้ภาวะแทรกซ้อนภายหลังการทำศัลยกรรมขนาดเล็กในช่องปาก ในพู้ป่วยที่รับยาต้านการเกิดลิ่มเลือดชนิดรับประทาน

ปรีชา เจษฎาชัย* สุภาพร วิธิยะจิรกุล* วราวุธ อัมพรวิโรจน์กิจ** อุษาศิริ ศรีสกุล***

บทคัดย่อ

วัตถุประสงค์: เพื่อเปรียบเทียบอัตราการเกิดภาวะเลือดออก และอัตราการเกิดภาวะแทรกซ้อนทางระบบ จากการทำศัลยกรรมขนาดเล็กในช่องปาก ระหว่างผู้ป่วยที่หยุดยาและไม่หยุดยาต้านการเกิดลิ่มเลือด

วัสดุอุปกรณ์และวิธีการ: การศึกษาเชิงสังเกตการณ์แบบย้อนหลังจากเวชระเบียนผู้ป่วยที่โรงพยาบาล ตากสิน ระหว่าง มกราคม 2560 ถึง ธันวาคม 2561 โดยบันทึกข้อมูลในผู้ป่วยที่เข้าทำศัลยกรรมขนาดเล็ก ในช่องปาก และมีประวัติการหยุดยา หรือไม่หยุดยาต้านการเกิดลิ่มเลือด ก่อนทำหัตถการ และมีข้อมูลติดตาม การเกิดภาวะแทรกซ้อนหลังทำหัตถการ ได้แก่ ภาวะเลือดออก และโรคกล้ามเนื้อหัวใจขาดเลือด หรือโรคหลอด เลือดสมอง

ผลการทดลอง: จำนวนผู้ป่วย ทั้งหมด 574 คน มีอายุเฉลี่ย 66.89 ± 10.63 ปี และร้อยละ 54.20 เป็นเพศชาย ผู้ป่วยส่วนใหญ่มีประวัติเป็นโรคความดันโลหิตสูง (ร้อยละ 68.30) โรคเบาหวานชนิดที่ 2 (ร้อยละ 48.60) ใช้ยาแอสไพรินเดี่ยว (ร้อยละ 77.40) เข้ารับการถอนฟัน (ร้อยละ 92.70) และหยุดยาต้านการเกิด ลิ่มเลือดก่อนทำหัตถการ (ร้อยละ 54.18) ผลการวิเคราะห์พบว่า ผู้ป่วยที่ไม่หยุดยาต้านการเกิดลิ่มเลือด มีอุบัติการณ์เกิดภาวะเลือดออกหลังทำหัตถการใกล้เคียงกับผู้ป่วยที่หยุดยาต้านการเกิดลิ่มเลือด (ร้อยละ 1.14 และ 1.29 ตามลำดับ) และมีค่า adjusted OR เท่ากับ 0.446 (95% CI 0.080-2.494 และ p = 0.358) ในทางกลับกัน พบว่า ผู้ป่วยที่หยุดยาต้านการเกิดลิ่มเลือดมีแนวโน้มเล็กน้อยต่อการเกิดภาวะแทรกซ้อนทางระบบ หลังการทำ หัตถการ มากกว่าผู้ป่วยที่ไม่หยุดยาต้านการเกิดลิ่มเลือด (ร้อยละ 2.57 และ 1.90 ตามลำดับ) และมีค่า adjusted OR เท่ากับ 1.422 (95% CI 0.454–4.452 และ p = 0.545)

สรุป: การไม่หยุดยาต้านการเกิดลิ่มเลือดไม่เพิ่มความเสี่ยงต่อการเกิดภาวะเลือดออกหลังการทำหัตถการ อย่างไรก็ตาม การหยุดยาต้านการเกิดลิ่มเลือดอาจเพิ่มความเสี่ยงต่อการเกิดภาวะลิ่มเลือดอุดตันในหลอดเลือด ดังนั้น ควรประเมินความเสี่ยงและประโยชน์ของผู้ป่วยแต่ละรายอย่างรอบคอบก่อนพิจารณาการหยุดยาต้านการ เกิดลิ่มเลือดก่อนทำหัตถการ

คำสำคัญ: ยาต้านการเกิดลิ่มเลือด ยาต้านการแข็งตัวของเลือด ยาต้านเกล็ดเลือด ศัลยกรรมขนาดเล็กในช่องปาก ภาวะลิ่มเลือดอุดตันในหลอดเลือด ภาวะแทรกซ้อนทางระบบ

^{*}กลุ่มงานทันตกรรม โรงพยาบาลตากสิน 543 ถนนสมเด็จเจ้าพระยา แขวงคลองสาน เขตคลองสาน กรุงเทพมหานคร 10600 **กลุ่มงานอายุรกรรม โรงพยาบาลตากสิน 543 ถนนสมเด็จเจ้าพระยา แขวงคลองสาน เขตคลองสาน กรุงเทพมหานคร 10600 ***คณะเภสัชศาสตร์ มหาวิทยาลัยสยาม ถนนเพชรเกษม แขวงบางหว้า เขตภาษีเจริญ กรุงเทพมหานคร 10160

Postoperative Complications from Oral Minor Surgery in Patients on Oral Antithrombotic Drugs

Preecha Chesdachai* Supaporn Viriyajirakul* Warawut Umpornwirojkit** Usasiri Srisakul***

Abstract

Objectives: To compare of bleedings and systemic complications after minor oral surgery between patients with interrupted antithrombotic drugs (IAT) and continued antithrombotic drugs (CAT).

