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Does cryoneurolysis result in persistent motor deficits? A controlled study using a rat peroneal nerve injury model

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Received 11 November 2019

Revised 13 January 2020

Accepted 14 January 2020

Published Online First

29 January 2020

ABSTRACT

Background Cryoneurolysis of peripheral nerves uses localised intense cold to induce a prolonged block over multiple weeks that has the promise of providing potent analgesia outlasting the duration of postoperative pain following surgery, as well as treat other acute and chronic pain states. However, it remains unclear whether persistent functional motor deficits remain following cryoneurolysis of mixed sensorimotor peripheral nerves, greatly limiting clinical application of this modality. To help inform future research, we used a rat peroneal nerve injury model to evaluate if cryoneurolysis results in persistent deficits in motor function.

Methods Male Lewis rats (n=30) had their common peroneal nerves exposed bilaterally at the proximal lateral margin of the knee and subsequently underwent cryoneurolysis on one limb and sham treatment on the contralateral limb. Outcomes were evaluated on days 3, 14, 30, 90 and 180. The primary end point was motor function, based on ankle dorsiflexion torque. In addition, sensory function was tested based on von Frey's filament sensitivity to the peroneal sensory distribution. A subset of animals was sacrificed following functional testing at each time point, and general tissue morphology, connective tissue deposition, and axon counts were evaluated.

Results Motor deficits in treated limbs were observed at 3 and 14 days but had resolved at time points beyond 1 month. Bilateral sensory deficits were also observed at 3 and 14 days, and also resolved within 1 month. Consistent with motor functional deficits, axon counts trended lower in treated nerves compared with contralateral controls at 3 days; however, axon counts were not significantly different at later time points.

Conclusions When applied to a mixed sensorimotor nerve, cryoneurolysis did not result in persistent motor deficits.

Cryoneurolysis is a non-opioid analgesic modality used to treat both acute and chronic pain involving the application of extremely low temperatures to a peripheral nerve to induce Wallerian degeneration and consequent abrogation of nerve conduction.² Axons regenerate along the undisturbed epineurium, perineurium, and endoneurium from the site of ablation towards their distal targets at about 1–2 mm/day,³ resulting in a nerve block with a variable duration, but typically measured in multiple weeks.⁴ While cryoanalgesia has been used for decades to treat primarily chronic pain,^{4–8} the advent of percutaneous cannulas and ultrasound guidance dramatically increases the potential applications for its use in treating acute pain as well.^{9–11}

To date, most published research involves the functional and histological recovery of sensory nerves^{12–16}; however, mixed motor-sensory nerves may also be targeted.^{4 10 17 18} Because cryoneurolysis affects both sensory and motor neurons, concern exists regarding residual and persistent motor weakness due to either incomplete regeneration or motor unit clustering. Given that interest in cryoneurolysis is increasing due to an increased awareness of the pervasiveness of inadequately-controlled postoperative pain,¹⁹ current public health efforts to decrease opioid use,²⁰ and recognition that chronic, persistent postoperative pain is associated with inadequately-controlled perioperative pain,²¹ it is imperative that any long-term cryo-induced motor deficits be identified.

In this controlled study, we used a sensitive and validated rat common peroneal (fibular) nerve injury model to evaluate if cryoneurolysis results in persistent motor function deficits, based on recovery of ankle dorsiflexion torque.

