1 2	<b>Title:</b> Beneath the Cuff: Often Overlooked and Under-Reported Blood Flow Restriction Device Characteristics and their Potential Impact on Practice
3	Short title: Cuff Design Variables Impacting BFR
4	Nicholas Rolnick <sup>1*</sup> . Kyle Kimbrell <sup>2</sup> . Victor de Oueiros <sup>3</sup>
-	Thenolas Romiek ', Ryte Rimoren', Vietor de Quenos
5	1 – The Human Performance Mechanic, CUNY Lehman College, New York, United States
6	2 - Owens Recovery Science, San Antonio, TX, United States; e-mail: Kyle@ors.io
7 8	3 - Graduate Program in Health Sciences, Federal University of Rio Grande do Norte (UFRN), Natal-RN, Brazil; e-mail: victor.sabino.121@ufrn.edu.br
9	*Nicholas Rolnick, The Human Performance Mechanic, 250 Bedford Park Blvd W, Department of Exercise
10	Science, CUNY Lehman, Bronx, NY, 10468, USA. E-mail: <u>nick@thehumanperformancemechanic.com</u>
11	Nicholas Rolnick – 0000-0003-0430-5015
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## 56 Abstract

Exercise with blood flow restriction (BFR) has been shown to be a useful technique to improve 57 58 muscle mass, muscle strength and a host of other physiological benefits in both healthy and injured populations using low intensities (20-30% 1-repetition maximum or < 50% VO<sub>2max</sub>). However, as 59 BFR is gaining popularity in both practice and research, there is a lack of awareness for potentially 60 important design characteristics and features associated with BFR cuff application that may impact 61 62 the acute and longitudinal responses to training as well as the safety profile of BFR exercise. While cuff width and cuff material have been somewhat addressed in the literature, other cuff design and 63 features have received less attention. This manuscript highlights additional cuff design and features 64 and hypothesizes on their potential to impact the response and safety profile of BFR. Features 65 including the presence of autoregulation during exercise, the type of bladder system used, the 66 shape of the cuff, the set pressure versus the interface pressure, the ratio of bladder to cuff width, 67 and the bladder length will be addressed as these variables have the potential to alter the responses 68 to BFR training. As more devices enter the marketplace for consumer purchase, investigations 69 specifically looking at their impact is warranted. We propose numerous avenues for future research 70 to help shape the practice of BFR that may ultimately enhance efficacy and safety using a variety 71 72 of BFR technologies.

Key words: BFR training, safety, autoregulation, bladder, kaatsu, occlusion training

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#### Introduction

Interest in blood flow restriction (BFR) training has led to its increased adoption in fitness and rehabilitation 84 settings (1-3) due to the numerous musculoskeletal, cardiovascular and performance benefits observed 85 following chronic use. However, varied methodologies (e.g., applied pressures, repetition schemes, and 86 87 cuffs and their various design feaures used to provide the BFR stimulus) reported in the literature increase 88 uncertainty interpreting the magnitude of the effects of BFR exercise due largely to use of non-personalized pressures (4) and minimal reporting of rationales for the applied pressure (5). Similar issues exist within 89 90 the exercise science and rehabilitation literature from insignificant reporting (6–9), although interpretation 91 of BFR interventions are exacerbated by the heterogeneity of BFR cuff prescription factors. A recent article 92 attempted to provide BFR application guidance based upon a consensus of researchers and clinicians (10), 93 recommending that personalized pressures be implemented across research and clinical practice because it 94 accounts for many of the variables (e.g., cuff width, blood pressure and limb circumference) that have been 95 associated with impacting arterial occlusion pressure (AOP), the minimum pressure needed to occlude both 96 arterial and venous return. Use of personalized pressures allows for a better determination of the relative 97 intensity of BFR because when standardized to a relative %AOP, blood flow is similar at rest despite 98 differences in cuff width (11) indicating a similar restrictive stimulus. While cuff width and material have 99 been reported in the literature, there are other cuff design and features that may impact the restrictive 100 stimulus and/or the cardiovascular and perceptual experiences of the exerciser, hindering extrapolation to practice. This is especially relevant considering the numerous reported devices used in administration of 101 102 BFR (3) that have varied capabilities to produce a personalized BFR stimulus. Without consideration and 103 reporting of these cuff-specific designs/features, it may lead to poorly designed research studies with limited 104 utility as practitioners are guiding their BFR prescriptions from the literature.

