A Clinical Appraisal of Post-operative Bleeding in Routine Extractions in Patients on Uninterrupted Low-dose Aspirin Therapy- An in Vivo Study

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Abstract

Antiplatelet drugs are recommended globally for long term prevention of serious vascular events in high risk patients. Temporary withdrawal of these drugs seems to be an attractive option in dental office prior to minor surgical procedures to prevent a bleeding hazard. The benefit of averting anti-platelet drug induced hemorrhage needs to be balanced against the risk of potentially fatal thrombosis that may occur due to rebound phenomenon after withdrawal of the anti-platelet drug. This in-vivo prospective study was thus designed to assess post-operative bleeding in patients on aspirin therapy after extractions. The study sample comprised of 50 patients between 50 to 75 years of age on long term aspirin therapy requiring extractions were assessed for bleeding at 30 mins, 1, 2, 3, 24 and 48 hours post operatively. Results showed 4 patients had mild bleeding at 30 mins, 2 patients at 1hour, 1 patient at 2 hours and no patients with bleeding at 3 hours. In our study we had an 8% risk of bleeding after extractions in patients on long term low dose aspirin therapy. Given the low incidence and severity of bleeding which can be easily managed by local measures only, we conclude that patients need not discontinue taking aspirin prior to dental extractions.

Keywords: Acetylsalicylic acid, tooth extraction, exodontia, post-operative bleeding.

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INTRODUCTION

Advances in science and technology have contributed to ease the workload on mankind. Present day sedentary and physically inactive lifestyles, combined with poorly executed diets and habits like smoking precipitate lifestyle related diseases amongst which hypertension and cardiovascular diseases are most common. Atherosclerotic plaques build up along the lining of blood vessels over many years in response to injury caused by high blood pressure, abnormal blood sugar levels, high blood cholesterol levels and toxins contained in tobacco [1-3]. Lipid-rich atherosclerotic plaques are vulnerable and their rupture can cause the formation of a platelet and fibrin-rich thrombus leading to myocardial infarction or a cerebrovascular accident [4].

In normal physiologic state a dynamic equilibrium exists between coagulation-anticoagulation and fibrinolysis- antifibrinolysis. Disturbance in this equilibrium with shift favouring any of the mechanism will result in either bleeding or thrombosis [5].

Medical science has always been in a state of flux and with increasing research, newer medicines are being discovered to combat disease, thus increasing the lifespan of human beings. Antiplatelet agents like aspirin, clopigrdogrel, ticloidipine, cilostazol, dypyriramole, ticagrelor and prasugrel have been extensively researched and are used routinely for the prevention of arterial thrombosis in patients with conditions such as ischemic heart disease, prosthetic heart valves and coronary artery stents as well as those at risk of experiencing ischemic cerebrovascular accidents [6-8].

Although antiplatelet drugs are extensively used in cardiovascular diseases, it also increases the potential risk of bleeding particularly when performing minor surgical procedure like dental extraction [9].
fear of uncontrolled or excessive bleeding prompts clinicians to stop or alter these drugs before surgical procedures [10-12]. Stopping antiplatelet therapy may expose patients to the risk of thromboembolism, myocardial infarction or cerebrovascular accidents [13-26]. However, considerable debate has been generated with regard to balancing the risk of a post-surgical hemorrhage with that of precipitating a thromboembolic event [27].

Temporary withdrawal of anti-platelet therapy therefore seems an attractive option in dental office wherein minor surgical procedure may pose a bleeding hazard. However, the benefit of averting anti-platelet drug induced hemorrhage needs to be balanced against the risk of potentially fatal thrombosis that may occur due to rebound phenomenon after anti-platelet drug withdrawal [28-32]. Thus the following study was conducted to evaluate the bleeding following dental extraction in patients who are on long term low dose aspirin therapy.

MATERIALS AND METHODS

Our study was approved by the Institutional Ethics Committee of our Institute. We followed the World Declaration Guidelines of Helsinki. 50 patients referred to our Department of Oral and Maxillofacial Surgery indicated for extraction of teeth between the age group of 50 –75 years on long term aspirin therapy (75 mg – 150 mg per day) were selected for the study. A written informed consent was obtained from all the study participants.

Patients, with teeth in the area having local infection, systemic conditions like liver or kidney disease and acquired or congenital bleeding disorders with a potential for bleeding, history of gastrointestinal bleeding or intracranial hemorrhage, undergoing vascular surgery, on anticoagulant therapy like warfarin, heparin, enoxaparin, allergic to lignocaine or any local anesthetics, diabetes mellitus, bone marrow disorders and pregnancy were excluded from the study.