Materials and Methods: An observational retrospective cohort study retrieved from patient medical records at Taksin hospital between January 2017 to December 2018. Data were collected from patients who underwent oral minor surgery with a history of IAT or CAT perioperatively. Post-operative complications including bleeding, myocardial infarction, and stroke, were evaluated.

Results: A total of 574 patients, (54.2% male) with mean age of 66.89 \pm 10.63 years old. were included. In general, there were patients with hypertension (68.30%), type 2 diabetes mellitus (48.60%), aspirin monotherapy (77.40%), dental extraction (92.70%) and IAT (54.18%). The results showed that patients in the CAT group had a similar incidence of post-operative bleedings to the IAT group (1.14 vs. 1.29%, respectively) with an adjusted OR of 0.446 (95% CI 0.080–2.494, p = 0.358). In contrast, the IAT group had a slight trend without statistical significance towards post-operative systemic complications compared with CAT group (2.57 vs. 1.90%, respectively) with an adjusted OR of 1.422 (95% CI 0.454–4.452, p = 0.545).

Conclusions: Continuing the antithrombotic drugs did not increase the post-operative bleeding risk. However, interrupting the antithrombotic drugs may influence the risk of systemic complications. Therefore, any individual patient should be carefully assessed for the risks and benefits of antithrombotic drugs before consideration of perioperative antithrombotic interruption.

Keywords: Antithrombotic drug, Anticoagulants, Antiplatelets, Oral minor surgery, Thromboembolism, Systemic complication

^{*}Dental department, Taksin Hospital, 543 Somdet Chaopraya Road, Khlong San Sub-district, Khlong San District, Bangkok 10600, Thailand.

^{**}Internal Medicine Section, Taksin Hospital, 543 Somdet Chaopraya Road, Khlong San Sub-district, Khlong San District, Banakok 10600, Thailand.

^{***}Faculty of Pharmacy, Siam University, Petchkasem Road, Bang Wa Sub-district, Phasi Charoen District, Bangkok 10160, Thailand.

Introduction

Due to increasing life expectancy, patients who are taking oral antithrombotic drugs for prevention of cardiovascular events and undergoing dental extraction have been increased. Dental providers must consider the actual risks of post-operative bleedings and thromboembolic complications from any potential interaction between oral minor surgery and antithrombotic therapy. This is a dilemma for dental providers whether to continue or discontinue antithrombotic drugs perioperatively and what is the best decision-making process.

Aspirin (acetylsalicylic acid; ASA) is a nonsteroidal anti-inflammatory drug (NSAIDs). It also has an antiplatelet activity by inhibiting platelet aggregation that induced from thromboxane A2. Based on this function, it is indicated for primary or secondary prevention of cardiovascular diseases.(1,2) Clopidogrel is an antiplatelet drug that irreversibly binds to platelet purinergic P2Y12 receptor which is activated by adenosine diphosphate (ADP). Like ASA, clopidogrel is indicated mainly for secondary prevention of atherothrombotic diseases. (3) Warfarin, an oral vitamin K antagonist (VKA), prolongs prothrombin time by antagonizing the vitamin K-dependent clotting factors (II, VII, IX, X). (4) Warfarin is the most common oral anticoagulant used for prevention and treatment of thromboembolisms, such as atrial fibrillation, mechanical heart valve, and venous thromboembolism. (5,6)

The dental procedure is classified under a minor surgery, which the risk of thromboembolisms associated with discontinuation of the antithrombotic drugs is low and the risk of bleedings is far out-weight when continuing the antithrombotic drugs. (7-12) Warfarin has a long half-life of 48 to 72 hours and a delay-onset of 76 to 96 hours after cessation as well as re-initiation of the treatment. (13-21) Dual antiplatelet (DAPT) combined of low-dose ASA and one P2Y12 inhibitor (clopidogrel, ticagrelor or prasugrel), which is recommended as a first-line treatment for secondary prevention in patients with coronary artery disease. (22,23) There are studies in dental procedures concluded that the risk of thromboembolic events from discontinuation of ASA or DAPT is more harm than the risk of post-operative bleedings from the continuation of antiplatelets. (10,11) Additionally, discontinuation of ASA or DAPT particularly clopidogrel is not recommended in the setting of dental procedures. (24, 25)

This study aimed to compare the incidences of post-operative complications including bleedings and systemic complications between patients with interrupted antithrombotic drugs and continued antithrombotic drugs.