105 For example, a recent publication attempted to compare two commercially available BFR devices (B-Strong<sup>TM</sup> and Delfi Personalized Tourniquet device) against a heavy load strength training control group 106 107 on muscle excitation and training-related perceptual factors (rate of perceived exertion and muscle pain) in a fixed repetition design (12). However, the research design was poorly constructed due to a failure to 108 109 consider the differences in occlusive capabilities of the devices, resulting in a disparate comparison as one 110 condition was likely exercising significantly closer to failure than the other, augmenting the perceptual 111 experience of the exerciser in the Delfi Personalized Tourniquet device condition (13). The researchers did 112 not consider that the B-Strong<sup>™</sup> cuff is designed to be very difficult to achieve full occlusion (width – 5 cm; multi-chambered bladder system) (14) whereas the Delfi Personalized Tourniquet device is designed 113 114 to produce full limb occlusion (width - 11.5 cm; single-bladder system) (15). The authors did not report this important design characteristic of the B-Strong<sup>TM</sup> cuff (multi-chambered bladder) and as such, the 115

study's conclusions stated that it was more tolerable than the Delfi Personalized Tourniquet device while 116 117 providing similar electromyographic activation of the quadriceps. Nonetheless, oversights like this impact application of BFR because practitioners may assume the B-Strong<sup>™</sup> cuff is just as effective as the Delfi 118 119 Personalized Tourniquet device with better participant tolerability and similar myoelectric activity. As 120 accelerated muscle fatigue is likely the primary way BFR induces its beneficial effect on muscle (16), the 121 design of Bordessa et al. (2021) gives limited guidance to the potential efficacy of the B-Strong<sup>™</sup> cuff compared to the Delfi Personalized Tourniquet device as both exercised in a work-matched fashion, limiting 122 123 our understanding of the proximity to failure between conditions and related perceptual factors.

124 To date, there have been no effort-matched studies that can provide insight as to the degree of fatiguability (e.g., repetitions to failure) experienced between different pneumatic cuff types (e.g., nylon vs. elastic; 125 126 single vs. multi-chambered bladder systems) set at different pressure schemes (e.g., %AOP vs. arbitrary 127 pressure application based on limb circumference). In part, this oversight is due to the rapid growth of BFR 128 in the literature and a lack of awareness of potentially important cuff designs and features that may impact 129 the BFR prescription and participant experience. Thus, the purpose of this manuscript is to discuss and 130 hypothesize on the potential impact of these BFR cuff designs and features to allow for better translation 131 of research to practice and propose numerous avenues for future research.

#### 132

#### Why Personalizing Pressure Application is Likely the Best Approach to BFR Prescription

BFR exercise repetitions to failure (17), neuromuscular responses (e.g., torque and myoelectric activity) 133 134 (18), and the perceptual experience during exercise have been shown to be impacted by the pressure applied to the exercising limb and the type of BFR cuff used. For example, one study compared the acute muscular 135 and perceptual response to a bout of 4 sets of biceps curl exercise performed with either a 3-cm wide 136 137 Kaatsu® elastic cuff inflated to an arbitrary 160 mm Hg applied pressure or a 5-cm wide Hokanson nylon 138 cuff inflated to 40% AOP (19). Despite similar cellular swelling, electromyographic amplitudes and post-139 exercise torque production, the nylon cuff condition reported greater number of repetitions performed 140 during sets 2 and 3, lower rate of perceived exertion during set one and lower rate of perceived discomfort 141 during all sets compared to the elastic cuff condition. This is particularly important as both device selection 142 and perceptual responses have been identified as major barriers to successful BFR training implementation 143 (20). The discrepancy between conditions in perceptual responses and repetitions to failure may be explained by the higher relative applied pressure of the elastic cuff (~65  $\pm$  19% AOP) compared to the 144 nylon cuff (40% AOP). As higher applied pressures have been shown to reduce repetitions to failure, 145 146 increase cardiovascular responses (21), and elevate perceptual experiences compared to lesser applied 147 pressures (22), personalizing the pressure may reduce the excessive physiologic and perceptual responses 148 associated with higher pressures, increasing compliance in a long-term periodized program (20). Moreover,