METHODS

Study intervention

Thirty 300 g male Lewis rats underwent survival surgery under isoflurane anaesthesia with analgesia provided with preoperative subcutaneous injection of 0.3 mg sustained release buprenorphine (Buprenorphine SR-LAB, ZooPharm, Windsor, Colorado, USA). Common peroneal (fibular) nerves were exposed bilaterally at the proximal lateral margin of the knee joint using a technique previously described.^{22 23} The fascial sheath overlying the nerve was left intact. On the left limb, cryoneurolysis was performed with a console based

INTRODUCTION

Pain following surgery is frequently treated with local anesthetic-based peripheral nerve blocks to improve analgesia, decrease/negate opioid reliance, and decrease the incidence/severity of persistent postsurgical pain. However, the duration of surgical pain is often far longer than the duration of even the longest-acting single-injection peripheral nerve block; and, perineural local anesthetic infusion is limited to less than 1 week in most cases.¹

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To cite: Shah SB, Bremner S, Esparza M, et al. *Reg Anesth Pain Med* 2020;**45**:287–292.

clinical cryoneurolysis machine (PainBlocker, Epimed International, Farmers Branch, Texas, USA). An 18 G trocar with a tapering tip was positioned immediately adjacent to the target nerve and three 2 min cryoneurolysis cycles separated by 1 min thaw periods were applied using nitrous oxide. This protocol is a typical clinical protocol,³ and given the size and exposure of the rat peroneal nerve, may be viewed as aggressive. On the right limb, sham cryoneurolysis was performed, with exposure, tip contact, and time of contact as above, but no freezing.

Multiple criteria were considered to retain consistency within our cryoneurolysis protocol. Frosting of the cannula tip as well as formation of an ice ball encompassing the nerve was confirmed for each cycle. Repeatability in freezing technique was confirmed based on: (i) consistent exposure of the peroneal nerve; (ii) device indicator confirming sufficiently cooled cannula tip; (iii) frosting of cannula; (iv) visual verification of contact of cannula tip directly on fascial sheath overlying peroneal nerve; (v) ice ball formation on tissue surrounding nerve; (vi) fasciculations on at least one of the freeze cycles. Increased local blood flow was typical during a freeze cycle. Cannulas were thawed completely before detachment from tissue to avoid imposing mechanical force on the nerve by cannula manipulation.

On the contralateral limb, the sham (control) treatment was administered by positioning the cryoablation cannula immediately adjacent to the target nerve for the same three 2 min on, 1 min off pattern as before, but without freezing (ie, no nitrous oxide). Overlying muscles were repaired by closing the fascia with 4–0 polyglycolic acid (PGA) sutures, and wound clips were used to close the skin incision.

Outcome measures

End points were assessed at 3, 14, 30, 90, and 180 days following treatment, a time frame over 3x longer than that during which a small population of persistent motor deficits were observed.²³ Motor function was assessed bilaterally *via* isometric ankle dorsiflexion torque, using a previously-described and validated method.²³ Briefly, following anaesthesia induction (2% isoflurane, 2.0 L/min), dorsiflexors were activated via the common fibular nerve using a nerve cuff (15 V at 100 Hz, 0.3 ms pulse duration and 650 ms train duration), while torque was measured using a custom-designed dynamometer.²² The dynamometer was created from a dual-mode servo motor (Aurora Model 305B, Aurora Scientific, Aurora, Ontario, Canada) which has a force resolution of 1 mN, a linearity of 99.8% and a step response time under 2 ms. Peak torque was defined as the absolute peak isometric torque elicited after the onset of stimulation. Sensory testing was performed bilaterally with von Frey filaments applied to the dorsal surface of the foot (peroneal nerve sensory distribution).²⁴ Rats were acclimated with one round of practice testing preoperatively. Rats were required to plant their entire foot (heel to toe) prior to testing, and paw withdrawal pattern was assessed using the up–down method.²⁵ Data are represented as the difference in post-treatment sensitivity (ie, force eliciting withdrawal) minus baseline sensitivity, with more negative values indicating hypersensitivity.

