Distal Venous Arterialization and Reperfusion Injury: Focus on Oxidative Status

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Abstract

In patients with unreconstructable arterial occlusive disease distal venous arterialization (DVA) seems to be a promising option in the treatment. The goals of this prospective study were to assess clinical efficiency and possible impact of DVA on tissue damage by estimating oxidative status of patients with critical limb ischemia treated with this procedure. The subjects were 60 randomized patients: 30 were undergoing DVA and 30 were treated with antiaggregation therapy. During the mean follow-up period (6.13 ± 4.32 months for DVA vs. 6.74 ± 0.5 months for antiaggregation therapy) survival (p < 0.01), limb salvage (p < 0.001), pain relief (p < 0.001) and wound healing (p < 0.001) rates were significantly different between the two groups of patients in favor of the DVA group. Ten minutes after declamping we observed a decreasing trend in the lactate level in the blood of the deep venous system (p < 0.001). Also, on postoperative day 7 digital systolic pressure and digital-brachial index were higher than before the operation (p < 0.001). In blood samples collected immediately before and successively at 1, 3, 5 and 10 min postoperatively, prooxidative status (thiobarbituric acid reactive substances, \( \text{O}_2^- \), \( \text{H}_2\text{O}_2 \) and nitric oxide) and antioxidative enzymes (superoxide dismutase, catalase and glutathione reductase) were determined spectrophotometrically. Using the nonparametric Friedman test, we noted statistically nonsignificant differences (p > 0.05) in values of both prooxidative parameters and enzymes of the antioxidative defense system, before and successively at 1, 3, 5 and 10 min after operation. These results indicate that there was no statistically significant reperfusion injury after revascularization, which could have been expected after this surgical procedure, thus confirming its validity in these patients.

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Despite confirmation of good early and late effects of revascularization by using distal venous arterialization (DVA) which do not only occur in specific cases, it is not a procedure which is regularly used by the modern-day vascular surgeon [2–5]. However, the development of DVA could move forward in the direction of researching the hemodynamics of the recipient microvascular bed, neovascularization and usage of the remodeled venous bed as an approach in adjuvant therapy, such as prostanooids and angiogenic growth factors. One of the stumbling blocks may be a risk of ischemic tissue damage after reperfusion.

On the other hand, an imbalance between oxidants and antioxidants, in favor of the oxidants potentially leading to damage, is termed 'oxidative stress' [6]. Oxidants are formed as a normal product of aerobic metabolism but can be produced at elevated rates under pathophysiological conditions. Antioxidant defense involves several strategies, both enzymatic and nonenzymatic ones. It is well known that during the ischemic period ATP is catabolized to yield hypoxanthine. During reperfusion, molecular oxygen is reintroduced into the tissue where it reacts with hypoxanthine and xanthine oxidase to produce a burst of superoxide anion and hydrogen peroxide. In the presence of iron, superoxide anion and hydrogen peroxide react via the Haber-Weiss reaction to form hydroxyl radicals. This highly reactive and cytotoxic radical then initiates a lipid peroxidation of cell membrane components, releasing various proinflammatory agents and consequently tissue damage [7].

Since there is a real possibility of reperfusion injury during revascularization, we would like to draw attention to the essential aim of this procedure – tissue reoxygenation. Regarding the fact that there are almost no articles covering this topic, we believe that this prospective randomized study might give us an answer as to whether DVA achieves useful oxygenation or worsens ischemia.

**Material and Methods**

From September 2009 to April 2011 at the Clinic for Vascular and Endovascular Surgery (Clinical Center of Serbia in Belgrade), 60 patients were treated for critical limb ischemia without the option for arterial reconstruction as a result of peripheral arterial occlusive disease with patent aortiolic segment. From an unconstructable cohort we chose 30 suitable patients for treatment with DVA. The remaining 30 patients were treated with the antiaggregation therapy (100 mg aspirin per day). Randomization was performed for sex, age, stage of disease, incidence of diabetes, hypertension and smoking.

For the determination of oxidative stress parameters blood samples of all 30 patients were collected (from the popliteal or femoral vein of the revascularized limb) immediately before and at 1, 3, 5 and 10 min postoperatively. The prooxidative status was estimated by determination of the index of lipid peroxidation (measured as thiobarbituric acid reactive substances, TBARS), superoxide anion radical (O₂⁻), hydrogen peroxide (H₂O₂) and nitric oxide (NO) in plasma. The antioxidative defense system was estimated by determination of antioxidative enzymes in erythrocytes: superoxide dismutase (SOD), catalase (CAT) and glutathione reductase (GSH). All parameters were determined spectrophotometrically (Analytic Jena Specord S 600).

