Intimal hyperplasia, the obstacle in bypass grafts
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CHAPTER 4

Inhibition of vein graft intimal and medial thickening by periadventitial application of a sulfated carbohydrate polymer

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ABSTRACT

Purpose: The purpose of this study was to determine whether the wall thickening observed in vein grafts after they were placed into the arterial circulation could be inhibited by periadventitial delivery of an insoluble sulfated polymer of β-cyclodextrin (P-CDS) capable of tightly binding heparin binding growth factors.

Methods: Thirty-four New Zealand White rabbits underwent implantation of reversed autologous jugular vein interposition grafts in the common carotid artery and were randomized to receive either 20 mg P-CDS (n=18) topically around the graft or no additional therapy (n=16). Before being killed at 28 days, animals were given bromodeoxyuridine to assess smooth muscle cell proliferation. Histomorphometric analyses were performed after perfusion fixation.

Results: Compared to controls, treatment with P-CDS was associated with reduced mean intimal thickness (24±3 vs 38±4 μm; mean SEM, p<0.01) and intimal area (0.25±0.03 vs 0.54±0.09 mm; p<0.01). There was also significantly less medial thickness in the P-CDS group (45±3 vs 63±3 μm, p<0.001). There was no significant difference in intimal or medial smooth muscle cell proliferation between P-CDS treated and control vein grafts at 28 days. The polymer persisted in the adventitia of all treated vein grafts with a mild foreign body reaction.

Conclusions: Periadventitial placement of a novel, insoluble, sulfated carbohydrate polymer, inhibits intimal and medial thickening of vein bypass grafts in this model of vein grafting. The persistence of P-CDS in vivo for prolonged periods, and the ease of topical application of P-CDS during vascular bypasses may have important implications for future use in vascular surgery.
INTRODUCTION

Vein grafts inserted into the arterial circulation develop intimal and medial thickening soon after implantation. Although wall thickening may be an appropriate healing process of the vein to new conditions and forces in the arterial environment, it may also result in pathologic lumen narrowing finally resulting in vein graft failure. Vein graft wall thickening is considered to be a leading cause of late vein graft failure. Recent studies demonstrate the pivotal role of growth factors in the formation of intimal thickening after arterial injury, growth factors may also play an important role in the normal physiologic condition and repair mechanisms of the vessel wall. The healing process of autologous vein grafts implanted into the arterial circulation is likely to be, at least in part, in response to growth factor-mediated stimulation. Therefore agents that interact or interfere with growth factor activity may be beneficial for controlling the wall thickening observed after vein grafts are placed in the arterial circulation.

Heparin has been shown to inhibit proliferation and migration in vitro and inhibits intimal thickening in models of arterial injury, although systemic continuous administration at higher levels is usually required. However, heparin’s action on smooth muscle cell growth appears independent of its anticoagulant function. To avoid problems with systemic administration, local administration to the periadventitial space has also successfully used to inhibit intimal thickening after arterial injury but often requires implantation of delivery vehicle.

To provide periadventitial therapy without an additional delivery vehicle, we developed an insoluble, sulfated carbohydrate polymer of β-cyclodextrin (P-CDS) that has several properties that make it an attractive candidate to prevent intimal thickening associated with vein graft arterilization: (1) it has no anticoagulant properties, (2) it is stable for prolonged periods when implanted periadventitially in animals without toxicity, (3) it has the capacity to tightly bind heparin-binding growth factors (HBGF), and (4) it inhibits smooth muscle cell proliferation and migration in vitro.

The purpose of this study was to determine whether periadventitial therapy with P-CDS would inhibit vein graft wall thickening after implantation in a well-characterized rabbit model.

METHODS

Rabbit vein graft model. Thirty-four male New Zealand White rabbits (Hazelton Research Denver, PA) weighing 3.0 to 3.5 kg at the time of the operation were anesthetized by intramuscular injection with a solution composed of ketamine (50 mg/kg) and xylazine (7 mg/kg). The rabbits underwent vein bypass graft interposition in the left carotid artery. Surgery was performed under ster-
ile conditions. A ventral midline incision was made in the neck and the left jugular vein and common carotid artery were carefully exposed and dissected free from surrounding tissues. Heparin (1000 U; Sima Chemical Co., St. Louis, Mo.) was given intravenously. The flow through the artery was interrupted with microvascular clips and then cut 15 mm segment of the external jugular vein was excised, reversed, and interposed into the divided artery. Proximal and distal anastomoses were performed end to end by the use of interrupted 7-0 polyprolene sutures (Ethicon, Inc., Somerville, N.Y.). After graft implantation the experimental group received P-CDS as described below. The skin was closed with 2-0 polyglactin suture.