Histological outcomes

A subset of animals was sacrificed under anaesthesia following functional testing at each time point *via* intracardiac injection of pentobarbital sodium and phenytoin sodium (150 mg/kg). In addition, three animals were euthanised immediately after freezing, for histological analysis of acute structural changes resulting from cryoneurolysis. For each animal, the common

peroneal nerve was harvested bilaterally, and snap frozen using liquid nitrogen cooled isopentane.²⁶ Transverse sections of the nerve underlying the distal edge of the cryoablation site and axial sections of tibialis anterior (TA) muscle ~5 mm distal to its insertion at the knee were taken per previously published methods, and slides were stored at –80°C until use.^{23,26} Sections were stained with H&E, Masson's trichrome (Polysciences, Warrington, Pennsylvania, USA), Wheat Germ Agglutinin (WGA) conjugated to AlexaFLuor 594, DAPI, and/or β 3-tubulin+laminin. The neuron-specific β 3-tubulin was used as an axonal marker for axon counting, laminin provided an outline for delineating distinct nerve fibers, and WGA outlined muscle fibers. Sections were permeabilised with 0.2% Triton-X, blocked with blocking buffer (10% goat serum, 3% bovine serum albumin (BSA) in phosphate buffered saline (PBS)) then incubated with primary antibodies mouse anti- β 3-tubulin (Sigma-Aldrich; 1:500 dilution) and rabbit anti-laminin (Sigma-Aldrich; 1:1000 dilution) for 1 hour. Slides were then incubated with AlexaFluor 488 goat-anti-mouse and AlexaFluor 594 goat-anti-rabbit secondary antibodies, then coverslipped with Vectashield Hardset with 4',6-diamidino-2-phenylindole (DAPI; Vector Labs, Burlingame, California, USA).

Fluorescent images were acquired using a Leica DM6000 microscope (Leica Microsystems, Buffalo Grove, Illinois, USA) equipped with a Leica DFC365 FX camera (Leica Microsystems) using a 10X or a 63X oil immersion objective, and bright-field images were acquired at 20X using a Leica SCN400 slide scanner (Leica Microsystems). Axon counts were estimated from two sections, based on manual counting of β 3-tubulin-positive puncta in selected 63X images covering at least 35% of the nerve cross-sectional area, and subsequent scaling based on cumulative fascicular area measured from 10X images.²⁶ This stereological sampling approach procedure provided sufficient coverage to provide confidence in axon counts, while also enabling the inclusion of sections with minor tears or regions of inconsistent staining. All histological analysis was blinded to the data analyst.

Statistics

The primary null hypothesis was that a clinically-relevant cryoneurolysis protocol does not induce persistent motor functional deficits. This hypothesis was tested based on measurement of ankle dorsiflexion torque as a primary end point. We achieved statistical power of 0.86 for $n=6$ /group, for a moderate effect size of 0.5 and $\alpha=0.05$. Secondary null hypotheses were that a clinically-relevant cryoneurolysis protocol did not induce persistent mechanical sensory functional deficits or structural changes, as measured by von Frey's hair testing and comparison of axon counts, respectively. Parametric statistical testing was performed for all comparisons. Normality was confirmed using the Shapiro-Wilk normality test. Mean motor, sensory and structural outcomes were compared using mixed design 2-way analysis of variance (ANOVA), which examined main and interaction effects of time (unpaired factor) and intervention (paired factor). Tukey's Honestly Significant Difference (HSD) post-hoc test was used to compare individual means and accounted for multiple comparisons. Prism 8 (GraphPad, La Jolla, California, USA) was used for all statistical testing.

RESULTS

As determined by our six criteria, consistent freezing was confirmed for each experiment. General histological assessment using H&E showed clear changes to tissue surrounding the epineurium immediately following cryoneurolysis, including

Figure 1 Cryoneurolysis results in immediate tissue changes exterior to, but not within, the ablated nerve (H&E label). (A) Control nerve in cross-section. Note scale of comparatively thin epineurium. Some tissue has folded over to the left side of the image. (B and C) Cross-sections of two different nerves in region of cryo-ablation. Note clear capsule formation around epineurium (asterisk *), and connectivity to underlying muscle tissue following ablation protocol. Bars 100 μ m. (D) Longitudinal section of nerve in region of cryo-ablation. Note seemingly normal nerve tissue within the epineurium, but apparent freeze damage and connectivity to underlying muscle outside of the nerve (asterisk *). The 'crack' in the nerve (arrow) is likely to be an artifact of sectioning. Bar 100 μ m.