Materials and methods

Study design and population:

We conducted an observational retrospective cohort study by retrieving data from electronic medical records of all patients who aged at least 18 years old, underwent minor oral surgery, and received oral antithrombotic drugs between January 2017 to December 2018 at Taksin hospital. The study protocol was approved by the Bangkok metropolitan administration human research ethic committee (BMAHREC) No.S007h/ 63 22 May 2020. Patients were categorized into two groups as interrupted antithrombotic drug

(IAT) groups as and continued antithrombotic drug (CAT) group. Antithrombotic drugs in this study included antiplatelets (ASA or clopidogrel), or anticoagulant (warfarin). The IAT group referred to patients with discontinuation of an antithrombotic drug at least 3 days before minor oral surgery for ASA or clopidogrel, or at least 1 day for warfarin. The CAT group referred to patients with continuation of ASA, clopidogrel, or warfarin before and on the day of minor oral surgery without any interruption. The minor oral surgery was defined as tooth extraction, surgical removal, and torectomy. All patients were given a post-operative self-care instruction by continuing to bite on the gauze pack for 1 hour to stop bleeding and taking good care of oral hygiene. In the case of patients on warfarin, surgical wounds were applied with local hemostatic agents, such as oxidized regenerated cellulose, and sutured with silk 3-0.

Outcomes:

Primary outcome was frequency of postoperative bleeding defined as any level of bleedings that occurred immediately after surgical procedures or up to 3 days after surgery. Bleeding definition and classification were modified from the bleeding academic research consortium (BARC) (26,27), as followed: Level 1 is any bleeding that stopped spontaneously without any additional treatment; Level 2 is any bleeding that required additional treatment by a health care professional with materials or tools, such as stitches to stop the bleed within 10 minutes on an observation; Level 3 is any overt bleeding that required additional wound suture with hospitalization for blood transfusion and intravenous fluid resuscitation. Secondary outcomes were systemic complication events including myocardial infarction or stroke that occurred after dental surgery within 30 days or longer.

Statistical analysis:

The baseline characteristics of patients with IAT and CAT were analyzed using the independent student t-test for continuous variables and the Chi-square test for categorical variables. Multiple logistic regression was conducted for comparing the risk of bleedings or systemic complications after surgery between two groups and calculated for odds ratio (OR) with 95% confidence intervals at alpha value of 0.05 (twosided). Potential covariates were controlled in the models, such as age, sex, medical history, dental procedures, antithrombotic drugs. All the analyses were performed by a statistical computer software.

Results

Patients:

A total of 574 patients were identified with a history of minor oral surgery and a prescription of antithrombotic drugs. Followed-up data of post-operative complications were available for all patients. Antithrombotic drugs uses were categorized as IAT group in 54.18%, and CAT group in 45.82% of patients. The characteristics of age, sex, medical history, antithrombotic drug regimens with warfarin dose, international normalized ratio (INR), and dental procedures are shown in table 1.

| Characteristics | Total | IAT | CAT | p-value | |
|--|---------------|---------------|---------------|---------|--|
| | (n = 574) | (n = 311) | (n = 263) | | |
| Age – year (mean ± SD)* | 66.89 ± 10.63 | 66.70 ± 10.88 | 67.11 ± 10.34 | 0.640 | |
| Male sex - no. (%) [†] | 311 (54.20) | 164 (52.70) | 147 (55.90) | 0.449 | |
| Medical history – no. (%)† | | | | | |
| Type 2 diabetes mellitus | 279 (48.60) | 147 (47.30) | 132 (50.20) | 0.485 | |
| Hypertension | 392 (68.30) | 211 (67.80) | 181 (68.80) | 0.802 | |
| Coronary artery disease | 123 (21.40) | 72 (23.20) | 51 (19.40) | 0.274 | |
| Stroke | 117 (20.40) | 64 (20.60) | 53 (20.20) | 0.899 | |
| Atrial fibrillation | 45 (7.80) | 27 (8.70) | 18 (6.80) | 0.414 | |
| Heart valve disease | 18 (3.10) | 10 (3.20) | 8 (3.00) | 0.905 | |
| Hyperlipidemia | 217 (37.80) | 100 (32.20) | 117 (44.50) | 0.002 | |
| Renal failure | 41 (7.10) | 25 (8.00) | 16 (6.10) | 0.365 | |
| Neurologic disease | 109 (19.00) | 60 (19.30) | 49 (18.50) | 0.840 | |
| Antithromboticregimens – no. (%)† | | | | | |
| Aspirin | 444 (77.40) | 246 (79.10) | 198 (75.30) | 0.277 | |
| Clopidogrel | 25 (4.40) | 14 (4.50) | 11 (4.20) | 0.852 | |
| Warfarin | 71 (12.40) | 35 (11.30) | 36 (13.70) | 0.377 | |
| Aspirin + Clopidogrel | 25 (4.40) | 8 (2.60) | 17 (6.50) | 0.023 | |
| Aspirin + Warfarin | 9 (1.60) | 8 (2.60) | 1 (1.40) | 0.035 | |
| Warfarin (n = 80)* | | | | | |
| Dose (mg/week) - mean ± SD | 20.21 ± 9.50 | 22.47 ± 9.84 | 16.92 ± 8.07 | 0.021 | |
| INR - mean ± SD | 2.07 ± 0.86 | 2.05 ± 0.84 | 2.10 ± 0.90 | 0.800 | |
| Dental procedures – no. (%) [†] | | | | | |
| Tooth extraction | 532 (92.70) | 285 (91.60) | 249 (93.90) | 0.297 | |
| Surgical removal | 2 (0.30) | 1 (0.30) | 1 (0.40) | 0.905 | |
| Torectomy | 43 (7.50) | 27 (8.70) | 16 (6.10) | 0.234 | |

Table 1. Patient Characteristics at Baseline.