at 30% 1RM, lower relative applied pressures have been shown to be equally as effective in producing 149 150 muscle hypertrophy as higher applied pressures (23) with decreased perceptual demands (24,25). In addition, when cuffs of different widths and materials are standardized to a %AOP, the physiologic and 151 152 perceptual responses are largely equivocal (26-28) indicating that much of the differences observed 153 following arbitrary pressure application protocols are due to varied degrees of relative personalized 154 pressures. Last, there is also a potential safety concern when using non-personalized pressures as different prescriptions of pressures do not provide similar levels of relative restriction and may predispose the 155 exerciser to full occlusion (23) or an ineffective pressure (17). Further, the assessment of AOP via handheld 156 doppler has been shown to be a surrogate for systolic blood pressure (29). As clinicians do not often 157 perform blood pressure screenings in outpatient settings (30), routinely assessing AOP potentially may 158 159 identify important hemodynamic status changes in patients/clients that could warrant referral; further increasing the contribution of BFR to the medical community on the whole. Given the aforementioned, it 160 161 is prudent for practitioners to implement BFR prescriptions using personalized pressures to account for 162 differences in anthropometry of the participants as well as some differences in cuff design. As lower applied 163 pressures may be equally as effective as higher pressures given a minimum threshold of load (31), cuff 164 characteristics that may reduce the amount of applied pressure needed to achieve AOP should gain more recognition and be specifically outlined in research designs moving forward. 165

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# The Impact of Lesser-Known Cuff Characteristics & Features on Personalized Pressure Application

Cuff width and cuff material's impact on AOP has been described elsewhere (28,32) and is more 168 commonly reported in the recent literature. Generally, wider cuffs reduce total AOP with greater 169 170 effectiveness of transmission to the limb and broader pressure plateaus (e.g., more distributed applied 171 pressure) compared to narrower cuffs. This aspect becomes relevant for prescribing BFR exercise because when the same pressure (% brachial systolic pressure) is applied, resistance exercise performed with 172 wider (non-elastic) cuffs can promote higher cardiovascular and perceptual responses (33). Lesser applied 173 pressures with wider cuffs may provide a safer restrictive stimulus to the neurovasculature (34,35) that 174 175 may enhance the safety profile of BFR exercise. Cuff material (e.g., nylon vs. elastic) appears to not 176 matter in altering the acute BFR exercise response as long as AOP can be determined (28), although only 177 one study exists in this area. Below we address additional cuff designs and features that may play a role in 178 further enhancing the safety profile of BFR exercise that are rarely (if ever) reported.

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180 *Autoregulation of Applied Pressure* 

Autoregulation refers to the capability of a device to maintain a consistent applied pressure on an exercising 181 182 limb (Figure 1). Conversely, manual pneumatic cuffs (e.g., non-autoregulated) do not adjust for the phase of muscular contraction and may increase the hemodynamic and perceptual responses to BFR exercise, 183 184 potentially impacting safety profile (36). Therefore, whether a BFR device is autoregulated may be an 185 important variable to report as it may impact perceptual experience and hemodynamic response to exercise. To date, no published study has directly compared BFR exercise performed with- and without 186 autoregulation using a cuff of similar width. A study currently in review from the lead author attempted to 187 answer that question. We showed that autoregulation reduced risk of minor adverse events (e.g., feeling 188 faint, numbness in the leg, excessive pain) ~7x compared to the same exercise performed without 189 autoregulation and was associated with lower delayed onset muscle soreness and reduced perceptual 190 191 experiences. The results of this study appear to provide preliminary support for the use of autoregulation to enhance the safety profile of BFR exercise, but more research is needed to make firmer conclusions. In 192 193 addition, commercially available BFR devices vary in their capacity to provide quick adjustments to applied 194 pressure during exercise, limiting conclusions about autoregulation to a particular device and not the feature 195 itself. Future BFR research should specifically report the presence or absence of autoregulation.