thickening of the epineurial wall and adhesion to surrounding musculature (figure 1A–D). However, the interior of the nerve appeared grossly intact. Consistent with these observations, trichrome staining in cryoneurolysed—but not sham—nerves indicated connective tissue deposition around the nerve, with structurally intact interiors (figure 2). These changes persisted throughout the evaluation period (figure 2D). Axonal localization within nerves also appeared qualitatively normal, in all groups (figure 3A–D). At 3 days after surgery, but at no other time point, decreased axon counts (indicative of Wallerian degeneration) were observed in cryoneurolysed nerves compared with sham (figure 3E). No pain-associated behavioral changes were observed after surgery at any time point.

Motor function

To test the primary hypothesis that cryoneurolysis did not result in persistent motor deficits, ankle dorsiflexion torque was evaluated. Two-way ANOVA showed main effects of both time ($p < 0.001$) and treatment ($p < 0.05$) on peak dorsiflexion torque, but no interaction effect. Consistent with normal growth, motor function gradually improved in this group over the 180-day evaluation period. Sham cryoneurolysis did not result in motor impairment at any time point, validating that surgical approach did not affect the control group (figure 4A). *Post hoc* testing of individual means revealed that peak torque was not significantly different between cryoneurolysis and sham groups at any time point (figure 4A). Recovery of torque to statistically insignificant magnitudes was observed at 30 days ($p = 0.88$); and, in support of the primary hypothesis, torque was indistinguishable between treatment and control limbs at 3 and 6 months ($p > 0.99$,

Figure 2 Cryoneurolysis results in persistent tissue changes exterior to, but not within, the ablated nerve (trichrome label). (A–D) Sample trichrome labeling for sham (A,C) and cryoneurolysed (B,D) nerve at 3 days (A–B) and 180 days (C–D). Note more integration of frozen nerves with surrounding nerve bed, in contrast to sharper epineurial borders in sham-treated nerves. White arrows in (A) indicate muscle connective tissue, which is distinct from epineurial tissue. Bars=100 μ m. Note that fasciculation patterns appear different in each section; this reflects anatomical variation in internal branching from nerve to nerve, despite consistency in section location.

figure 4A). Consistent with these observations, no differences in fiber morphology, spacing between muscle fibers, or centralized nuclei were observed between sham and contralateral TA muscle 6 months after injury (figure 4B–E).

Sensory function

Two-way ANOVA revealed main effects of time ($p < 0.001$), but not treatment ($p = 0.45$) on mechanical sensitivity in the dorsal paw (peroneal sensory distribution). Post-hoc testing revealed no statistically significant difference between treated and control limbs at any time point (figure 5), with slight (1–2 filament) bilateral hypersensitivity at Day 3, possibly associated with surgical trauma. All hypersensitivity resolved by 1 month, and normal sensitivity was maintained at 3 and 6 months (figure 5).

DISCUSSION

This study provides quantitative evidence in an animal model that cryoneurolysis does not result in persistent functional deficits when applied to a mixed sensorimotor nerve of a rat. The expected motor deficits existing within the first two post-treatment weeks (days 3 and 14) resolved by 1 month, mirroring the histological evidence of Wallerian degeneration followed by regeneration. While we cannot extrapolate the timing of motor recovery in laboratory animals to a clinical setting, these results are reassuring in that cryoneurolysis, a modality that is approved by the United States Food and Drug Administration and used clinically for multiple decades—does not appear to induce persistent deficits.