IAT: interrupted antithrombotic drugs; CAT: continued antithrombotic drugs; SD: standard deviation; no.: number of patients; INR: international normalized ratio; N/A: not available; *Independent t-test; [†]Chi-Square test.

The total mean age was 66.89 ± 10.63 years old, which was similar to the both groups (p = 0.640). Almost half of patients were male in both groups (p = 0.449). A significantly higher proportion of patients with CAT had hyperlipidemia (p = 0.002). Antithrombotic regimens seemed to be similar in both groups. There were slightly more patients in the CAT group had ASA combined with clopidogrel (p = 0.023), although the IAT group had a slightly higher proportion of ASA combined with warfarin (p = 0.035). Warfarin was prescribed in 80 patients (13.94%), 43 patients were in the IAT group and 37 patients were in the CAT group. In addition, the mean warfarin weekly dose in the IAT group was higher than the CAT group with a statistical difference (22.47 ± 9.84 vs. 16.92 ± 8.07 mg/week, respectively, p = 0.021). However, the mean INR was comparable between IAT and CAT groups $(2.05 \pm 0.84 \text{ and } 2.10 \pm 0.90, \text{ respectively},$ p = 0.800). More than 90 percent of patients in both groups had tooth extraction. There were similar proportions of patients in the both groups underwent surgical removal or torectomy.

Outcomes:

The Incidence of postoperative bleedings was reported in 7 patients (1.20%) as shown in table 2. Comparison of post-operative bleedings between the IAT and CAT groups was analyzed by using the Chi-square test. The result revealed no statistically significant difference between the IAT and the CAT groups [4/311 (1.29%) vs. 3/263 (1.14%), respectively, p = 0.874]. However, there was a trend for more pronounced post-opertaive bleedings in patients who interrupted warfarin (4/80 patients, 5.00%) than continued antiplatelets (1/503 patients, 0.20%), respectively. The most frequent bleedings were classified in level 2 (4/7 patients), 2 patients interrupted warfarin, 1 patient

continued warfarin, and another patient continued antiplatelets. All patients who experienced postoperative bleedings were received re-sutured and packed with surgicel. There were 2 patients who had bleedings in level 3 and were admitted into the hospital and given additional treatments with transamine mouth rinse and wound stenting. Additionally, there was 1 patient who continued warfarin and had an INR level of 2.00 before undergoing torectomy, experienced bleeding level 2 that was successfully managed by re-sutured, packed with surgicel, and transamine mouth rinse.

Incidence of total systemic complications after minor oral surgery was 2.26% (13/574). The incidence of stroke (7/574, 1.22%) and myocardial infarction (6/574, 1.05%) were similar as shown in table 2. All systemic complication events found in patients who were taking antiplatelets. The incidence of systemic complications in patients who interrupted antiplatelets compared with patients who continued antiplatelets was not statistically significant [8/503 (1.59%) vs. 5/503 (0.99%), respectively, p = 0.625]. The incidence of myocardial infarction in patients who interrupted antiplatelets was low, yet tended to be higher than patients who continued antiplatelets without statistically significant difference [4/503 (0.80%) vs. 2/503 (0.40%), respectively, p = 0.695]. The comparison of stroke events between patients who interrupted and continued antiplatelets was similar without statistical significance [4/503 patients (0.80%) vs. 3/503 patients (0.60%), respectively, p = 1.000]. For time to systemic complication events, mostly occurred in duration of longer than 30 days after minor oral surgery. The systemic complications that occurred within 30 days after procedure were myocardial infarctions which were recorded from 2 patients, one patient interrupted antiplatelets and another patient continued antiplatelets.