**Surgical Technique**

The operations were performed under regional anesthesia with prophylactic antibiotics and intravenous heparin administration. The in situ technique was used in most of the cases. It consisted of the following steps: after dissection of the median marginal vein of the foot and great saphenous vein, we ligated all tributaries caudally up to the ankle and created lateroterminal anastomosis between the great saphenous vein and the in-flow artery. Valvulotomy was performed by introducing an expandable LeMaitre valvulotome through the more distal segment of the median marginal vein. Reverse vein bypass graft with the position of a distal anastomosis on the junction of the dorsal venous arch and the superficial vein of the thumb was used in 6 patients. In this way arterial blood flow was made possible in two directions, cranially, without valve resistance up to the ligated trunk of the great saphenous vein at the level of the medial malleolus, and caudally through the dorsal venous arch after valvulotomy with a Fogarty catheter and a probe.

**Biochemical Assays**

For the determination of oxidative stress parameters blood samples of all 30 patients were collected (from the popliteal or femoral vein of the revascularized limb) immediately before and at 1, 3, 5 and 10 min postoperatively. The prooxidative status was estimated by determination of the index of lipid peroxidation (measured as thiobarbituric acid reactive substances, TBARS), superoxide anion radical (O₂⁻), hydrogen peroxide (H₂O₂) and nitric oxide (NO) in plasma. The antioxidative defense system was estimated by determination of antioxidative enzymes in erythrocytes: superoxide dismutase (SOD), catalase (CAT) and glutathione reductase (GSH). All parameters were determined spectrophotometrically (Analytic Jena Specord S 600).

**Determination of Index of Lipid Peroxidation (TBARS)**

Lipid peroxidation product malondialdehyde (MDA) concentration in plasma is determined by measuring TBARS using 1% thiobarbituric acid in 0.05 NaOH, incubated with plasma at 100°C for 15 min and read at 530 nm [8]. At high temperature and low pH value, MDA reacts with 2-thiobarbituric acid via nucleophilic addition. The product of this reaction is a colored substance whose concentration of MDA correlates with the color intensity of this mixture. Distilled water was used as a blank probe. Thio-barbituric acid extract was obtained by combining 0.8 ml plasma and 0.4 ml trichloroacetic acid. After that samples were put on ice for 10 min, and centrifuged for 15 min at 6,000 rpm.

**O₂⁻ Determination**

Determination of O₂⁻ plasma concentration was measured using a spectrophotometric method based on the reaction of O₂⁻ with nitro-blue tetrazolium in Tris buffer which formed nitro-blue formazan [9]. Absorbance is registered at 550 nm.
The protocol for measurement of H₂O₂ in plasma is based on oxidation of phenol red in the presence of horseradish peroxidase [10]. There were combined (1:20) 200-µl samples with 800-µl phenol red solution and 10-µl horseradish peroxidase (POD). The level of H₂O₂ was measured at 610 nm.

Determination of NO in Plasma

NO decomposes rapidly to form stable metabolite nitrate (NO₃⁻)/nitrate products. The method for detection of the plasma nitrate and NO₃⁻ levels is based on the Griess reaction. NO₃⁻ was determined as an index of NO production with Griess reagent [11]. 0.1 ml 3 N perchloric acid, 0.4 ml 20 mmol/l ethylenediaminetetraacetic acid (EDTA) and 0.2 ml plasma were put on ice for 15 min, then centrifuged for 15 min at 6,000 rpm. After pouring off the supernatant, 220 µl K₂CO₃ was added. NO₃⁻ was measured at 550 nm. Distilled water was used as a blank probe.

Determination of Antioxidant Enzymes

Isolated RBCs were washed 3 times with 3 vol of ice-cold 0.9 mmol/l NaCl. Hemolysates containing about 50 g Hb/l (prepared according to McCord and Fridovich [12]) were used for the determination of CAT activity. CAT activity was determined according to Beutler [13]. Lysates were diluted with distilled water (1:7 v/v) and treated with chloroform-ethanol (0.6:1 v/v) to remove hemoglobin [14]. Then 50 µl CAT buffer, 100-µl samples and 1 ml 10 mmol/l H₂O₂ were added to the samples. Detection was performed at 360 nm. Distilled water was used as a blank probe. SOD activity was determined by the epinephrine method of Misra and Fridovich [15]. 100 µl lysate and 1 ml carbonate buffer were mixed, and then 100 µl of epinephrine were added. Detection was performed at 470 nm.