Our data should be placed in the context of several previous studies that examined effects of cryoneurolysis on nerve structure and neuromuscular function. Whittaker published a comprehensive assessment of nerve morphology and ultrastructure after freezing the ventral epithelium of the golden hamster tongue.^{27 28} Significant disruption of myelination as well as

Figure 3 Nerve morphology and axon counts are not affected by cryoneurolysis. (A–D) Sample B3-tubulin (axons, green), laminin (fiber borders, red), and DAPI (nuclei, blue) labeling for sham (A,C) and cryoneurolysed (B,D) nerve at 3 days (A–B) and 180 days (C–D). Bars=100 μ m. (E) Quantification of axon counts (mean $\pm$ SEM) shows no effect of time or treatment, and NO statistically significant difference in axon number (sham vs cryoneurolysis) at any time point (mixed-design two-way ANOVA and Tukey's HSD post-hoc testing). A trend towards fewer axons at 3 days, but no other time point, is observed, consistent with motor recovery. Bars=100 μ m.

signatures of axon degeneration were observed within an hour of freezing, with substantial myelin disruption and absorption as well as axolemmal disappearance. By day 14, early signs of regeneration, including axonal regrowth within early myelinating sheaths, were apparent (although functional testing was not performed). While we did not perform electron microscopy on ablated nerves, we observed structural damage to the nerve at 3 and 14-day time points, although not as comprehensive as in the lingual nerve model. Our timeline of functional recovery by 1 month also suggested structural regeneration at an earlier time point; however, differences in species and scale preclude a direct comparison.

Deleo, Wagner and colleagues published a subsequent series of studies on outcomes following cryoneurolysis of the rat sciatic nerve with the intent of validating a new model of pain.^{29–33} In these studies—as for our investigation—adverse sensory outcomes were observed in both the cryo-treated and control limbs.³⁴ However, sensory deficits persisted beyond 28 days suggesting more severe damage, in contrast to our observations of sensory recovery by 1 month. The authors speculated that the persistence of bilateral deficits was indicative of a central nervous system contribution to neuropathy.³⁴ This persistence may also have a methodological basis: in addition to freeze damage, the nerve was decompressed and raised from its bed during cryoneurolysis which may have created a non-physiological heat (cold)-transfer and mechanical environment more conducive to damage (Wagner, personal communication 2019).

A third set of studies by Hsu and Stevenson directly tested motor functional recovery.^{35–36} In this case, a tibial nerve injury

Figure 4 Motor function and muscle structure are not persistently affected by cryoneurolysis. (A) Peak isometric torque revealed ipsilateral motor deficits on the ipsilateral (ablated) side at early time points, which resolved by 30 days (main effects of time ($p<0.001$) and treatment ($p<0.05$) based on mixed-design two-way ANOVA. No significant differences between individual means (sham vs cryoneurolysis) at any time point based on Tukey's HSD post-hoc testing). Values are provided as mean $\pm$ SEM. (B) Control and (C) cryoneurolysed TA muscle at 180 days show no differences in gross morphology (H&E labeling). (D) Control and (D) cryoneurolysed TA muscle at 180 days show no differences in muscle fiber morphology, no widening of intra-fiber spacing, and NO centralised nuclei (WGA and DAPI labeling). Bars=100 μ m. TA, tibialis anterior; WGA, Wheat Germ Agglutinin.

was created using a single 1 min long cryoneurolysis application (which in one study was repeated three times at 2-month intervals). Motor function was evaluated using toe spreading

