University Library

MINERVA ACCESS

A gateway to Melbourne's research publications

Minerva Access is the Institutional Repository of The University of Melbourne

Author/s:
Pelot, NA;Vaseghi, M;Reznikov, L;Osborne, PB;Conde, SV

Title:
Editorial: Multiscale anatomy and biophysics of the autonomic nervous system: implications for neuromodulation

Date:
2023-01-01

Citation:
Pelot, N. A., Vaseghi, M., Reznikov, L., Osborne, P. B. & Conde, S. V. (2023). Editorial: Multiscale anatomy and biophysics of the autonomic nervous system: implications for neuromodulation. Frontiers in Neuroscience, 17, https://doi.org/10.3389/fnins.2023.1289177.

Persistent Link:
https://hdl.handle.net/11343/345490

License:
CC BY

frontiers | Frontiers in Neuroscience

TYPE Editorial
PUBLISHED 01 November 2023
DOI 10.3389/fnins.2023.1289177

Check for updates

OPEN ACCESS

EDITED AND REVIEWED BY
Jasenka Zubcevic,
University of Toledo, United States

*CORRESPONDENCE
Nicole A. Pelot
✉ nikki.pelot@duke.edu

RECEIVED 05 September 2023
ACCEPTED 10 October 2023
PUBLISHED 01 November 2023

CITATION
Pelot NA, Vaseghi M, Reznikov L, Osborne PB and Conde SV (2023) Editorial: Multiscale anatomy and biophysics of the autonomic nervous system: implications for neuromodulation. *Front. Neurosci.* 17:1289177. doi: 10.3389/fnins.2023.1289177

COPYRIGHT
© 2023 Pelot, Vaseghi, Reznikov, Osborne and Conde. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

Editorial: Multiscale anatomy and biophysics of the autonomic nervous system: implications for neuromodulation

Nicole A. Pelot[1]*, Marmar Vaseghi[2], Leah Reznikov[3], Peregrine B. Osborne[4] and Silvia V. Conde[5]

[1]Department of Biomedical Engineering, Duke University, Durham, NC, United States, [2]UCLA Cardiac Arrhythmia Center, Division of Cardiology, Department of Medicine, University of California, Los Angeles, Los Angeles, CA, United States, [3]Department of Physiological Sciences, University of Florida, Gainesville, FL, United States, [4]Department of Anatomy and Physiology, University of Melbourne, Melbourne, VIC, Australia, [5]NOVA Medical School, Faculdade de Ciências Médicas, Universidade NOVA de Lisboa, iNOVA4Health, Lisboa, Portugal

KEYWORDS

autonomic nervous system, neuroanatomy, neuromodulation, neural stimulation, imaging, image analysis, cardiovascular disease

Editorial on the Research Topic
Multiscale anatomy and biophysics of the autonomic nervous system: implications for neuromodulation

Introduction

Using peripheral neuromodulation to target the autonomic nervous system is a rapidly growing therapeutic approach in which electrical, ultrasonic, or optical signals are delivered invasively or non-invasively. These signals alter nerve activity to treat multiple visceral system disorders, including cardiovascular, respiratory, gastrointestinal, and urogenital pathologies. Given that the neuromodulation targets include multiple fiber types that innervate multiple organ systems, there is a critical need to better understand the relevant neuroanatomy and physiology of this network to develop more effective clinical therapies.

This Research Topic introduces studies that advance our knowledge of neuroanatomy, including functional organization of the pig vagus nerve, innervation of the kidneys in mice, and a review of cardiac vagal afferents. Additional studies provide analytical tools to quantify the structure of autonomic nerves, including efficient segmentation of microCT of the human vagus nerve and statistical descriptions of the spatial arrangement of axons. Finally, neuromodulatory approaches for treatment of ischemia are presented.

Functional organization of the pig vagus nerve

Implanted vagus nerve stimulation (VNS) is most commonly used to treat epilepsy and is under investigation for many other conditions. Thompson et al. used complementary approaches to reveal the functional organization of the left cervical vagus nerve (VN) in anesthetized pigs: electrical impedance tomography (EIT), electrical stimulation using small

contact pairs, and microCT of post-mortem nerve samples. Microcomputed tomography (microCT) uses X-rays for 3D, non-destructive, high resolution (e.g., 150 μm voxels) imaging of stained tissues. Based on the microCT, at the mid-cervical level of the VN, fibers from pulmonary and recurrent laryngeal (RL) branches were mixed across many fascicles (i.e., bundles of axons), although they spanned mostly separate regions, while the cardiac fascicles remained distinct; the rotational location of the three functional groupings was approximately consistent across animals. In the EIT study, they delivered current through different contact pairs of a multi-contact cuff electrode; the voltage recorded on all contacts was used in image reconstruction to estimate the locations of neural activity in the nerve cross section. Correlating the locations of cardiac, pulmonary, and RL fascicles from microCT with changes in heart rate, respiratory rate, and laryngeal EMG, respectively, demonstrated that EIT and selective stimulation could approximately localize these functional groupings. Key remaining knowledge gaps include analogous data for the right VN, the organization of vagal fibers at the cervical level that innervate abdominal organs, and functional organization of the human VN, for which the fascicles are fewer, larger, and split/merge more frequently (Pelot et al., 2020; Upadhye et al., 2022).