196

## \*\*INSERT FIGURE 1 HERE\*\*

## 197 Multi- vs. Single-Chambered Bladder System

198 A tourniquet – by definition – is designed to occlude arterial flow (37) and this concept forms the basis for the majority of BFR cuffs on the marketplace and in research because it allows a personalization of applied 199 200 pressure (10). The bladder is the portion of the tourniquet that encircles and applies circumferential pressure 201 to the limb to eliminate arterial inflow and venous return. Typically, most BFR cuffs are single-chambered 202 and apply circumferential pressure around the limb that increase cardiovascular and perceptual responses 203 at a given workload (38) and may reduce hypertrophy of the muscles underneath the cuff in longitudinal regimens (39,40). Nonetheless, despite these potential limitations, the use of a single-chambered bladder is 204 205 commonplace in the BFR literature and is the most implemented bladder type in the BFR training literature 206 as its able to determine AOP as long as it is wide enough (15).

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### \*\*INSERT FIGURE 2 HERE\*\*

Recently, commercially available devices have entered the marketplace that consist of numerous sequential bladders that according to the manufacturer are designed to reduce the potential for arterial occlusion and result in a non-uniform circumferential pressure during exercise (Figure 2) (41). As the multi-chambered bladder system is not designed to occlude, AOP is largely unfeasible and arbitrary pressures by the manufacturer have been recommended for use in practice (250 mm Hg for the upper body and 350 mm Hg for the lower body) (41) limiting generalizability. To the authors' knowledge, only one training study utilizing the B-Strong<sup>™</sup> cuff has been published and it does not help in answering the potential efficacy of the bladder type given that exercise was conducted to failure, no low-load comparison group was included, and no volume load was reported (41). This is important because low-load exercise with- and without BFR has been shown to improve muscle mass and strength to a similar degree (42,43). To date, no study has investigated longitudinal musculoskeletal outcomes and volume load when exercise is performed to failure between low-loads with and without different BFR bladder designs and heavy loads (> 70% 1-repetition

221 222 maximum).

223 Within the current BFR body of literature, there are three published studies comparing the acute responses 224 of a multi-chambered bladder system to a single-bladder system (12,14,44). All studies have similar methodological issues due to the multi-chambered cuff construction preventing researchers from making 225 226 pressures relative to that induced by the single chamber systems. Presumably, this results in a greater magnitude of AOP achieved by the single chambered systems, affecting acute measures and leading to 227 228 potentially faulty conclusions on safety risk and/or longitudinal outcomes. For example, one walking study 229 compared the acute perceptual and hemodynamic responses between the B-Strong<sup>TM</sup> cuff (5-cm cuff width) 230 and Hokanson rapid-inflator research device (18-cm cuff width) inflated to 300 mm Hg and 160 mm Hg, 231 respectively (44). The results appear to support the use of the B-Strong<sup>TM</sup> cuff for BFR aerobic exercise as 232 the Hokanson device promoted greater increases in heart rate, blood pressure, and double product during 233 exercise with elevated perceptual demands. Lactate levels were observed to be significantly greater in the Hokanson condition indicating that metabolic stress was greater for this condition than the B-Strong<sup>™</sup> 234 235 condition. This likely resulted in a larger stimulation of the afferents governing the muscle metaboreflex response, increasing cardiovascular and perceptual responses (45). Considering the width of the Hokanson 236 237 cuff (18cm), the magnitude of pressure used (160 mm Hg), and the demographics of the participants, the authors of this manuscript conjecture that most were exercising very near 100% AOP. For comparison, 238 Hughes et al. (36) used a narrower Hokanson cuff (13 cm v 18 cm width) and reported full arterial occlusion 239 in 18 subjects at 163.33 +/- 17.06 mm Hg (36). The Hughes et al. cohort likely had higher AOP values than 240 the Stray-Gunderson cohort given the subject pool was entirely male, had higher BMI values (23 +/-3 versus 241 242 28.94+/-3.28), and higher resting systolic blood pressure (116+/-11mm Hg versus 129+/-9 mm Hg), all factors that have been shown via direct or indirect evidence to influence AOP. Extrapolating safety and 243 244 potential longitudinal outcomes given the current dearth of longitudinal research on multi-chambered 245 bladder systems warrants caution given the likely minimal amount of pressure needed to induce beneficial 246 adaptations and the uncertainty regarding that pressure threshold without standardization to a personalized

pressure (17). Nonetheless, BFR appears to be safe across a variety of cuffs, pressure applications, and
protocols (46).

