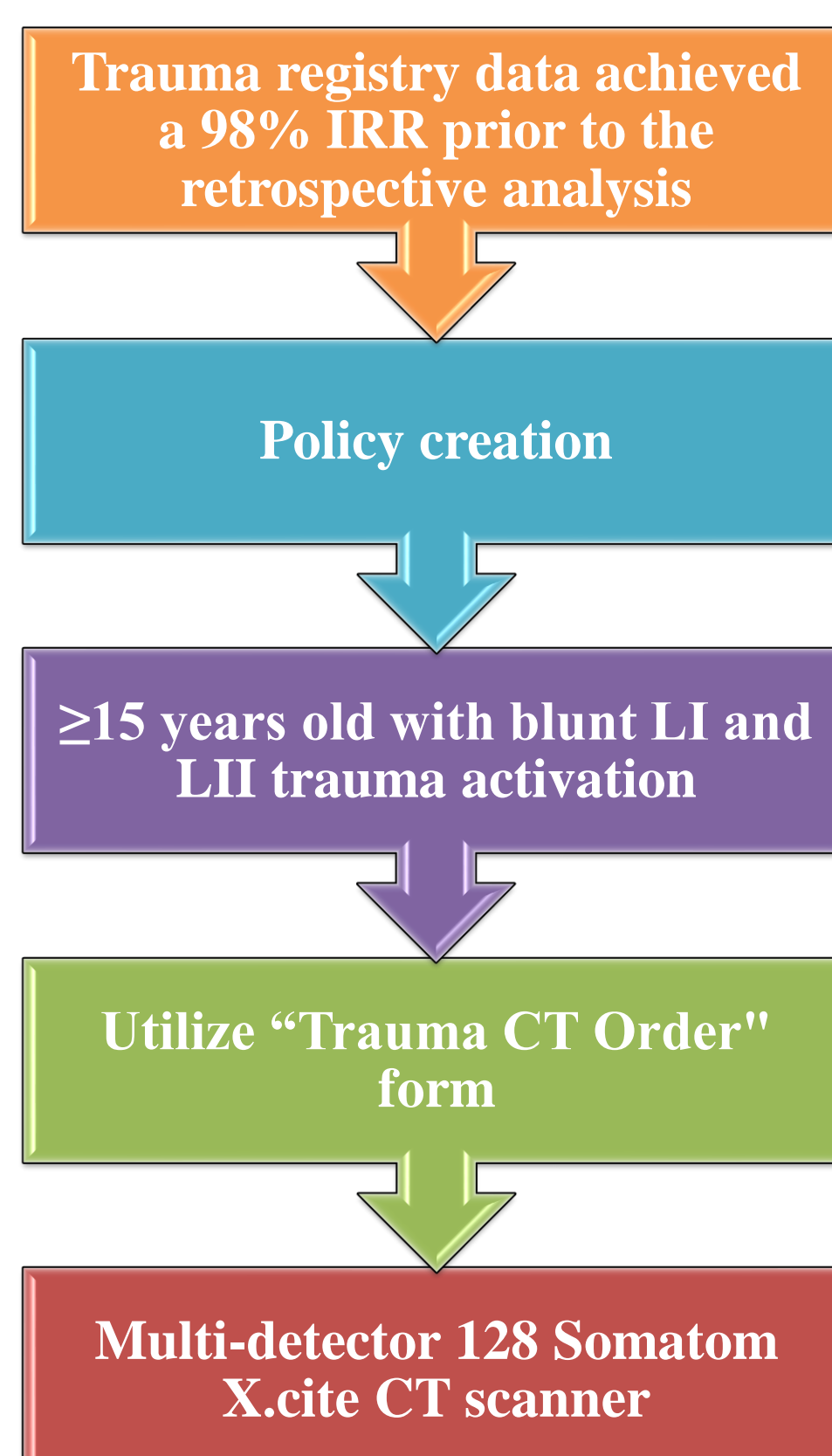
PROBLEM INTRODUCTION

- The project site uses clinical screening criteria (CSC) with extended Denver criteria (eDC), Denver criteria (DC), and the Memphis criteria (MC) to screen for BCVI in patients with blunt-force trauma.
- Relying solely on CSC will result in undiagnosed BCVI (Harper et al. (2022)).
- Literature supports the use of CTA neck to screen for BCVI (Ali et al., 2022; Black et al., 2020; Harper et al., 2022; Kim et al., 2020; Leichtle et al., 2020).
- The project aims to implement CTA neck during the initial blunt trauma assessment for early detection & treatment of BCVI in patients who sustained blunt force injuries.