Statistical Analysis

Analysis of the data was performed with the statistical evaluation, using Fisher’s exact test, Student’s t test, multivariate analysis of variance and Friedman’s nonparametric test with the support and use of computer program SPSS version 12.0 for Windows.

Results

In the experimental and control group, the majority of the patients were male with an average age of 65 with gangrene or unhealed painful ulceration of the foot. They had significant accompanying comorbidity. In both groups, one to two thirds of the patients had at least one of the following risk factors such as diabetes, hypertension and smoking. Around 30–40% of the patients had carotid and/or coronary artery disease.

Primary features of the patients and comorbidity are shown in tables 1 and 2.

To investigate the short-term clinical outcome of DVA we observed survival, limb salvage, pain relief and wound healing. All of these parameters were improved after DVA compared with results in the patients treated conservatively (fig. 1).

One of the ways to assess success of the surgical revascularization of the extremity is measurement of the segmental arterial blood pressure. For the purposes of this study 7 days after DVA a digital systolic pressure on the foot and a digital-brachial index were measured by using the Doppler machine. The use of the Student t test showed that there were no statistically significant differences in

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**Table 1. Demographic characteristics and risk factors of the patients**

<table>
<thead>
<tr>
<th>Group</th>
<th>Surgical procedure</th>
<th>Medication</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>21</td>
<td>17</td>
<td>0.211</td>
</tr>
<tr>
<td>Female</td>
<td>9</td>
<td>13</td>
<td></td>
</tr>
<tr>
<td>Age (mean ± SD)</td>
<td>65.37 ± 10.87</td>
<td>65.93 ± 11.27</td>
<td>0.844</td>
</tr>
<tr>
<td>Stage of disease¹</td>
<td>9</td>
<td>14</td>
<td>0.144</td>
</tr>
<tr>
<td>III</td>
<td>21</td>
<td>16</td>
<td></td>
</tr>
<tr>
<td>IV</td>
<td>11</td>
<td>15</td>
<td>0.217</td>
</tr>
<tr>
<td>Hypertension</td>
<td>21</td>
<td>17</td>
<td>0.211</td>
</tr>
<tr>
<td>Diabetes</td>
<td>16</td>
<td>13</td>
<td>0.303</td>
</tr>
<tr>
<td>Smoking</td>
<td>6</td>
<td>8</td>
<td>0.381</td>
</tr>
<tr>
<td>Hyperlipidemia</td>
<td>6</td>
<td>5</td>
<td>0.500</td>
</tr>
<tr>
<td>Obesity</td>
<td>6</td>
<td>5</td>
<td>0.500</td>
</tr>
</tbody>
</table>

¹ Stage of disease by Fontaine.

**Table 2. Comorbidity**

<table>
<thead>
<tr>
<th>Group</th>
<th>Surgical procedure</th>
<th>Medication</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coronary artery disease</td>
<td>12</td>
<td>12</td>
<td>0.604</td>
</tr>
<tr>
<td>Carotid artery disease</td>
<td>9</td>
<td>12</td>
<td>0.294</td>
</tr>
<tr>
<td>Renal failure</td>
<td>5</td>
<td>4</td>
<td>0.500</td>
</tr>
<tr>
<td>COPD</td>
<td>5</td>
<td>4</td>
<td>0.500</td>
</tr>
</tbody>
</table>

COPD = Chronic obstructive pulmonary disease.
Distal Venous Arterialization and Reperfusion Injury

preoperative values of digital pressures (table 3) between the groups of patients studied (p > 0.05).

A significant increase of digital systolic pressure values at 68.4 ± 16.21 mm Hg (p < 0.001) and digital-brachial index at 0.487 ± 0.149 (p < 0.001) was detected after surgical foot revascularization by using DVA (fig. 2, 3).

Immediately before revascularization, i.e. arterial clamping, a high lactate level in the blood of the deep venous system with the gradual fall in values within the first 10 min after the surgery, from 2.43 ± 0.49 to 1.143 ± 0.329 mmol/l, was registered in patients treated with DVA, as can be seen in figure 4.

The usage of the multivariate analysis of variance showed statistically significant difference (p < 0.001) not only between the registered preoperative value and every new measurement at 1, 3, 5 and 10 min postoperatively, but also after each successive measurement.