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Because the multi-chambered bladder system is very difficult to achieve arterial occlusion (12,14,41), 250 studies investigating the potential relevancy of a multi-chambered bladder system have accounted for the 251 252 lack of restriction capability by increasing volume performed (e.g., 3 sets of 30 repetitions compared to 253 commonly recommended 4 sets totaling 75 repetitions in single-bladder systems) (41). To reduce flaws in 254 comparisons between devices with different bladders, future studies should investigate the magnitude of post-exercise muscle fatigue (e.g., isometric/dynamic torque loss) following various application 255 parameters. Of most value to practice are acute studies that compare repetitions to failure between different 256 257 bladder types applied at recommended application settings (e.g., 250/350 mm Hg in multi-chambered bladder systems and 40-80% AOP in single-bladder systems) and longitudinal studies that track volume 258 259 load, relevant outcomes, and occurrence of adverse events in non-failure and failure repetition schemes. 260 These experimental designs will greatly increase practical relevancy, thus helping practitioners make 261 informed decisions regarding the device they choose to use with their clients and patients.

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#### 263 Contour vs. Straight Cuff

264 Cuff shape has been shown to impact the amount of applied pressure needed to determine AOP (Figure 3) (47). Contour cuff shapes are longer at the top and shorter at the bottom, creating a closer fit on the limb 265 266 due to differences in diameter. Contoured cuffs also can be manufactured with variable contour shape, a 267 design feature that allows for an even more secure fit to the limb as the device fastener apparatus can 268 account for small differences in extremity size and shape (48). Nonetheless, the difference in proximal to distal diameter of a contoured cuff reduces AOP slightly (~5.9 mm Hg) compared to a straight cuff (e.g., 269 270 cuff that is similar length on the top and the bottom) (49). Further, the occlusive stimulus may be different as straight cuffs are more likely to apply asymmetric pressures to the limb given the change in limb 271 272 circumference proximally to distally in the extremities (37). In at risk populations where pressures during BFR exercise may want to be minimized to theoretically enhance safety, the use of a contoured cuff may 273 be preferred to accommodate for the conical limb shape. To date, no study has directly compared the acute 274 and longitudinal responses to a BFR exercise regimen using cuffs of similar widths but varying in cuff 275 276 shape, so this area of research is largely unknown.

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#### **\*\*INSERT FIGURE 3 HERE\*\***

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### 281 Set Pressure Versus Pressure Applied to the Limb