All the patients were evaluated at 30 minutes, 1 hour, 2 hours, 3 hours after extractions for bleeding in our Department of Oral and Maxillofacial Surgery. On discharge from the clinic these patients were followed up telephonically. They were also recalled back to the clinic 24 hours and 48 hours post-operatively for follow up.

To record the intra-oral bleeding, the extraction site was observed without gauze piece in situ. At the end of 30 minutes and onwards any bleeding that extended beyond the crest of the socket (i.e., onto surrounding gingival tissues) during an observation period of 1 min was considered as bleeding present. The bleeding record was maintained using the following bleeding score.

<table>
<thead>
<tr>
<th>SCORING PATTERN FOR BLEEDING</th>
</tr>
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<tbody>
<tr>
<td>Score 0</td>
</tr>
<tr>
<td>Bleeding absent</td>
</tr>
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A detailed case history was obtained and details of the procedure were explained to the patient & the relatives. Pre-operative blood pressure was checked with manual sphygmomanometer by auscultatory method. Pre-operative investigations-blood sugar level, bleeding time, clotting time, platelet count, prothrombin time (PT), international normalized ratio (INR) were done. Bleeding time was checked using Duke’s method and clotting time by Capillary Tube method. Physician’s consent was obtained for every patient in lieu of blood pressure and cardiovascular status. An informed consent was obtained regarding the surgical procedure and follow up time for 3 hours and 24 hours and 48 hours post-operatively. Pre-operative intraoral periapical radiographs were obtained. Preoperative patient’s oral cavity was prepared using 2% povidone-iodine germicidal gargle and routine barrier protection techniques were used by the operating surgeon. After following all the necessary surgical aseptic protocol, patients were prepared and draped for the surgical procedure. 15% lignocaine topical anesthetic spray was applied on the mucosa of area to be injected. Pterygomandibular nerve block was given using of 2% lignocaine hydrochloride without adrenaline for mandibular teeth. Local infiltration injection was used for maxillary teeth. The subjective and objective signs and symptoms of the anesthesia were confirmed. Extraction of teeth was performed as per routine.

After extraction the socket was curetted and was compressed for two minutes. The patients were then asked to bite hard on a piece of sterile gauze for 30 minutes and re-evaluated for bleeding upto 3 hours post-operatively at regular intervals. Patients were discharged after 3 hours with a family member after giving routine post-operative instructions.

All patients were covered post-operatively with Amoxycillin-Trihydrate capsules 500mg 8 hourly and paracetamol tablets 500mg 8 hourly for five days.

RESULTS

Out of 50 patients, 29 were male and 21 were female. Age ranged from 50-75 years with a mean age of 59.90±7.11 years. The bleeding time of the patients ranged from 1.25 minutes to 2.85 minutes and clotting time ranged from 3.13 minutes to 5.53 minutes. Both the bleeding time and clotting time were within the normal range. Mean bleeding and clotting time were 1.89±0.43 minutes and 4.05 minutes respectively. In our study we observed bleeding beyond the crest of the socket in 4 patients at 30 minutes, out of which bleeding stopped in 2 patients after 30 minutes. 2 patients continued to bleed for 1 hour after extraction. In 1 of these patients, bleeding stopped after 1 hour, the
2nd patient experienced bleeding 2 hours post-operatively. Bleeding in all patients was controlled using a pressure pack.

Graph 1 show that no patients had any bleeding after 3 hours, 24 and 48 hours. None required blood transfusion, hospitalization for bleeding or major cardiovascular event.

Chi square test was used to show an association between bleeding and time interval \[P=0.046\]. Since \(P<0.05\), it indicates significant association between the status of bleeding and time point measurements. It implies that bleeding is absent as time progresses.

Graph 2 – Showing percentage distribution of bleeding with respect to time post-operatively

Graph-3: Shows that the overall risk of bleeding was 8% in our study

Graph-3: Showing the trend line in the Bar Chart

DISCUSSION

Excessive post-extraction bleeding is defined as continuous bleeding from the extraction socket beyond the post-operative window of clot formation period [33]. Excessive bleeding in the patients is not only distressing for the patient, but also hinders the completion of the procedure and can compromise wound healing [34-38]. Lifestyle related diseases like hypertension and other cardiovascular diseases are invariably associated with increased blood levels of lipids, cholesterol and free radicals which facilitate the formation of atherosclerotic plaque causing lumpy thickening of the blood vessel, narrowing of the arterial lumen and slowing down the blood flow which leads to the formation of thrombus [39-41].

Since platelets have an important role in coagulation and, in particular, arterial thrombosis, anti-platelet drugs are in widespread use for the prevention of morbidity and mortality from vascular disease [42].