LITERATURE REVIEW

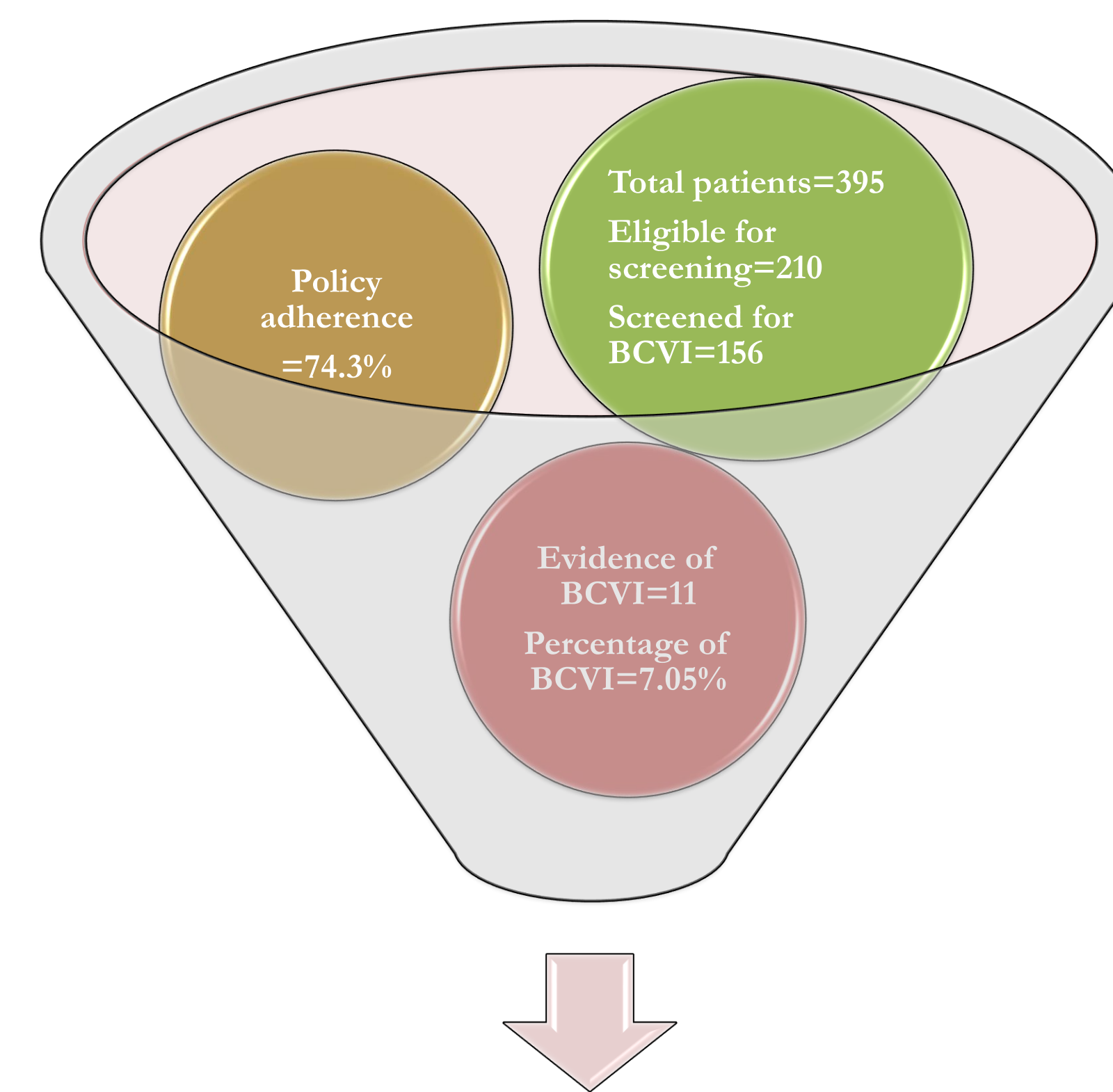