282 The pressure that is set for BFR (i.e., "the set pressure") may not be the same pressure that is applied to the limb, known as the "interface pressure" (Figure 4) (36). Hughes et al. (2018) showed that when the Delfi 283 284 Personalized Tourniquet device (automatic autoregulated; cuff width = 11.5 cm; contoured cuff shape) was inflated to 40% and 80% AOP, the interface pressure was  $8 \pm 4$  mm Hg and  $9 \pm 4$  mm Hg lower than the 285 286 set pressure. Conversely, when the manual cuff (Occlusion Cuff, cuff width = 8 cm; straight cuff shape) was inflated to similar relative pressures, the interface pressure was  $20 \pm 10$  mm Hg and  $37 \pm 13$  mm Hg 287 lower than the set pressure. Thus, despite personalizing the pressure to %AOP, the amount of applied 288 pressure to the limb varied significantly between devices and may influence both acute (e.g., repetitions to 289 290 failure, cardiovascular and/or perceptual experience) and longitudinal (e.g., muscle hypertrophy, muscle 291 strength and/or vascular adaptations) outcomes. Preliminary results from Hughes et al. (2018) also indicated 292 that cardiovascular and perceptual experiences were heightened in the manual cuff compared to the 293 automatic autoregulated cuff when the same exercise was performed. This is likely due to a much greater 294 interface pressure (> 37 mm Hg) compared to the set pressure during the four exercise sets, indicating that 295 the manual cuff was applying a greater pressure to the limb during exercise than during resting conditions. 296 In contrast, the Delfi Personalized Tourniquet device maintained the set and interface pressure during 297 exercise that did not exceed +15 mm Hg in any of the four sets measured. However, despite setting AOP 298 to a similar percentage based on the cuff, the comparison wasn't direct as cuff widths varied between 299 devices, the Delfi Personalized Tourniquet device is autoregulated, and their cuff shapes varied. Insomuch 300 as what's currently known from the devices in the consumer market, the Delfi Personalized Tourniquet 301 device has been shown to apply a pressure within measurement error  $(\pm 15 \text{ mm Hg})$  to the underlying limb, 302 ensuring a stimulus that is like the set pressure during resting and exercise conditions. If possible, future 303 studies should integrate measurements for determining interface pressures, particularly when novel devices 304 are being investigated. Special attention should be paid to studies using lower (40-50% AOP) pressures in 305 their lower body interventions as this may impact the clinical relevance given lower pressures in this range have been shown to be ineffective at accelerating fatigue accumulation in BFR exercise (17). If a cuff used 306 307 in a lower pressure intervention was shown to be ineffective, researchers should determine if it was ineffective due to the parameters set (e.g., lower pressure) or inadequate cuff restrictive capabilities. 308

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#### \*\*INSERT FIGURE 4 HERE\*\*

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# 312 Presence/Absence of an Internal Stiffener

A stiffener is a feature of a tourniquet that directs the pressure from the bladder onto the limb and helps maintain the cuff's position when inflated (48). The presence of an internal stiffener may impact the degree of AOP and/or the exerciser's perceptual experiences during exercise as its presence increases the resistance to cuff deformation with muscular contraction. With respect to BFR exercise, no study has investigated the impact of an internal stiffener on cuffs with similar widths to determine its effect on acute- and longitudinal training outcomes. Future studies should determine its relevance with BFR exercise as more devices are

- being purchased and used in practice (3).
- 320

# 321 Bladder to Cuff Width Ratio

A cuff characteristic that is rarely reported is the bladder to cuff width ratio (Figure 5). This ratio is the percent of the cuff that the bladder makes up. The ratio is important because irrespective of cuff width, if the bladder width is significantly narrower, it may increase the amount of applied pressure needed to achieve AOP and predispose a similar stimulus as narrow cuffs. Future research studies should specify the cuff width and bladder width as cuffs with the same width may have different size bladders, impacting the transmissibility of the pressure to the exercising limb as well as related cardiovascular and perceptual responses.

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#### **\*\*INSERT FIGURE 5 HERE\*\***

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# 331 Bladder Length - Circumferential vs. Partial Circumferential

332 The last cuff characteristic that can impact BFR exercise is the length of the bladder (Figure 6). In traditional tourniquets, the bladder circumferentially envelopes the limb. In partial circumferential bladders, the 333 334 bladder does not extend the length of the cuff, leaving areas without pneumatic pressure application that instead relies on compression from the sleeve of the device. In the only published study investigating the 335 336 influence of bladder location on AOP during resting blood flow restriction, Spitz et al. (2020) showed that 337 positioning the bladder on the outside of the thigh required greater applied pressures than when the bladder was positioned on the inside of the thigh (50). As the femoral artery is located anteromedially (e.g., "inside" 338 position) and not anterolaterally (e.g., "outside" position), a lower pressure was required to occlude the 339 limb in the inside position. As most, but not all (e.g., Airbands/SAGA Fitness Cuffs) BFR devices on the 340 341 marketplace have circumferential bladders, little is known about the acute responses associated with 342 differences in bladder length. If bladder positioning impacts AOP in cuffs with partial circumferential 343 bladders, this may have relevancy for clinical populations where limited applied pressure may enhance acute safety and/or longitudinal training responses. Future studies should specify positioning of the bladder 344 when utilizing partial circumferential BFR cuffs and determine the potential relevancy of this cuff feature 345 346 to BFR exercise.

