A New Technique to Induce Experimental Myointimal Hyperplasia

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Significance of the Study

- In this paper, we describe a new innovative technique to induce experimental myointimal hyperplasia using a dental flossing cachet. The technique is simple, inexpensive, and highly reproducible. It would be of interest to researchers studying the pathogenesis of myointimal hyperplasia.

Keywords

Angioplasty · Carotid stenosis · Intimal injury · Myointimal hyperplasia · Neointima formation

Abstract

\textbf{Background:} Arterial myointimal hyperplasia (MIH) has a significant impact on the long-term outcomes of vascular procedures such as bypass surgery and angioplasty. In this study, we describe a new and innovative technique to induce MIH using a dental flossing cachet in Wistar rats. 

\textbf{Methods:} The intimal damage in the common carotid artery was induced by inserting the tip of the dental flossing cachet through the external carotid artery into the common carotid artery and turning it on for 3 rounds of 20 s each (n = 10). After 2 weeks, the rats were anesthetized and the common carotid arteries of the experimental side and the contralateral side (control) were harvested and preserved for histopathological studies. 

\textbf{Results:} The experimental carotid arteries showed significant intimal proliferation and thickening compared to the controls. The intima/media ratio of the experimental and normal (control) common carotid arteries were 1.274 ± 0.162 and 0.089 ± 0.023 (mean ± SEM), respectively (p < 0.001).

\textbf{Conclusion:} This technique is simple, inexpensive, and highly reproducible and it induces sufficient MIH to study this phenomenon in animal models.

Introduction

The development of arterial myointimal hyperplasia (MIH) has a significant impact on the long-term outcomes of vascular procedures such as bypass surgery and angioplasty. Experimentally induced MIH in animal models is used for the study of the pathogenesis of this...
condition and to investigate ways to prevent or decrease its development. Most studies have used balloon angioplasty in animal models to induce MIH [1–3]. However, these methods are inherently tedious and require the use of one balloon catheter for each procedure, with the attached cost. In addition, these models were found not to be standardized because the pressure within the balloon may vary depending on the operator’s maneuver (i.e., the balloon is not inflated equally within the same carotid, and it is not the same in different applications) [4–8]. We report a new technique to induce MIH using a dental flossing cachet, which is not expensive compared to the balloon and can be easily purchased. Unlike experimental balloon angioplasty, which entails the complete denudation of the intima from the carotid artery, this technique is automated and induces injury only to the intima by the rotating tip of the flossing cachet, which would stimulate the endothelial cells to initiate the process of intimal hyperplasia, mimicking the clinical procedures of angioplasty or bypass anastomosis.

**Materials and Methods**

This study was approved by the Animal Protection Ethics Committee. Ten inbred Wistar rats (body weight ≥ 400 g) were maintained at the Animal Resources Centre of the Health Science Center, Kuwait University. The animals had free access to rat chow and tap water and were maintained in a standard 12-h light/dark cycle; the temperature was maintained at 25 ± 2 °C.

![Fig. 1. a Power Flosser device. b Technique of inducing myointimal hyperplasia in the carotid artery. The dotted ties on the common carotid artery (CCA) and the internal carotid artery (ICA) are the stay sutures used to temporarily interrupt the blood flow during the procedure. The loose tie (blue) at the origin of the external carotid artery (ECA) is used to tie the artery after completion of the procedure. The rotating tip is seen within the CCA via an incision in the ECA just beyond its take off from the CCA. For better clarity in the diagram, a space is shown between the rotating tip and the wall of the CCA; in real life the device is introduced with some difficulty into the lumen of the CCA and it sits snugly within the lumen. Upon starting the machine, the floss tip is in firm contact with the intima and thus makes the required damage.](image-url)
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as a control by dissecting it up to the angle of the jaw and by the application of loose stay sutures at the proximal and distal ends of the artery, which were ultimately removed before wound closure. At the end of the procedure, the wound was washed with warm sterile normal saline, and the retracted muscles were returned to their normal anatomic position in the midline. The skin incision was closed with continuous 3/0 silk sutures, and the animals were kept warm until recovery, transferred to a designated cage, and allowed free access to food and water.

After 2 weeks, the animals were anesthetized and the neck wounds opened. The experimental site was exposed first, and the CCA was dissected all along to its bifurcation. A stay suture was passed around the proximal CCA at the root of the neck. This suture was tied, immediately followed by injection of normal saline (using an insulin needle and syringe) into the CCA to flush out the blood. The CCA, along with its bifurcation, was then harvested and fixed in formalin. The contralateral (control) CCA was likewise flushed, harvested, and fixed. Finally, the animals were euthanized via an overdose of anesthesia. The tissue was fixed in 10% formalin and embedded in paraffin blocks, and cut and cross-sections of the harvested arteries were stained with a standard hematoxylin and eosin stain.

Morphometric Analysis

Digital images of the harvested arteries were captured using a computerized imaging system and analyzed. Morphological analysis was performed by tracing the interior surface of the lumen, the internal elastic lamina, and the external elastic lamina (EEL). The area inside of the internal elastic lamina is the intima, and the area between IEL and EEL is the media. Smooth muscle cell proliferation and migration (blue arrows) lead to neointima formation and an increase in the intima/media ratio (magnification ×40).

Statistical Analysis

All data are presented as means ± SEM. Comparison of histological findings between the control and experimental CCA groups was made via an unpaired Student t test. p < 0.05 was considered statistically significant. All statistical analyses were performed with GraphPad Prism version 6.0 (GraphPad Software Inc., San Diego, CA, USA).

Results

All rats appeared healthy and were moving all limbs before sacrifice, and no adverse effects were observed. All of the neck incisions healed well, with no infections observed.
The cross-section of the experimental carotid arteries (Fig. 2a, c) shows impressive intimal proliferation and thickening compared to the control carotid arteries (Fig. 2b, d). The MIH was noted to extend from the site of the insertion of the flosser’s tip in the ECA (Fig. 3) down to the whole length of the CCA, i.e., the site of intimal contact with the rotating tip of the flosser. Figure 3 shows the bifurcation of the CCA, in which the ECA lumen is nearly completely occluded due to the MIH. Figure 4 shows that an extensive MIH was induced in the experimental carotid arteries as indicated by the higher intima/media ratios (1.274 ± 0.162) compared to the control common carotid arteries (0.089 ± 0.023) (mean ± SEM); this difference was highly significant (p < 0.001).

Discussion and Conclusion

Experimental MIH has been induced previously in rat carotid arteries using a balloon catheter; however, this technique does not mimic the real-world situation because it denudes the whole intimal layer, which does not occur during clinical angioplasty [1–3]. Moreover, the small Fogarty (balloon) catheter is used only once per animal and it is therefore expensive. We opted to explore the use of a new method to induce MIH in the rat carotids, which would mimic the clinical situation (i.e., injury to the arterial intima) rather than denuding the whole intima.

In our study, the automated flossing tip induced impressive intimal proliferation and thickening compared to the control. To prove the reproducibility and validity of the technique, the procedure was performed on 10 rats by 2 coauthors (A.S. and F.A.-O.). In all animals, MIH was consistently induced without variability along the CCA. In comparison, many reports have shown that the balloon injury model in rats is not standardized because the value of the pressure in the balloon is not automated and may vary depending on the operator’s maneuver [4–8]. Thus, our model of intimal injury is simple, inexpensive, and highly reproducible and induces sufficient MIH to study this phenomenon in the animal model.

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