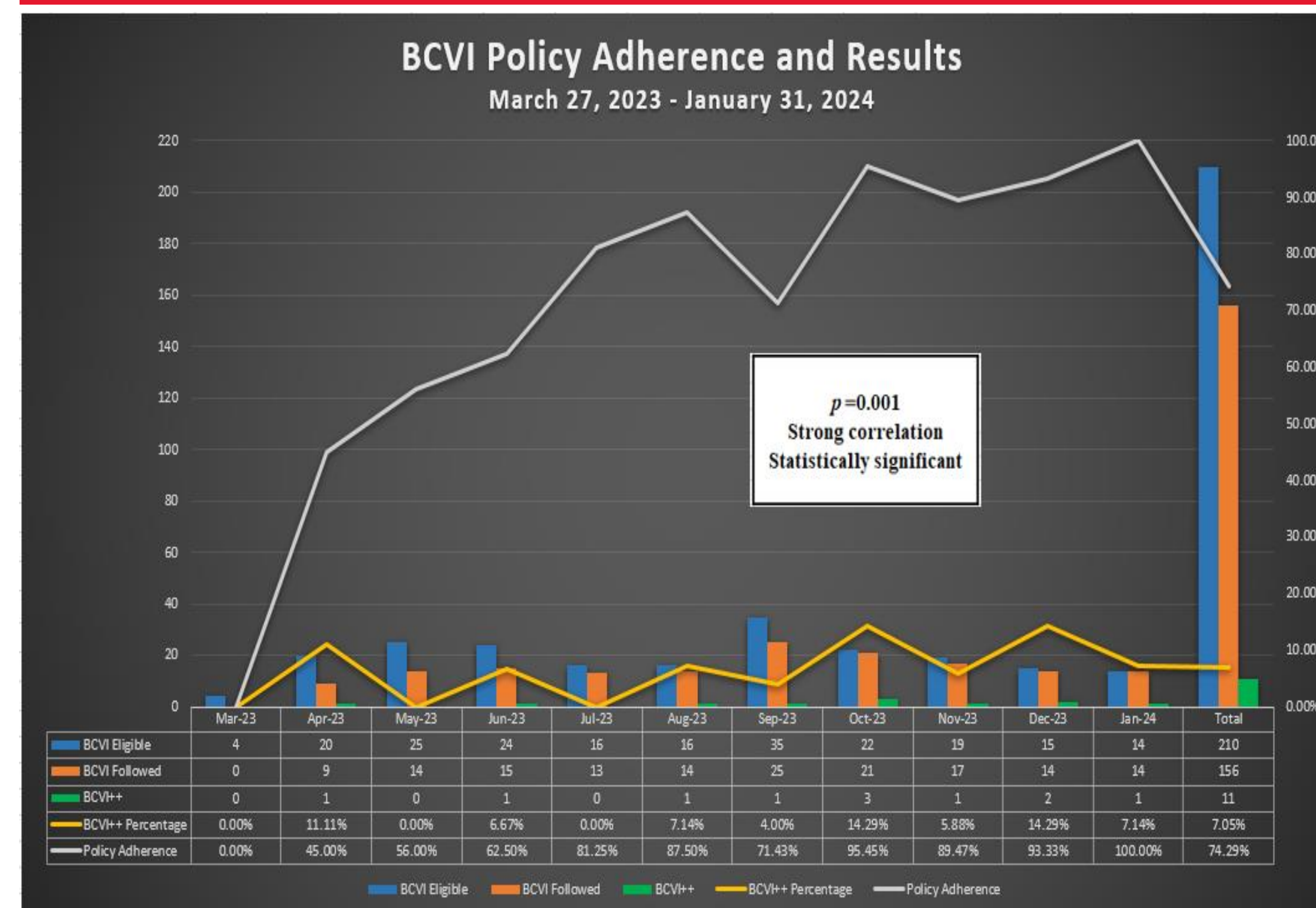