| | Aspirin or Clopidogrel | | | | Warfarin | | |
|-----------------------------|------------------------|-------------|-----------|---------|-------------|-----------|---------|
| Post-operative | Total | (| n = 503) | | | (n = 80) | |
| Complications | (n = 574) | Interrupted | Continued | p-value | Interrupted | Continued | p-value |
| Bleedings – no. (%)* | 7 (1.20) | 0 | 1 (0.20) | 0.451 | 4 (5.00) | 2 (2.50) | 0.681 |
| Bleeding level – no. (%)*† | | | | 0.451 | | | 0.776 |
| Level 1 | 1 (0.20) | 0 | 0 | | 1 (1.25) | 0 | |
| Level 2 | 4 (0.70) | 0 | 1 (0.20) | | 2 (2.50) | 1 (1.25) | |
| Level 3 | 2 (0.35) | 0 | 0 | | 1 (1.25) | 1 (1.25) | |
| Dental procedures - no. (%) | * | | | | | | |
| Tooth extraction | 6 (1.05) | 0 | 1 (0.20) | N/A | 4 (5.00) | 1 (1.25) | 0.333 |
| Surgical removal | 0 | 0 | 0 | N/A | 0 | 0 | N/A |
| Torectomy | 1 (0.20) | 0 | 0 | N/A | 0 | 1 (1.25) | 0.333 |
| Bleeding Management*- | | | | | | | |
| no. (%)* | 7 (1.22) | 0 | 1 (0.20) | 0.451 | 4 (5.00) | 2 (2.50) | 0.681 |
| Re-sutured | 7 (1.22) | 0 | 1 (0.20) | 0.451 | 4 (5.00) | 2 (2.50) | 0.509 |
| Pack with surgicel | 2 (0.35) | 0 | 0 | N/A | 1 (1.25) | 1 (1.25) | 1.000 |
| Transamine mouth rinse | 2 (0.35) | 0 | 0 | N/A | 1 (1.25) | 1 (1.25) | 1.000 |
| Systemic Complications - | | | | | | | |
| no. (%)* | 13 (2.26) | 8 (1.59) | 5 (0.99) | 0.625 | 0 | 0 | N/A |
| Myocardial infarction - | | | | | | | |
| no. (%)* | 6 (1.05) | 4 (0.80) | 2 (0.40) | 0.695 | 0 | 0 | N/A |
| After procedure < 30 days | 2 (0.35) | 1 (0.20) | 1 (0.20) | 0.713 | 0 | 0 | N/A |
| After procedure > 30 days | 4 (0.70) | 3 (0.60) | 1 (0.20) | | 0 | 0 | N/A |
| Stroke – no. (%)* | 7 (1.22) | 4 (0.80) | 3 (0.60) | 1.000 | 0 | 0 | N/A |
| After procedure < 30 days | 0 | 0 | 0 | 1.000 | 0 | 0 | N/A |
| After procedure > 30 days | 6 (1.05) | 3 (0.60) | 3 (0.60) | | 0 | 0 | N/A |
| Unknown | 1 (0.20) | 1 (0.20) | 0 | | 0 | 0 | N/A |

Table 2. Incidence of Post-operative Complications and Antithrombotic Drugs.

no.: number of patients; N/A: not available; *Chi-Square test or Fisher's Exact test as appropriate; †Postoperative bleeding level – level 1 is bleeding that stopped spontaneously without any additional treatment; Level 2 is bleeding that required additional treatment by a health care professional with materials or tools such as stitches to stop the bleed within 10 minutes observation; Level 3 is overt bleeding that required additional wound suture with hospitalization for blood transfusion and intravenous fluid resuscitation. In the table 3 showed multivariate analysis of post-operative complications. Results found that the CAT group was not associated with an increased risk of bleedings after minor oral surgery (adjusted OR 0.446, 95% Cl 0.808–2.494, p = 0.358). In contrast, the IAT group had a small trend of increased systemic complications without statistical significance (adjusted OR 1.422, 95% CI 0.454– 4.452, p = 0.545). Similarly, the systemic complication risks in patients with IAT had small trends without statistical significances to increase myocardial infarction (adjusted OR 1.583, 95%CI 0.279–8.986, p = 0.604), and stroke (adjusted OR 1.241, 95%CI 0.265–5.814, p = 0.784).

| Post-operative Complications | No. (%) | OR* | Adjusted OR [†] | p-value |
|------------------------------|-----------|-----------------------|--------------------------|---------|
| | | (95% CI) | (95% CI) | |
| Bleedings | | | | |
| Total (n = 574) | 7 (1.20) | | | |
| Interrupted (n = 311) | 4 (1.29) | 1.129 (0.250 – 5.091) | 1.227 (0.213 – 7.058) | 0.819 |
| Continued (n = 263) | 3 (1.14) | 0.886 (0.196 - 3.993) | 0.446 (0.080 - 2.494) | 0.358 |
| Systemic complications | | | | |
| Total (n = 574) | 13 (2.26) | | | |
| Interrupted (n = 311) | 8 (2.57) | 1.362 (0.440 – 4.216) | 1.422 (0.454 – 4.452) | 0.545 |
| Continued (n = 263) | 5 (1.90) | 0.734 (0.237 – 2.271) | 0.682 (0.218 – 2.138) | 0.511 |
| Myocardial infarction | | | | |
| Total (n = 574) | 6 (1.05) | | | |
| Interrupted (n = 311) | 4 (1.29) | 1.700 (0.309 – 9.358) | 1.583 (0.279 – 8.986) | 0.604 |
| Continued (n = 263) | 2 (0.76) | 0.588 (0.107 - 3.237) | 0.597 (0.104 – 3.425) | 0.563 |
| Stroke | | | | |
| Total (n = 574) | 7 (1.20) | | | |
| Interrupted (n = 311) | 4 (1.32) | 1.129 (0.250 – 5.091) | 1.241 (0.265 – 5.814) | 0.784 |
| Continued (n = 263) | 3 (1.14) | 0.886 (0.196 - 3.993) | 0.815 (0.172 – 3.865) | 0.797 |

Table 3. Multivariable Analysis for Post-operative Complications.

