Arteriogenesis and ischemia impair functional vasodilation in resistance arteries

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Introduction
Patients with peripheral arterial occlusive (PAOD) disease have impaired vasodilation. Identifying the mechanism of this impaired could lead to improved therapies. Co-morbidities common to PAOD, e.g., hypercholesterolemia, are known to impair vasodilation, but arterial occlusion alone also impairs vasodilation. Although impaired vasodilation in lower-leg arterioles is well described in animal models of PAOD, the impact of arterial occlusion on collateral and resistance arteries is unknown. Collateral resistance is the largest determinant of downstream flow and resistance arteries contribute up to half of the resistance to skeletal muscle blood flow, so identifying the impact of arterial occlusion on these vessels will likely provide the greatest potential therapeutic benefit. Therefore, the goal of this project is determine the impact of arteriogenesis and ischemia on collateral artery and resistance artery vasodilation, respectively. The long-term goal of this work is identify potential therapeutic targets for impaired vasodilation in patients with PAOD.

Experimental Model – Intravital Microscopy
We measured feed arteriomy diameter using side-stream Dark Field (SDF) intravital microscopy. SDF collects the reflected light from oblique 530nm pulses, which is the isobestic point for hemoglobin, so the vasculature appears dark and the parenchymal and stromal tissue appears light. Our intravital imaging station is shown in Figure 2. Example images of the muscular branch artery (at rest and following functional vasodilation) are shown in Figure 3.

Intravital Microscopy- Functional Vasodilation
We electrically stimulate muscle contraction in the gracilis muscles to assess the endogenous vasodilation pathways in the profunda femoris artery. The (-) stimulating electrode is placed in the superficial surface of the gracilis muscles and the (+) ground electrode is placed in the skin. The motor end plate is stimulated with 1mA pulses of 200-500μs duration at 8Hz for 90sec. Example resting and dilated profunda femoris arteries are shown in Figure 2.

Surgical Model I - Arteriogenesis
To determine the impact of elevated flow and arteriogenesis on collateral artery vasodilation, we utilized an experimental model that involved ligation of the femoral artery between the profunda femoris and popliteal branches (proximal ligation, Figure 3a) or just distal to the popliteal artery (distal ligation, Figure 3b). These surgeries lead to arteriogenesis in the profunda femoris and its development as the stem region of a collateral circuit.

Surgical Model II - Ischemia
To determine the impact of ischemia on feed artery vascular reactivity, we utilized an experimental model that involved resection of the femoral artery-vein pair from just upstream of the profunda femoris to the distal saphenous. Figure 6. This surgery presumably produces ischemia in the profunda femoris artery and the downstream gracilis muscle microcirculation.

Ischemia Delays Return to Baseline Diameter Following Intense Muscle Contraction
As seen in Figure 9, a more intense muscle contraction can cause a normal vasodilation in the ischemic profunda femoris artery. However, restoration of resting diameter seemed to be delayed in these studies. Therefore, we examined the time course of diameter before and after the electrical stimulation of the gracilis muscles with tungsten microelectrodes with 1mA pulses of 500μs duration at 8Hz for 90sec, Figure 10.

Ischemia Impairs Endothelial & Smooth Muscle-dependent Vasodilation
To test the hypothesis that impaired vasodilation following ischemia was due to impaired endothelial-dependent vasodilation, measured profunda femoris artery diameter at day-14 following resection, before and after electrical stimulation of the gracilis muscles with tungsten microelectrodes with 1mA pulses of 200μs duration at 8Hz for 90sec, Figures 7 & 8.

Ischemia does not Affect Functional Vasodilation during Intense Muscle Contraction
To further examine the impact of ischemia on profunda femoris artery functional vasodilation, we measured functional vasodilation at day-14 following resection, before and after electrical stimulation of the gracilis muscles with tungsten microelectrodes with 1mA pulses of 200μs duration at 8Hz for 90sec, Figure 9. The increased stimulus duration produces a more robust muscle contraction and presumably a greater stimulus for vasodilation.

Figure 11. Profunda femoris artery diameter at rest, following vasodilation to acetylcholine and sodium nitroprusside, and following reduction to noradrenaline. Top Panel- absolute diameter; Bottom Panel- percent change from baseline.

Summary
• Arteriogenesis transiently reduces functional vasodilation in collateral resistance arteries. The degree of reduction appears to be impacted by the arteriogenesis stimulus, Figures 4 & 5.
• Ischemia reduces functional vasodilation in resistance arteries, Figures 7 & 8. The extent of this reduction is impacted by the muscle contraction intensity and presumably vasodilation stimulus, Figure 9. This reduction appears to be due to reduced smooth muscle responses, Figure 11.
• Ischemia delays the restoration of resting diameter in resistance arteries following intense muscle contraction, Figure 10. This delay is not due to impaired responsiveness to noradrenaline, Figure 11.

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Figure 1. Intravital microscopy imaging station for functional vasodilation measurements.

Figure 2. Still images of the muscular branch artery at rest (left) and following 90 seconds of BHZ muscle contraction (right).

Figure 3. Medial aspect of the mouse hindlimb showing site of femoral artery ligation and primary collateral flow paths.

Figure 4. Left panel, profunda femoris artery diameter at rest and following muscle contraction in sham-operated animals and at day-7 and day-14 following distal femoral artery ligation. Right panel, percent increase in profunda femoris artery diameter following muscle contraction, same groups as left panel. * p < 0.05 vs sham vasodilation, # p <0.05 vs sham resting.

Figure 5. Left panel, profunda femoris artery diameter at rest following femoral artery resection in response to increasing stimulus frequency: 2, 4, or 8Hz and 1mA, 500μs, 90sec.

Figure 6. Medial aspect of the mouse hindlimb. The region of femoral artery-vein resection is indicated with a dashed line.

Figure 7. Profunda femoris artery diameter at rest (black bar) and immediately following the cessation of moderate muscle contraction (grey bar), stimulated at 1mA, 200μs, 8Hz, 2min. * p < 0.05 vs sham.

Figure 8. Functional vasodilation (% change from resting or 100%) of the ischemic profunda femoris artery with moderate intensity muscle contraction for sham (black) and day-14 ischemic (grey).

Figure 9. Muscular branch artery diameter at day-14 following femoral artery resection in response to increasing stimulus frequency: 2, 4, or 8Hz and 1mA, 500μs, 90sec.

Figure 10. Functional vasodilation recovery in the ischemic profunda femoris artery following high intensity contraction. Sham operated (black circles) and day-14 ischemic arteries (grey circles). * p < 0.05 vs sham.