Complete or near-complete inhibition of platelet cyclo-oxygenase [COX-1] can be achieved with low doses of aspirin (75-150 mg) given once daily. In contrast, inhibition of COX-2-dependent pathophysiologic processes (eg, hyperalgesia, inflammation) requires larger doses of aspirin and a much shorter dosing interval because nucleated cells rapidly re-synthesize the enzyme. Thus, 10- to 100-fold higher daily doses of aspirin are required when the drug is used as an anti-inflammatory agent rather than as an anti-platelet agent. Lemkin et al. [43] and Mc Gaul et al. [9] have documented that there was an increased postoperative bleeding after dental extraction and recommended to discontinue aspirin. For most elective surgeries, few authors have recommended that the patient should stop taking aspirin 7 to 10 days before the procedure [44]. Some authors recommended discontinuing aspirin 3 days prior to the surgery since the platelet.

Inhibition achieved with aspirin, although irreversible for target platelets, lasts until a significant pool of new platelets is synthesized and complete recovery of platelet aggregation may occur in 50% of cases by day 3 and in 80% of cases by day 4 [45]. Some
investigators even suspect the existence of a biological platelet aggregation “rebound phenomenon”. Rapid withdrawal of aspirin may cause abnormally high levels of blood markers reflecting an increase of thromboxane -A2 which may have possible hazardous effects in patients with cardiovascular disease [46]. The delicate balance between adequate antithrombotic effect and risk of bleeding remains a sensitive matter, which often influences a patient's or a physician's choice to discontinue daily anti-platelet therapy[47].

The best screening test for the effect of aspirin on coagulation is the platelet function analyzer (PFA-100) [13]. This test mimics the clotting process in vitro and allows for a more accurate determination of platelet function [44]. If this is not available, then the bleeding time can be used. Although aspirin affects platelets and the coagulation process through its effect on platelet release, it does not usually lead to a significant bleeding problem unless the bleeding time is greater than 20 minutes. If surgery is to be performed under emergency conditions and the bleeding time is in excess of 20 minutes, 1-desamino-8-D-arginine vasopressin (DDAVP) can be used to shorten the bleeding time [13].

In our study, patients were evaluated at 30 minutes, 1 hour, 2 hours, 3 hours, 24 hours and 48 hours from the time of extraction for bleeding. To record the intraoral bleeding, the extraction site was observed without gauze in place. Any bleeding that extended beyond the crest of the socket (i.e., onto surrounding gingival tissues) during an observation period of 1 minute was considered as bleeding present. In our study we observed bleeding beyond the crest of the socket in four patients at 30 minutes out of which bleeding stopped after 30 minutes in two patients. Two patients continued to bleed for 1 hour after extraction. None of the patients had bleeding after 2 hours, 3 hours, 24 hours and 48 hours. There were no cases of transfusion, hospitalization for bleeding or major cardiovascular events. The consequences of possible hemorrhage in non-compartment surgeries is greatly outweighed by the risk associated with cessation of anti-platelet therapy, which can result in acute coronary syndrome in serious cases [39].

Burger et al. [48] stated that, in patients on aspirin, the average risk of intra-operative bleeding increases by a factor of 1.5 [45]. In our study the overall risk of bleeding observed after extractions in patients on aspirin therapy was 8%. Probably the few cases of bleeding reported in our study could be contributed to the fact that in our study lignocaine was used without adrenaline and suturing was not included in our dental extraction protocol.

The American College of Chest Physicians Evidence-Based Clinical Practice Guidelines 2012 made the following recommendations, like in patients who are receiving ASA for the secondary prevention of cardiovascular disease and are having minor dental or dermatologic procedures or cataract surgery, we suggest continuing ASA around the time of the procedure instead of stopping ASA 7 to 10 days before the procedure (Grade 2C-high risk patients ).stopping anti-platelet agents is NOT recommended prior to most dental procedure. Dent! !ts should obtain a medication history including any currently prescribed antiplatelet agents and when last taken. Given the low incidence and severity of bleeding following dental surgery, use of local measures (e.g. absorbable gelatin sponges and sutures) is adequate to control bleeding. Elective procedures with significant risk of bleeding should be postponed [13].

On the basis of our study and above recommendations we conclude that patients need not discontinue taking aspirin prior to dental extractions.

**CONCLUSION**

The main objective of our study was to evaluate the risk of bleeding in patients on long term aspirin therapy undergoing dental extractions.

In our study we had a 8% risk of bleeding after extractions in patients on long term low dose aspirin therapy. These patients were treated as any other normal patients and local measures were used to achieve hemostasis.

This study highlights that patients who are on long term low dose aspirin therapy need not discontinue their medications prior to dental extractions. Considering the small risk of post-operative bleeding it will be prudent to pack the extraction socket with absorbable gel foam and place single interrupted sutures to achieve hemostasis postoperatively.

**REFERENCES**


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