

## Background

- Cangrelor is a reversible intravenous (IV) antiplatelet that inhibits the P2Y<sub>12</sub> receptor<sup>1,2</sup>
- It is used in Percutaneous Coronary Interventions (PCIs) due to its quick onset of action of approximately two minutes. Platelet function returns to normal reactivity about an hour after infusion discontinuation<sup>1,2</sup>
- Cangrelor is dosed as a 30 mcg/kg bolus followed by a 4 mcg/kg/min infusion<sup>3</sup>
  - Cangrelor is recommended to be infused for two hours, or the entire duration of the PCI, whichever is the longest<sup>3</sup>
- The main adverse event with cangrelor is bleeding and it is contraindicated in patients with an active bleed<sup>4</sup>
- The rate of bleeding was about 0.56% with cangrelor in the CHAMPION PHOENIX trial, which was not significantly increased from clopidogrel alone<sup>5</sup>
- BJC HealthCare has approved the use of cangrelor in the cardiac catheterization lab if the patient meets one of the following criteria:
  - High risk stable elective PCI cases with complex anatomy not loaded with an oral P2Y<sub>12</sub> inhibitor within 2 hours of the procedure
  - NSTEMI/UA undergoing urgent PCI and not loaded with an oral P2Y<sub>12</sub> inhibitor and will go to the cath lab within 2 hours for PCI
  - STEMI patient in which GIIbIIIa inhibitors are not planned to be used, bivalirudin or heparin will be used, and the patient was not loaded with an oral P2Y<sub>12</sub> inhibitor prior to PCI

## Purpose

- To evaluate if the patients met the BJC approved criteria for use to receive cangrelor during a PCI at Memorial Hospital Belleville and Memorial Hospital East

## Methods

- Study design: retrospective chart review
- Inclusion criteria:
  - Age ≥ 18 years old
  - Cangrelor administered for use during a PCI between 12/1/18 to 6/1/19 at Memorial Hospital Belleville and Memorial Hospital East
- Exclusion criteria:
  - Cangrelor pulled from the automated dispensing cabinet under the patients name, but was never used and later returned
  - Cangrelor used for bridging therapy while holding oral antiplatelets
- A report was generated for all patients that had cangrelor pulled from the automated dispensing cabinet during his or her stay
- Primary outcome: appropriate usage of cangrelor based on BJC HealthCare's criteria for use
- Secondary outcomes:
  - Bleed occurrence during PCI
  - Bleed occurrence within 72 hours of procedure
  - Death due to MI/ACS/CVA or bleeding during hospital stay
  - Readmission to Memorial due to MI/ACS or bleeding associated events within 30 days of discharge
- Provider trends
- Events not meeting bleed definition

## Results

Table 1. Patient Information	Total, No. (%)
All Patients	(n=47)
<b>Patients with cangrelor administered in the cath lab</b>	<b>(n=42)</b>
Patients with high risk stable elective PCI	12 (28.6)
Patients with NSTEMI	12 (28.6)
Patients with STEMI	18 (42.9)
<b>Patient location during PCI</b>	
Memorial Hospital Belleville	26 (61.9)
Memorial Hospital East	16 (38.1)
<b>Appropriate criteria for cangrelor use met</b>	
Yes	41 (97.6)
No	1 (2.4)

Table 2. Provider Trends	Total, No. (%)
<b>Number of PCI procedures performed using cangrelor</b>	
Provider A	17 (40.5)
Provider B	12 (28.6)
Provider C	2 (4.8)
Provider D	10 (23.8)
Provider E	1 (2.4)
<b>Appropriate dosing of cangrelor</b>	
Yes	38 (90.5)
No	3 (7.1)
Unknown	1 (2.4)
<b>Duration of cangrelor infusion (n=41)</b>	
Less than 2 hours	1 (2.4)
2 hours	30 (75)
2-3 hours	4 (9.8)
Greater than 3 hours	6 (14.6)
<b>Average infusion duration per provider</b>	
Provider A	2.4 hours
Provider B	3.1 hours
Provider C	2.9 hours
Provider D	1.9 hours
Provider E	2 hours
<b>P2Y<sub>12</sub> inhibitor used post procedure (n=37)</b>	
Ticagrelor	37 (100)
<b>Dual antiplatelet therapy prescribed at discharge (n=35)</b>	
Yes	33 (94.3)
No	2 (5.7)

## Results

Table 3. Outcomes	Total, No. (%)
<b>Bleed occurrence during PCI</b>	1 (2.4)
<b>Bleed occurrence within 72 hours of procedure</b>	2 (5)
<b>Death due to MI/ACS/CVA during hospital stay</b>	2 (4.8)
<b>Death due to bleeding associated event during hospital stay</b>	0 (0)
<b>Readmission to Memorial due to MI/ACS within 30 days of discharge</b>	1 (2.5)
<b>Readmission to Memorial due to bleeding associated event within 30 days of discharge</b>	0 (0)

Table 4. Events not meeting bleed definition	Total, No. (%)
<b>Hemoglobin change</b>	
Hgb drop of greater than or equal to 3 g/dL with no identified source	7 (17)
<b>Adverse Events</b>	
Pseudoaneurysm without a Hgb drop greater than or equal to 3 g/dL	2 (4.8)
Retroperitoneal bleed without a Hgb drop greater than or equal to 3 g/dL	1 (2.4)

## Conclusions

- Only one patient received cangrelor without meeting approved criteria for use due to the patient being on an oral P2Y<sub>12</sub> inhibitor prior to admission
- Education to nursing staff is recommended regarding returning cangrelor vials not used, crediting patients' profiles and ensuring documentation of dosing is consistent
- Education to physician staff is recommended regarding appropriate criteria for use and infusion duration for cangrelor

## References

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