No.: number of patients; OR: odds ratio; CI: confidence interval; *simple logistic regression; †Multiple logistic regression with adjusted for antithrombotic drugs, dental procedures, age, sex, and medical history.

Univariate analysis of post-operative bleedings as shown in table 4. The analyses demonstrated that warfarin was a statistically significant risk factor that associated with increased post-operative bleedings (OR 46.338, 95% CI 5.492-390.990, p < 0.001), which was stronger than ASA (OR 0.471, 95% CI 0.006 -0.391, p = 0.005). Interestingly, atrial fibrillation demonstrated astatistically significant association with increased post-operative bleedings (OR 9.375, 95% CI 2.031-43.277, p = 0.004). This result might be influenced by the highly frequent warfarin use for atrial filbrillation. There were 3 post-operative bleeding events from patients with atrial fibrillation and taking warfarin. Overall, the INR levels revealed no statistically significant association with post-operative bleedings. However, when compared with INR in subtherapeutic levels of less or equal to 1.99 as reference, the INR levels in therapeutic ranges of 2.00 to 2.49 seemed to have a slightly higher risk of postoperative bleedings without statistical siginificance (OR 6.643, 95% CI 0.634–69.621, p = 0.114) than the INR in supratherapeutic levels of higher or equal to 3.00 (OR 6.200, 95% CI 0.332-115.918, p = 0.222), and the INR in therapeutic levels of 2.50 to 2.99 (OR 2.385, 95% CI 0.138-41.079, p = 0.550), respectively. In addition, comorbid conditions that found no association with postoperative bleedings included heart valve disease, hyperlipidemia, and chronic kidney disease.

| Risk Factors | Bleeding (n = 7) | | leeding = 567) | 0R [†] (95% CI) | p-value |
|---|---------------------|-----|-------------------|-----------------------------|---------|
| Continued antithrombotic drug - no. (%) | 3 (42.86) | 260 | (45.86) | 0.886 (0.196–0.993) | 0.874 |
| ASA | 1 (14.29) | | | | |
| Warfarin | 2 (28.57) | | | | |
| Dental procedures - no. (%) | | | | | |
| Tooth extraction | 6 (85.71) | 526 | (92.77) | 0.468 (0.055–3.978) | 0.487 |
| Surgical removal | 0 | 2 (| (100.00) | N/A | N/A |
| Torectomy | 1 (14.29) | 42 | (7.42) | 2.079 (0.245–17.677) | 0.503 |
| Antithrombotic regimens – no. (%) | | | | | |
| Aspirin | 1 (14.29) | 443 | (78.13) | 0.471 (0.006–0.391) | 0.005 |
| Clopidogrel | 0 | 25 | (4.41) | N/A | N/A |
| Warfarin | 6 (85.71) | 65 | (11.46) | 46.338 (5.492-390.990) | < 0.001 |
| Aspirin + Clopidogrel | 0 | 25 | (4.41) | N/A | N/A |

| Table 4. Univariate | Analysis | for Post-operative | Bleedings. |
|---------------------|----------|--------------------|------------|
|---------------------|----------|--------------------|------------|

| Risk Factors | Bleeding | No Bleeding | g OR [†] | p-value |
|---------------------------|-------------------|------------------|-----------------------|---------|
| | (n = 7) | (n = 567) | (95% CI) | |
| Warfarin | (n = 6) | (n = 63) | | |
| INR - mean ± SD | 2.55 ± 1.39 | 2.02 ± 0.79 | 1.819 (0.783–4.229) | 0.164 |
| INR level – no. (%) | | | | |
| <1.99 | 1 (16.67) | 31 (49.21) | reference | |
| 2.00 - 2.49 | 3 (50.00) | 14 (22.22) | 6.643 (0.634–69.621) | 0.114 |
| 2.50 – 2.99 | 1 (16.67) | 13 (20.63) | 2.385 (0.138–41.079) | 0.550 |
| > 3.00 | 1 (16.67) | 5 (7.94) | 6.200 (0.332–115.918) | 0.222 |
| Warfarin dose (mg/week) - | | | | |
| mean ± SD | 20.60 ± 10.99 | 20.16 ± 9.46 | 1.005 (0.913–1.106) | 0.923 |
| Age – year (mean ± SD) | 63.57 ± 7.09 | 66.93 ± 10.66 | 0.973 (0.914–1.037) | 0.402 |
| Male sex - no. (%) | 5 (71.43) | 306 (53.97) | 0.469 (0.090–2.437) | 0.368 |
| Medical history - no. (%) | | | | |
| Type 2 diabetes mellitus | 1 (14.29) | 278 (49.03) | 0.173 (0.021–1.448) | 0.106 |
| Hypertension | 3 (42.86) | 389 (68.61) | 0.343 (0.076–1.550) | 0.164 |
| Coronary artery disease | 1 (14.29) | 122 (21.52) | 0.608 (0.073–5.098) | 0.646 |
| Stroke | 0 | 117 (20.63) | N/A | N/A |
| Atrial fibrillation | 3 (42.86) | 42 (7.41) | 9.375 (2.031–43.277) | 0.004 |
| Heart valve disease | 1 (14.29) | 17 (3.00) | 5.392 (0.615–47.289) | 0.128 |
| Hyperlipidemia | 4 (57.14) | 213 (37.57) | 2.216 (0.491–9.997) | 0.301 |
| Chronic kidney disease | 1 (14.29) | 40 (7.05) | 2.196 (0.258–18.687) | 0.472 |
| Neurologic disease | 1 (14.29) | 108 (19.05) | 0.708 (0.084–5.945) | 0.751 |

Table 4. (cont.)