Figure 5 Mechanical sensitivity is not persistently affected by cryoneurolysis. von Frey's testing revealed slight (difference of 1–2 filaments) bilateral increases in sensitivity in the peroneal sensory distribution (dorsal aspect of the paw) at early time points, which resolved by 30 days. (main effects of time ($p<0.001$) but not treatment ($p=0.45$) based on mixed-design two-way ANOVA. No significant differences between individual means (sham vs cryoneurolysis) at any time point based on Tukey's HSD post-hoc testing). Values are provided as mean $\pm$ SEM

and a semi-quantitative walking scale, and reversibility of motor deficits was observed within 50 days. While these results were promising, there were several important limitations to this work that precluded a definitive conclusion. Importantly, a relatively benign cryoablation protocol was used, which may not represent a clinically-typical and more invasive multi-cycle cryoneurolysis protocol.⁴ Most critically, motor functional assessments were semi-quantitative and low in sensitivity, and minimal quantitative assessment of nerve morphology was performed. These confounding factors were compounded by the decision to treat the tibial nerve: while this nerve indeed plays an important role in ankle plantar-flexion and toe abduction, it does not innervate toe extensors and ankle dorsi-flexors which are essential for proper stepping during walking and the uncurling of toes. Thus, their reported semi-quantitative motor score which incorporated qualitative assessments of both gait and toe curling, may not have accurately or adequately described the impacts of their particular injury model. Lastly, a small (though not statistically significant) cohort of animals displayed continued motor deficits at the latest time point.^{35 36} These observations suggested that higher-resolution measurements may be needed to detect more subtle—but persistent—motor deficits. It was for this reason that we undertook the current study using a far more sensitive model of functional motor deficits. Our more sensitive and quantitative functional motor outcomes were qualitatively consistent with these prior studies in that motor deficits were indeed reversible.

Our peroneal nerve injury model and measurement of dorsi-flexion joint torque, which have been validated in multiple studies perturbing various aspects of neuromuscular function,^{22 23 37 38} offer several advantages to prior studies, and provide additional confidence in interpretation of our results. Peroneal nerve injury stands in contrast to sciatic nerve injury, in which both the tibial and peroneal branches are affected. Without a more sensitive outcome specific to each branch, it is difficult to confidently interpret outcomes. For example, following sciatic nerve injury, poor functional recovery may be a result of incomplete or imbalanced regeneration into the two branches. In our model, we injured only one branch, and the corresponding sensory and motor outcomes tested were directly related to the peroneal distribution—that is, joint dorsiflexion torque and sensitivity of the dorsal surface of the foot. Thus, there is no ambiguity, compared with observational assessments of motor function or behavioral outcomes such as Rotarod or grip testing. In addition, we applied a protocol of multiple freeze cycles separated by shorter thaw periods which better-represents a typical clinical application protocol that is intended to provide analgesia and not induce pain. Additionally, the nerve was left attached and unperturbed in its underlying tissue (ie, not decompressed), simulating the clinical reality of percutaneous cryoneurolysis. Thus, we are confident that the reversibility of sensory and motor deficits is not artifactual and is specific to the nature of the nerve injury.

The transient and minor nature (<2 von Frey hair's difference) of bilateral sensory hypersensitivity on both the treated and untreated limb may simply reflect discomfort due to bilateral surgical trauma, which may also contribute to the higher variability in motor and sensory function at the 3-day time point (figure 5). However, bilateral sensory hypersensitivity has also been noted in a variety of unilateral nerve injury models, including nerve freezing.^{34 39 40} A number of hypotheses have been posed to explain these phenomena, including systemic factors⁴¹ and central processing of pain.^{41–43} The current study design could not distinguish among these possibilities; as the primary outcome of this study was to evaluate motor function, an important extension of this work is the need for deeper evaluation of sensory outcomes.