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#### \*\*INSERT FIGURE 6 HERE\*\*

348	
349	Conclusion
350	This manuscript attempted to contextualize the potential importance of infrequently reported BFR cuff
351	characteristics and features and hypothesize their potential impact on BFR training. As BFR continues to
352	expand into practice, researchers should be aware of not only the importance of AOP assessment and its
353	impact on BFR exercise responses, but of the ways that physiological responses may vary between cuffs
354	despite standardization to %AOP. Cuffs that are unable to be standardized to a %AOP (e.g., multi-
355	chambered bladder systems) may have clinical utility, but the current body of evidence on their efficacy is
356	lacking and should be a focal area of future research - particularly if similar beneficial results are obtained
357	with reductions in adverse events.
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364	NR wrote the initial draft of the manuscript. KK and VSQ provided critical review and helped edit the
365	manuscript for content and flow. All authors agreed to the final version of the manuscript and the statements
366	made in the article.
367	
368	Conflict of Interest
369	NR is the founder of The BFR PROS and teaches BFR training workshops to fitness and rehabilitation
370	practitioners using a variety of BFR training devices. KK is a clinical instructor for Owens Recovery
371	Science, a BFR education company that also distributes the Delfi Personalized Tourniquet Device. VSQ
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# FIGURE LEGEND

571	Figure 1. Autoregulation of Applied Pressures. Autoregulation is a design feature that accommodates for
572	the changes in limb circumference as a result of muscular contraction. In current available
573	devices, the BFR cuff is attached to a pneumatic air compressor via an air tubing that adjusts
574	according to the pressure sensed at the cuff-limb interface. The speed at which this adjustment
575	occurs varies across devices, making it a cuff-specific feature. Autoregulation may enhance the
576	acute safety of BFR exercise.
577	Figure 2. Multi-Chambered Versus Single-Chambered Bladder Cuff Design. As opposed to traditional
578	tourniquets whose function is to occlude arterial flow, multi-chambered bladders are composed of
579	sequential bladders that when inflated, leave regions where minimal compression occurs. This
580	cuff feature reduces the ability for the device to occlude arterial flow making it difficult to obtain
581	a personalized pressure. The inability to occlude has been hypothesized to enhance safety during
582	BFR exercise.
583	Figure 3. Differences in Limb Fit Between Contoured and Straight BFR Cuffs. Contour cuffs provide a
584	more secure fit due to the conical shape of the limb compared to a straight cuff. This may
585	enhance the safety profile of BFR exercise.
586	Figure 4. Set Pressure Versus Interface Pressure. The set pressure is the pressure that the pneumatic cuff
587	is inflated to by the clinician/exerciser/researcher whereas the interface pressure is the amount of
588	pressure actually applied to the limb from the cuff. Cuffs that can maintain a similar set and
589	interface pressures may enhance acute safety of BFR exercise.
590	Figure 5. Bladder to Cuff Width Ratio. The cuff width is the diameter of the cuff whereas the bladder
591	width is the diameter of the bladder. In some cuffs, the cuff width and the bladder width are not
592	identical. As the bladder is the portion of the BFR cuff applying pressure to the limb, if there is a
593	large difference between the cuff width and the bladder width, this may alter the AOP and the
594	applied pressure as a %AOP, influencing the acute safety profile of BFR exercise.

595	Figure 6. Partial Circumferential Versus Circumferential Bladder Length. In traditional tourniquets, the
596	bladder extends the length of the cuff (Right Image) whereas in some BFR cuffs, the bladder
597	extends partially not covering the entirety of the length of the cuff (Left and Center Illustrations).
598	Studies implementing BFR cuffs with partial circumference bladders should specify the position
599	of the bladder because its placement may impact acute responses to BFR exercise.
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AOP will be greater in the cuff with a smaller bladder !

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PARTIAL BLADDER

FULL CIRCUMFERENTIAL BLADDER