Table 3. Preoperative digital systolic pressure (DSP) and digital-brachial index (DBI)

<table>
<thead>
<tr>
<th>Pressure</th>
<th>Group</th>
<th>n</th>
<th>Mean</th>
<th>SD</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>DSP</td>
<td>DVA</td>
<td>30</td>
<td>30.67</td>
<td>7.512</td>
<td>0.676</td>
</tr>
<tr>
<td></td>
<td>Control</td>
<td>30</td>
<td>31.40</td>
<td>5.917</td>
<td></td>
</tr>
<tr>
<td>DBI</td>
<td>DVA</td>
<td>30</td>
<td>0.233</td>
<td>0.060</td>
<td>0.582</td>
</tr>
<tr>
<td></td>
<td>Control</td>
<td>30</td>
<td>0.225</td>
<td>0.051</td>
<td></td>
</tr>
</tbody>
</table>

Average value of the pressure (mm Hg) and index.

![Fig. 1. Clinical outcomes after DVA and antiaggregation therapy.](image1.png)

![Fig. 2. Hemodynamic changes after DVA.](image2.png)

![Fig. 3. Digital-brachial index improvement after DVA.](image3.png)

![Fig. 4. Decreasing trend of the lactate level after DVA.](image4.png)
Tables 4 and 5 show values registered at the measurement of the oxidative stress in 10 patients immediately before DVA and successively at 1, 3, 5 and 10 min postoperatively. Table 4 shows levels of estimated prooxidant parameters before DVA and at 1, 3, 5 and 10 min postoperatively. Using the nonparametric Friedman test ($p < 0.05$), it was observed that there was no significant difference in any prooxidant molecule at all times of measurement. Table 5 shows data of the enzymes of the antioxidative defense system before DVA and at 1, 3, 5 and 10 min after this procedure. Again, using the nonparametric Friedman test ($p < 0.05$), it was noted that there was no significant difference in any antioxidant enzymes between all times of measurement.

Surgical complications in a narrow sense started developing in 8 (26.6%) patients (Table 6). These complications meant infection of operative incision, graft thrombosis and bleeding. The early postoperative process was complicated with the development of pneumonia in 1 patient, with a cardiac decompensation in 2 patients and with the fatal outcome in 1 patient.

Discussion

Revascularization of critically ischemic limbs unsuitable for standard reconstructive surgical procedure is a surgical dilemma with a long prehistory. At the beginning of the 20th century, the only method of treatment of patients with gangrene of the extremities caused by atherosclerosis was high amputation. In 1896, the first clini-
cal report about arteriovenous reversal was published [16]. In 1912, Halstead and Vaughan [17] reviewed 42 cases and revealed only 1 patient with apparent perioperative pulsation of the foot veins who survived for 4 months without extension of the gangrene. The concept of reverse flow has been revised in 1951, when Szilagyi et al. [18] reported nine failures in 9 patients. They created a fistula between the common femoral artery and superficial femoral vein, reaffirming the view that the reverse flow is possible without destruction of venous valves. The reason for the failure in all patients can be explained with the small sample; they agree with Halstead and Vaughan that anoxia and the presence of fistulas are important for the formation of collateral flow. In 1975, Lengua [1] introduced the concept of a more distal anastomosis and achieved success in 3 of 8 patients. Originally, he did not perform valvulotomy preserving the dorsal venous arch and attributed success to the valves commonly absent from the superficial foot veins. Almost 20 years later, the same author published a series of 26 patients using terminolateral anastomosis between the great saphenous vein and the medial marginal vein, with mechanical valve destruction [19]. But this time after an average follow-up period of 1 year and 3 months, an average graft flow of 73% was reached. A similar success in 50% of the 6 patients was achieved by Sheil [20], by applying the technique of distal arterialization with destruction of the dorsal venous valve portion. After 18 months of follow-up, 20% of the fistula was left with a surprising functional limb preservation rate of 50%, indicating the viability of collateral circulation after graft occlusion. The best results of critically ischemic limb treatment using DVA were achieved by Pokrovsky et al. [3] in the largest published study so far. Metallic olives were used to break down the valves without direct visualization. They reported 85 cases with 80% limb salvage. In recent years the number of papers with optimistic results has been increasing [21–23]. In everyday surgical practice, most vascular surgeons in patients without the option for femoro-distal reconstructions consider the application of drugs (antiplatelet agents or prostanoids), spinal cord stimulation, and hyperbaric oxigenotherapy, but these are still limited and with modest results [24, 25].