Treatment group groups. Rabbits were randomized to treatment with P-CDS (n=18) or to control group (n=16) that underwent similar graft implantation and manipulation without polymer placement. P-CDS was provided as a dry, white powder by American Maize, Inc (Hammonton, Ind). With the vein graft in place, P-CDS was sprinkled topically to coat the graft. After placement of the dry polymer, three drops of normal saline solution were added to the treated area, making a gel-like substance to assure that the entire circumference of the vein graft and proximal and distal anastomoses had adequate contact with the polymer. The P-CDS and saline solution mixture has a consistency much like wet sugar, and, because it has no architectural strength, it did not prevent the vein graft from distention of the vein graft. All animal care was complied with the “Principles of Laboratory Animal Care,” and the “Guide for the care and Use of Laboratory Animals,” (NIH Publication No.80-23, revised 1985).

Harvesting procedure. Four weeks after vein graft implantation, at 17, 9, and 1 hour before being killed, the animals received 3 doses (1ml/kg) of 30mg/ml 5-bromo-2’deoxy-uridine solution (Boehringer Mannheim, Indianapolis, Ind). The rabbits were anesthetized and intravenously injected with Evans blue dye (25 mg/kg) and heparin (1000 units). Laparotomy was performed for placement of an abdominal aortic perfusion catheter and an inferior vena cava drainage catheter. The animals were killed, and the vascular system was washed out with lactated Ringer’s solution via a retrograde cannula in the thoracic aorta. Perfusion fixation was performed with 10% formalin in neutral 0.1 M phosphate buffer at a pressure of 100 mm Hg in the standard fashion, with the vein graft undisturbed in situ. After perfusion fixation, with the head, neck, and upper thorax fixed, the left common carotid artery with the vein graft, and segments of the right carotid artery and the right jugular vein were removed and immersion fixed for an additional 48 hours.

Structure. Samples of the midsection of the vein graft were used for histologic,
morphometric and immunohistochemical analyses. Specimens were embedded in paraffin, cut in 4 µm sections and stained with hematoxylin and eosin, Verhoeff van Gieson stain, or Richardson’s stain. Morphometric measurements were obtained with computer based representation measuring. (Biometrics, Nashville, In.). The inner intimal boundary was the luminal surface, and the intima-media boundary was identified by the color gradient with the Verhoeff van Gieson stain and the Richardson stain. The outer boundary was defined by the perivascular capillaries. The thickness of the vein graft wall compartments were determined at 8 separate, equally spaced sites along the circumference of the vein. These values were averaged. The vein graft wall circumference was measured, and the cross-sectional vein wall compartment areas were identified as the product of the perimeter and the corresponding compartment thickness. Alternatively, sections were used for immunohistochemistry to detect BrdU-positive cells as described previously.\(^1\)\(^3\)

These sections were counterstained with hematoxylin-and-eosin. The number of BrdU-positive cells and the total number of cells per unit area in both the intima and media of control and P-CDS treated vein grafts were counted manually. An optical graticule was used, and nuclei were counted per unit area at eight even intervals around the circumference. The intimal and medial boundaries were determined by comparison the correspond-
The distribution of intimal thickening present in the vein grafts was somewhat variable. The P-CDS vein grafts showed more uniformity of intimal thickening. Four P-CDS experiments showed a striking reduction in intimal thickening, and five had moderate to low intimal thickening. However, one of the P-CDS grafts displayed severe intimal thickening.

In both groups the smooth muscle cells in the intima appeared to be arranged in an irregular pattern with extracellular matrix between the cells. The media of control and P-CDS-treated vein grafts underwent concentric thickening and was less cellular in nature than the intima. Verhoeff van Gieson staining demonstrated elastin in the intima of control and P-CDS-grafts, with a predominance of collagen but not elastin staining in the media.

The light microscopical appearances of the cells and the extracellular matrix in P-CDS and control grafts showed some variability and no consistent qualitative differences at 28 days. There was no histologic evidence of toxic effects of P-CDS in terms of endothelial cell and smooth muscle cell structure or extracellular matrix appearance, and multinucleated giant cells were seen around the P-CDS particles, suggestive of a mild foreign body reaction. Evans blue staining revealed 100% confluent endothelial lining in all
Finally, we assessed the degree of smooth muscle cell proliferation at the time the animals were killed (28 days). At this time point, the overall degree of proliferation (~5%) was clearly higher than normal control jugular vein (~1%). However, there was no significant difference in the percentage of intimal and medial smooth muscle cell that were proliferating between P-CDS-treated and control vein grafts (Table I).

**DISCUSSION**

In this study we have demonstrated that periadventitial delivery of polymeric β-cyclodextrin sulfate significantly reduces the development of wall thickening in experimental vein bypass grafts at 28 days. This is the first report of locally applied polymer, which itself both therapeutic and nontoxic to reduce experimental vein graft wall thickening.