PROJECT METHODS



- UNIVERSAL SCREENING GUIDELINE:**
- Major blunt force trauma requiring a CT head and C-spine will get Universal BCVI screen with one of the following:
 - CT Head/Face without contrast
 - CTA Neck or CTA Head/Neck (scan at about 20 seconds)
 - Inject 100mL of Omnipaque 350
 - If pt is under 100 lbs then Omnipaque 300 (1mL per pound).
 - Do C-spine reconstructions off of the CTA neck
 - CT Chest/Abd/Pelvis (scan portal venous phase at 1 minute)
 - Do T/L spine reconstructions off of the CAP
 - Delay CT Abd/Pelvis
- OR If scan includes CTA CAP:**
- CT Head/Face without contrast
 - CTA Neck or CTA Head/Neck
 - Inject 70mL of Omnipaque 350 (1st injection)
 - If pt is under 100 lbs then Omnipaque 300, 1mL per pound.
 - Do C-spine reconstructions off of CTA neck
 - CTA Chest/Abd/Pelvis
 - Inject 100mL of Omnipaque 350 (2nd injection)
 - If pt is under 100 lbs then Omnipaque 300 (1mL per pound).
 - Do T/L spine reconstructions off of CTA CAP
 - Delay CT Abd/Pelvis

EVALUATION

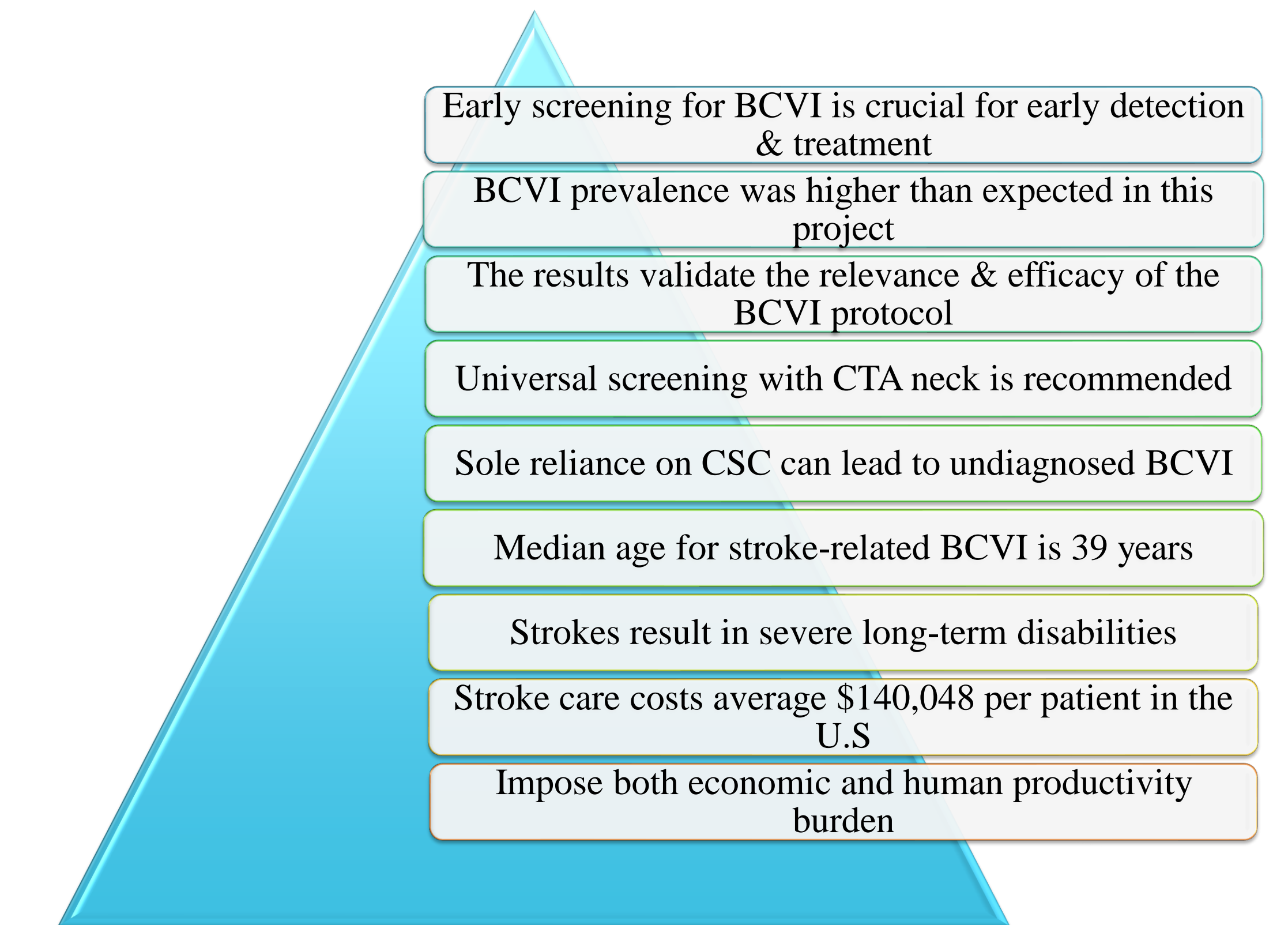


$r=0.97$ showed positive linear relationship between "BCVI Followed" and "Positive BCVI"
 $p<0.001$ emphasized the reliability of the observed correlation