OR: odds ratio; CI: confidence interval; no.: number of patients; INR: international normalized ratio; SD: standard deviation; reference: bleeding events from patients with INR < 1.99 (the lowest range) was set as a comparator when calculated OR for the higher INR levels; N/A: not available; [†]simple logistic regression.

The univariate analysis of post-operative systemic complications as shown in table 5. We found no statistically significant association of interrupted antiplatelets to increase the post-operative systemic complications (OR 1.362, 95% CI 0.440-4.216, p = 0.592). However, ASA tended to have a higher risk of post-operative systemic complications without statistical significance

(OR 3.583, 95% CI 0.462-27.820, p = 0.222) than clopidogrel (OR 1.865, 95% CI 0.233-14.934, p = 0.557). Additionally, there were some variables that found no associations with increased postoperative systemic complications included IAT, ASA, clopidogrel, type 2 diabetes mellitus, hypertension, coronary artery disease, heart valve disease, and hyperlipidemia.

| Risk Factors | Systemic | No Systemic | ort | p-valu |
|--|------------------------------------|---------------|----------------------|--------|
| | Complication Complication (95% CI) | | | |
| | (n = 13) | (n = 561) | | |
| Interrupted antithrombotic drug-no.(%) | 8 (61.54) | 260 (54.01) | 1.362 (0.440-4.216) | 0.592 |
| Aspirin | 7 (58.33) | | | |
| Clopidogrel | 1 (7.69) | | | |
| Antithrombotic regimens – no. (%) | | | | |
| Aspirin | 12 (92.31) | 432 (77.01) | 3.583 (0.462-27.820) | 0.222 |
| Clopidogrel | 1 (7.69) | 24 (4.28) | 1.865 (0.233-14.934) | 0.557 |
| Warfarin | 0 | 71 (12.66) | N/A | N/A |
| Aspirin + Clopidogrel | 0 | 25 (4.46) | N/A | N/A |
| Aspirin + Warfarin | 0 | 9 (1.60) | N/A | N/A |
| Dental procedures – no. (%) | | | | |
| Tooth extraction | 13 (100.00) | 519 (92.51) | N/A | N/A |
| Surgical removal | 0 | 2 (0.36) | N/A | N/A |
| Torectomy | 0 | 43 (7.68) | N/A | N/A |
| Male sex – no. (%) | 9 (69.23) | 302 (53.83) | 0.518 (0.158-1.703) | 0.279 |
| Age – year (mean ± SD) | 64.46 ± 9.947 | 66.94 ± 10.65 | 0.980 (0.933-1.028) | 0.403 |
| Medical history – no. (%) | | | | |
| Type 2 diabetic mellitus | 8 (61.54) | 271 (48.31) | 1.712 (0.553-5.298) | 0.351 |
| Hypertension | 9 (69.23) | 383 (68.27) | 1.046 (0.318-3.441) | 0.941 |
| Coronary artery disease | 3 (23.08) | 120 (21.39) | 1.102 (0.299-4.069) | 0.884 |
| Stroke | 1 (7.69) | 116 (20.68) | 0.320 (0.041-2.484) | 0.276 |
| Atrial fibrillation | 0 | 45 (8.02) | N/A | N/A |
| Heart valve disease | 1 (7.69) | 17 (3.03) | 2.667 (0.328-21.697) | 0.359 |
| Hyperlipidemia | 6 (46.15) | 211 (37.61) | 1.422 (0.472-4.287) | 0.532 |
| Chronic kidney disease | 0 | 41 (7.31) | N/A | N/A |
| Neurologic disease | 1 (7.69) | 108 (19.25) | 0.350 (0.045-2.717) | 0.315 |

Table 5. Univariate Analysis for Post-operative Systemic Complications.

OR: odds ratio; no.: number of patients; SD: standard deviation; N/A: not available; [†]simple logistic regression.

Discussion

Our analyses demonstrated that the relationship of CAT and post-operative bleeding was not statistically significant (adjusted OR 0.446, 95% CI 0.080-2.494, p = 0.358). This finding supported by Yang Set al. 2016 (28) conducted a meta-analysis included 6 randomized controlled studies or controlled clinical studies in 591 patients taking oral anticoagulants and underwent dental extractions. The results revealed that patients who continued oral anticoagulants had a non-significant risk of post-operative bleedings compared with patients who discontinued oral anticoagulant therapy (risk ratio 1.31, 95% CI 0.79–2.14).