Wall thickening develops in all vein grafts within a short period after implantation into the arterial circulation. Vein graft intimal hyperplasia has been associated with vein graft stenosis and is a leading cause of late vein graft failure. Human autologous vein bypass graft studies and experimental vein graft models have shown that the healing process consists of vascular SMC undergoing an early proliferative phase followed by a synthetic phase resulting in a rapid intimal thickening. Smooth muscle cell proliferation and migration into the intima are consid-
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Considered as the major processes in the formation of vein graft intimal thickening. It is postulated that the healing process after vein graft implantation is driven by growth factors derived from vein graft smooth muscle cells and endothelium cells. This hypothesis is strengthened by the observation that in vein grafts in rabbits SMC and endothelium cells continue to proliferate in the absence of detectable endothelial denudation or platelet accumulation. It has been suggested that in particular heparin binding growth factors (HBGF) play a pivotal role in vessel remodelling after vascular injury. It is known that heparin and related glycosaminoglycans bind avidly to these HBGF and may affect their function. Heparin, P-CDS, and other heparin-like compounds, such as of β-cyclodextrin tetradecasulfate, inhibit SMC proliferation and migration in vitro and inhibit intimal thickening in animal models of arterial injury. Systemic administration of heparin, however, has yielded conflicting results with respect to its effect on wall thickening in experimental vein grafts. Makhoul et al reported that continuous intravenous administration of heparin significantly reduced intimal thickening in the rabbit carotid vein graft.

Kohler et al, using the same model of vein grafting, reported that heparin failed to reduce vein graft intimal thickening or SMC proliferation. Conflicting data of the effect of systemic heparin on vein graft wall thickening were also reported in studies using rat vein graft models. It is postulated that the mechanisms of intimal thickening between arterial injury model and vein graft model differ. Structural and functional differences between veins and arteries may also explain why certain therapeutic regimens succeeded in reducing intimal hyperplasia in models of arterial injury but failed to reduce intimal hyperplasia in models of vein grafting. Therefore the results of successful reduction of intimal hyperplasia in arterial injury models can not directly be extrapolated to models of vein grafting.

Multiple agents have been studied for their effects on vein graft remodeling with variable success. Mechanistically, blockade of growth factor function is an attractive option. Systemic administration of growth factor antagonists such as heparin may be harmful, since growth factors are widespread and not limited to the vasculature. Wound healing in general

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<th>Control</th>
<th>P-CDS</th>
<th>p</th>
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<tbody>
<tr>
<td>Media</td>
<td>0.04 ± 0.008</td>
<td>0.05 ± 0.01</td>
<td>NS</td>
</tr>
<tr>
<td>Intima</td>
<td>0.06 ± 0.01</td>
<td>0.04 ± 0.006</td>
<td>NS</td>
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Table I. Cell proliferation indexes of the media and intima of vein grafts at 28 days.
could be impaired, presenting a problem after surgical procedures. Furthermore, the usefulness of heparin is limited by its potential bleeding complications and its higher dose requirements for inhibiting SMC proliferation than the dose required for anticoagulation.

This stimulated the search for novel heparin mimics with the antiproliferative heparin characteristics of heparin but that could be applied locally. Local application of a pharmacologic compound may reduce its systemic effects and allow high concentrations at the target side. Monomeric β-cyclodextrin tetradecasulfate (CDS) is a universal heparin analog, paralleling and exceeding the demonstrated cell modulating capabilities of heparin without significant anticoagulant activities. We have previously shown that orally administered of this compound inhibits restenosis and intimal hyperplasia in a hypercholesterolemic rabbit angioplasty model. The polymer from P-CDS has biologic and therapeutic activity in that it inhibits SMC proliferation and migration in vitro and inhibits the arterial wall response to injury.

In this study, the polymer was applied to the adventitial side of the vein graft and both anastomoses in the form of dry, fine powder. Hydration of the polymer yields a soft gel that has no architectural strength and does not interfere with vessel dilatation. After hydration the polymer offers an internal heparin-like surface capable of absorbing HBGF proteins like basis fibroblast growth factor (bFGF), platelet-derived growth factors or vascular endothelial growth factor. Although the mechanism through which P-CDS reduces arterial and vein graft wall thickening is still unknown, we speculate that P-CDS may provide a sustained therapeutic effect by tightly binding and sequestering growth factors in the periadventitial space and that the cyclic structure of the carbohydrate polymer subunits would render it relatively resistant to degradation. Previous studies have demonstrated that the ability of P-CDS to tightly bind bFGF and prevent proliferation of rat SMC in vitro by bFGF, platelet derived growth factor-BB and epidermal growth factor. Whether growth factors generated in the wound site are sequestered and prevented from gaining access to their cognate receptors remains to be investigated. However, wound response in general was not impaired by the presence of this polymer in this model. At 28 days after injury there was no significant difference in medial and intimal proliferation indexes between control and P-CDS treated vein grafts. This is concordant with a study by Schwartz et al., who reported that smooth muscle cell proliferation in this model of vein grafting appears to begin within 24 hours and to peak at 2 days but then returns to normal at 14 days. Future studies are needed to determine whether P-CDS limits the early SMC proliferative burst soon after the implantation into the arterial circulation, which may in part account for the reduction in
intimal thickening.

In summary, this is the first report of locally applied novel heparin-like polymer without an additional delivery vehicle to reduce experimental vein graft wall thickening. The ease of local application of this polymer at the time of vascular surgery, especially at the anastomoses where intimal thickening is a particular problem, will facilitate its application in human vascular bypasses. Further investigations into its mechanism of action seem warranted to optimize its utility as a therapeutic agent.

References

Inhibition of vein graft intimal and medial thickening