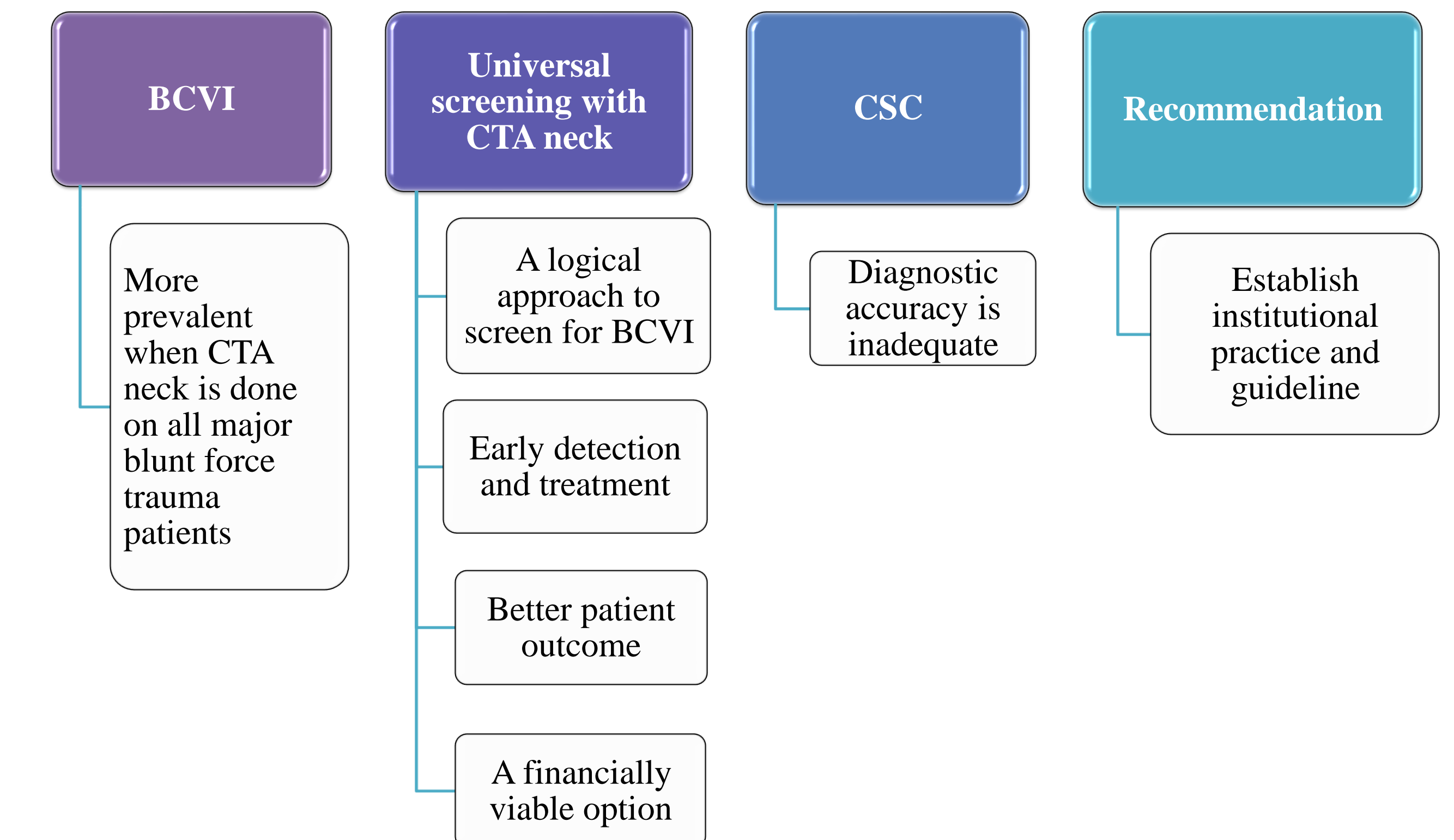
LIMITATIONS

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- Hemodynamically unstable patients did not undergo a complete initial radiographic workup
 - Transfer patients who already had CT scans
 - Human memory resulted in poor compliance during the initial phase of the project.
- Improved by sending bi-weekly scorecards to the trauma surgeons

IMPACT ON PRACTICE



CONCLUSIONS



REFERENCES

- Ali, A., et al. (2022). Cost effectiveness of universal screening for blunt cerebrovascular injury: A markov analysis. *Journal of the American College of Surgeons*. Publish Ahead of Print. <https://doi.org/10.1097/xcs.0000000000000490>
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