Regarding motor deficits, we observed variable muscle weakness ranging from normal to complete loss of function in the active treatment limbs. This may reflect variable and/or incomplete nerve injury and is consistent with both histological and clinical observations. Structural changes to the nerve bed were visible immediately after cryoneurolysis and persisted throughout the 6 month recovery period. However, the degree and rapidity with which surrounding tissue thickened after the initial cycle of cryoneurolysis may impact the delivery of consistent freeze damage within the nerve in subsequent freeze cycles.⁴ Consistent with functional loss at 3 and 14 days, axon counts generally trended downward at 3 days and recovered within 1 month; however, the variability in axonal reduction was also substantial. A vascular contribution to functional deficits has also been proposed,⁴ although such a mechanism is not intuitively apparent from our study. However, our observation of mesoneurial deposition of connective tissue (figure 3) may contribute to entrapment at longer time points, which may result in neurovascular compromise.^{44 45}

Limitations

Due to potentially confounding effects of surgical exposure required for high-sensitivity functional testing, we were unable to follow the functional recovery of a single rat longitudinally. Thus, we relied on comparisons to contralateral sham controls, in which there may be some inherent baseline differences in structure and function.^{46 47} We powered our study to evaluate the primary outcome of motor function and, in this effort, we were successful. However, this sample size was likely underpowered for more subtle structural outcomes, including distinguishing between sensory and motor-specific histological outcomes. We also acknowledge that cryoneurolysis was not superposed on a model of pain, and so we could not evaluate its impact on pain relief. Finally, while modern clinical deployment of cryoneurolysis is frequently ultrasound guided and percutaneous, we acknowledge that the peroneal nerve was exposed during cryoneurolysis, an approach more commonly performed by surgeons in earlier studies.⁵ Percutaneous insertion of currently-available cryo-delivery cannulas would have been more challenging at the small scale of rat limbs; our technique enabled more consistent placement of the cannula on its target, providing us more confidence in our subsequent results. This approach may also be viewed as more conservative, as no motor deficits were observed *in spite of* the more invasive approach on a very small nerve target.

CONCLUSIONS

We combined a typical clinical cryoneurolysis protocol and a sensitive, validated rat peroneal nerve injury model to assess the effect duration of cryoneurolysis on motor function. When applied to a mixed sensorimotor nerve, cryoneurolysis did not result in persistent functional deficits.

Summary statement

When applied to a mixed sensory-motor nerve in Lewis rats, cryoneurolysis did not result in persistent structural or functional motor deficits.

Acknowledgements The authors appreciate the invaluable assistance of Rochelle Wagner, PhD (Senior Medical Science Liaison, TerSera Therapeutics, Lake Forest, Illinois) for her help in designing the protocol and manuscript review.

Contributors SS: Planning, design, conduct, acquisition of data, data analysis and interpretation, and manuscript preparation. SB, ME, SD, EO, CH: Conduct, acquisition of data and manuscript revision. BMI: Conception, planning, design, data interpretation and manuscript preparation. RAG: Planning, design, conduct, data

interpretation and manuscript revision. SW: Conception, planning, design, conduct, data interpretation and manuscript revision.

Funding Funding and equipment for this project provided by Epimed International (Farmers Branch, Texas).

Competing interests SS: The following companies have provided material and/or funding for his research directly to SS's institution: Epimed International (Farmers Branch, Texas). BMI: The following companies have provided material and/or funding for his research directly to BMI's institution: Epimed International (Farmers Branch, Texas); Myoscience (Fremont, California); SPR Therapeutics (Cleveland, Ohio); Infutronics (Natick, Massachusetts); Ferrosan Medical (Szczecin, Poland); and Heron Pharmaceuticals (San Diego, California). RAG: The following companies have provided material and/or funding for his research directly to RAG's institution: Epimed International (Farmers Branch, Texas); Myoscience (Fremont, California); SPR Therapeutics (Cleveland, Ohio); Infutronics (Natick, Massachusetts); and Ferrosan Medical (Szczecin, Poland). SW: The following companies have provided material and/or funding for his research directly to SW's institution: Epimed International (Farmers Branch, Texas); DeNova Inc (Chicago, IL).

Patient consent for publication Not required.

Ethics approval All animal work was prospectively approved by the University of California, San Diego Institutional Animal Care and Use Committee (La Jolla, California).

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request from the authors. Please contact SS, PhD (sbshah@ucsd.edu).

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