Interesitngly, we found that warfarin was a strong risk factor for post-operative bleedings (p < 0.001). In contrast, ASA was less likely to cause bleedings post-operatively (p = 0.005). Our results supported by Lu et al., 2018 reported that continuation of warfarin had a strong trend of higher incidence of bleeding after dental extraction than a continuation of single or dual and antiplatelet regimens (9.1% and 1.82%, respectively, p = 0.0599) (29). Results from Bajkin et al., 2014 showed that patients who continued dual antiplatelet regimen had a 1.7% incidence of bleeding after dental extraction, and no bleeding was reported in continuation of single antiplatelet treatment (30). Furthermore, we were the first to report the association of atrial fibrillation as a statistically significant comorbid condition to increase post-operative bleedings after oral minor surgery (OR 9.375, 95% CI 2.031-43.277, p = 0.004). However, the data showed with a low precision due to the wide range of 95% confidence intervals and should be determined in a larger study.

This study showed trends of INR in the therapeutic levels towards post-operative bleedings without statistical significance. These results are supported by previous studies, Phanpaisan et al., 2015 They demonstrated that the incidences of post-operative bleeding were comparable between patients with INR in therapeutic and supratherapeutic levels (4.88% and 3.66%, respectively) (31). Morimoto et al., 2008 showed that patients with target INR levels had a higher incidence of bleedings than patients with out-of-target INR levels (10.09% and 2.50% respectively) and reported the INR was not correlated to post-operative bleedings (p = 0.06) (12).

Our analyses demonstrated that an interruption of antithrombotic drugs had a trend towards post-operative systemic complications without statistical significance (adjusted OR 1.422, 95% CI 0.454-4.452, p = 0.545). Although the total incidence of systemic complications was low as 2.26% and all the events occurred in patients who taking antiplatelets. We found that ASA had a trend without statistical significance (OR 3.583, 95% CI 0.462-27.820, p = 0.222) to increase systemic complications than clopidogrel (OR 1.865, 95% CI 0.233-14.934, p = 0.557). Maulaz et al., 2005 reported from patients in a stroke unit that the interruption of ASA at least 3 days was associated with an increased risk of ischemic stroke or transient ischemic attack (OR 3.40, 95% Cl 1.08-10.63) (32). There are several previous studies reported that discontinuing the antiplatelet drugs can increase the risk of thromboembolic events particularly myocardial infarction, stent thrombosis, and stroke. (33-39) Therefore, overestimation of severe post-operative bleeding risk from dental extraction which

considered as low potential, may increase the risk of thromboembolic event and potentially harm to develop consequences of morbid complications.

Interestingly, there was no systemic complication reported in patients who interrupted warfarin in our study, which might due to longer half-life of warfarin compared to antiplatelets. (13–21) However, the systemic complications that occurred in duration of longer than 30 days after procedure were likely to be confounded from other unmeasured factors than the interruption of antiplatelets for minor oral surgery.

Limitations

The mean age of patients in this study was relatively old, which may not reflect the younger population. As an observational retrospective study, potential confounding factors that were unable to control or measure may exist, leading to underestimate or overestimate the outcomes particularly other clinical factors related to bleedings or systemic complications. Moreover, the quality and variety of data in medical charts or missing data could impact the analyses. In addition, an influent factor that was unable to measure is patient adherence to interrupt or continue the antithrombotic drugs as recommended by physicians or dentists, as well as the adherence to re-initiate the antithrombotic therapy. This study included torectomy, a higher risk of bleeding than tooth extraction, but our result showed no association with post-operative bleeding that might due to low incidence. Because the antithrombotic drugs in this study are limited to ASA, clopidogrel, and warfarin, the results could not be extrapolated to newer

agents such as ticagrelor, prasugrel, or direct oral anticoagulants (DOAC). Finally, the limited number of patients and low incidences of postoperative complications might affect the precision of statistical results, which should be examined in a prospective and larger study.

Conclusions

Our findings demonstrated that the incidence of postoperative bleeding is low and not related to continuation of antithrombotic drugs, and more frequent in patients taking warfarin. Interruption of antithrombotic drugs was not related to the occurence of systemic complications. Although, the low incidence of systemic complications was reported, the result showed a slight trend without statistical significance of antiplatelet interruption towards systemic complications that should be determined in the future study. Continuing education should be encouraged to carefully evaluate an individual patient for bleeding and thrombosis risks before considering of interruption the antithrombotic therapy.

Acknowledgments

The authors would like to express our appreciation to our colleagues, and patients at the dental department, Taksin hospital.

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ติดต่อบทความ

ทพ.ปรีชา เจษฎาชัย กลุ่มงานทันตกรรม โรงพยาบาลตากสิน 543 ถนนสมเด็จเจ้าพระยา แขวงคลองสาน เขตคลองสาน กรุงเทพมหานคร 10600, โทรศัพท์: 089 128 2310 อีเมล์: Chesdachai42@gmail.com

Corresponding author:

Dr. Preecha Chesdachai Dental department, Taksin Hospital 543 Somdet Chaopraya Road, Khlong San Sub-district, Khlong San District, Bangkok 10600, Tel: (668) 9128 2310 E-mail: Chesdachai42@gmail.com

Received Date: Apr 30, 2020 Revised Date: May 12, 2020 Accepted Date: Jul 20, 2020