What certainly justifies the performance of DVA as a limb salvage procedure are the acceptable morbidity and mortality rates which follow this procedure. In this study the mortality rate after DVA was 3.3%, while in conservatively treated patients it reaches 33.3% after an average 6-month follow-up period. According to our experience in a smaller group of patients, the outcome was significantly improved [5]. Despite the similar comorbidity rate in both groups, only 1 patient treated with DVA died of cardiac decompensation. The patient had patent graft and wound healing on the foot. Most of the patients in the control group, 8 out of 10, died of the septic complications of the foot gangrene.

Most of the patients in the DVA group with a patent graft, 22 out of 25, achieved pain relief at rest and improved walking capacity, therefore improving the functional status of the extremities. In 3 patients with a patent graft the gangrene process was not stopped, which required an urgent high amputation of the extremities. In 2 patients with control angiography a technical error was detected, which is illustrated in figures 5 and 6. After ligature of the overlooked tributaries the clinical status in the first patient did not get better, and in the second case an additional procedure to improve run-off was not possible. Finally, high leg amputation was necessary.

An ischemic ulcer healed successfully in 21 of 24 patients, all of whom had a patent graft. One of the indicators of the successful reverse revascularization of the leg was the increase in the digital systolic pressure from an average of 30.67 ± 7.5 mm Hg before the operation to
68.4 ± 16.2 mm Hg 7 days postoperatively (p <0.001). Also, in the same period of time the systolic digital-brachial pressure index increased from 0.233 ± 0.60 before the operation to 0.497 ± 0.147 postoperatively (p <0.001).

The surplus of the ischemic region lactate which is the result of the anaerobic metabolism is being gradually eliminated within the first 10 min after revascularization. There are two ways of eliminating the surplus of the lactate from the ischemic region. One is the conversion of the lactate into glucose, by the process of the gluconeogenesis in the liver, known as the Cori cycle. Nevertheless, this can hardly explain the rapid fall in the lactate level in venous blood of the revascularized extremity. A more rational way of explaining this rapid fall of the lactate is the oxygenation into pyruvates, which normally occurs during exertion in well-oxygenated muscle cells. Pyruvate created in this way can easily join the well-known Krebs cycle for producing energy in the form of ATP. Therefore, we would argue that DVA improves not only skin circulation – the most endangered tissue in critical limb ischemia – but also circulation of the muscles of the foot.

As we know, revascularization of the ischemic tissues may be followed by microvascular dysfunction, which leads to local and remote cellular and organ failure. Pathophysiology of ischemia-reperfusion injury concerns impaired endothelium-dependent, NO-mediated relaxation of smooth muscle to all receptor-dependent vasodilators in arterioles; enhanced capillary fluid filtration and the plugging of capillaries by stiffer, activated leukocytes with capillary malperfusion; leukocyte-endothelial cell adhesion, transendothelial leukocyte migration, platelet-leukocyte aggregation, albumin extravasation and enhanced oxidant production in postcapillary venules, and NO-superoxide imbalance in endothelial cells tipped in favor of superoxide [26–29]. Unfortunately, this scenario is not predictable but might be modified in the process termed acute or classical preconditioning and delayed preconditioning or ischemic tolerance [30, 31].

On the other hand, it is interesting that, considering pathophysiological mechanism of reperfusion injury, there are almost no data about the effect of DVA on oxidative status and cell damage. Analyzing the dynamics of oxidative stress of parameters (TBARS, O$_2$–, H$_2$O$_2$ and NO$_2$–) as well as the reply of the antioxidant system (SOD, CAT and GSH) after surgical revascularization of the foot, we can observe that DVA did not cause significant variations of all estimated parameters. Intervention did not improve antioxidant defense (which can be hardly expected because the tissue is not healthy), but more importantly, it did not increase oxidative stress. This result could be an acute response of local circulation, suggesting that there was no significant reperfusion injury, which can be expected after this kind of surgical procedure. At this stage of research, we still cannot determine why this occurred. One of the answers could be the development of ‘controlled reperfusion’, which in this case (retrograde perfusion through superficial venous system) provides adequate oxygenation with minimum damaging of the cells. However, it is certain that these pathophysiological events require further research. In the latest medical database search (PubMed) only few papers covering this problem could be found. It has been shown that the prostacyclin analogue (PGA$_2$) iloprost decreases the system level of oxidative stress parameters in patients with peripheral vascular disease [32]. Also, preoperative administration of L-alanyl-glutamine (250 ml) caused protection of the local tissue after surgical revascularization in patients with peripheral vascular disease versus those without this therapy [33].

**Conclusion**

When comparing our results with other studies, we can conclude that, in our case, DVA did not induce reperfusion injury but has achieved the therapeutically goal-adequate reoxygenation of ischemic tissue.

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**References**


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