Afferent innervation of glomeruli in mouse kidneys

Recent advances in treatment of hypertension include ablation of nerves innervating the kidneys; however, anatomy and function of afferent fibers in renal nerves are largely unknown. Tyshynsky et al. analyzed the anatomical relationship between afferent nerve fibers and renal glomeruli in mice using higher-throughput imaging for population counts and higher-resolution imaging for distance measurements. Renal glomeruli are specialized blood vessels that filter blood in the kidneys to create urine. The afferents were visualized using immunolabeling of calcitonin gene-related peptide (CGRP+) in samples from wildtype and transgenic mice, as well as fluorescence in TRPV1 lineage cells (tdTomato+) of transgenic mice. Approximately half of glomeruli had nearby CGRP+ or tdTomato+ axons, with higher likelihood for glomeruli deeper in the cortex. Bowman's capsule surrounds the glomerular capillary loops to filter blood and provide passage toward the ureters and urinary bladder. The afferents seemed to travel along the surface of Bowman's capsule, and thus they may transmit changes in glomerular pressure for reflexive responses to adjust vascular diameter. Additional studies are needed to confirm direct innervation and to elucidate the transduction and signaling roles of afferents in renal (patho)physiology.

Review of cardiac vagal afferents

Visceral organ injury and disease can cause pathological changes in the autonomic nervous system, which can in turn, affect visceral organ function. These pathological changes present an important consideration when evaluating therapeutic strategies using bioelectronic medicine to target the autonomic system, as highlighted by Van Weperen and Vaseghi. In this review, the authors summarize the role of cardiac vagal afferent/sensory neurotransmission in health, and they highlight important reported alterations in signaling during cardiovascular disease. They further demonstrate how a detailed mechanistic understanding of cardiac physiology and pathophysiology, combined with a wide range of advanced, clinically relevant animal models and techniques (e.g., Veerakumar et al., 2022) can provide unique model systems for investigating how to increase the therapeutic efficacy of targeted autonomic neuromodulation strategies under clinically-relevant conditions.

Efficient segmentation of fascicles from microCT of human vagus nerves

MicroCT imaging plays a pivotal role in unraveling the intricate three-dimensional arrangement of the fascicles of the human vagus nerve, a critical element for anatomical exploration and the progress of neuromodulation therapies. However, conventional manual segmentation methods to identify the fascicle boundaries can be laborious. In this issue, Buyukcelik et al. developed a U-Net convolutional neural network to automate the segmentation of fascicles within microCT images of the human vagus nerve. The U-Net structure was originally designed for segmentation of biomedical images (Ronneberger et al., 2015). With this U-Net, the authors accurately segmented approximately 500 images of a cervical vagus nerve in just 24 s. This study enables the use of deep learning fascicle segmentation of microCT images by employing a conventional U-Net as a benchmark. Although it can be further optimized, the U-Net described by Buyukcelik et al. will contribute to generating data to enrich computational models, thereby facilitating the advancement of analysis and the strategic design of neuromodulation therapies tailored specifically for the human vagus nerve.

Spatial arrangement of axons in autonomic nerves

Shemonti et al. report statistical methods for quantifying the spatial arrangements of axons in peripheral nerves, accounting for their intensity, interaction, and regionality. They used a published dataset of rat pelvic and vagus nerves imaged with transmission electron microscopy from which unmyelinated axons were delineated through manual segmentation. They present methods for spatial point pattern construction (an analysis used in many fields to measure spatial distribution and locations in two- or three-dimensional space), feature configuration (a set of features), and Sinkhorn distance computation (a mathematical/statistical tool for optimizing transport/distance between objects), encompassed in a generalized pipeline to analyze the inhomogeneous structure of peripheral nerves. Though the differences noted between the vagus and pelvic nerves and between nerve samples from male and female rats are of unknown physiological and biological significance, their method provides an objective technique

applicable to quantifications of axonal spatial arrangements in any nerve.

Stimulation of the dorsal root ganglion to treat cardiac sympathetic hyperactivity

Sustained sympathoexcitation can cause ventricular arrhythmias and sudden cardiac death. Kuwabara et al. examined neuromodulation of the dorsal root ganglion (DRG) as a means to dampen cardiac sympathoexcitation and ventricular excitability. Using a porcine model of acute myocardial ischemia (i.e., obstructed blood flow to the heart muscle), they found that DRG stimulation reduced sympathoexcitation caused by cardiac ischemia, consistent with their observation of reduced markers of neuronal activation in the dorsal horn of the spinal cord. Combined, these findings lay the preclinical groundwork to suggest that DRG stimulation may have important cardioprotective effects in humans, though additional data are required to know the extent and degree of potential benefit.

Stimulation of the spinal cord to treat cardiac sympathetic hyperactivity

Using a similar model of porcine myocardial ischemia, Salavatian et al. examined the potential of preemptive spinal cord stimulation to reduce sympathoexcitation and ventricular excitability caused by myocardial ischemia. The authors found that preemptive stimulation of the dorsal horn of the spinal cord reduced activity of ischemia-sensitive neurons during myocardial ischemia. Preemptive stimulation also reduced the firing activity of neurons within the intermediolateral column of the spinal cord, and it prevented the myocardial ischemia-induced augmentation of synchrony between dorsal horn and intermediolateral column neurons—where the intermediolateral column contains the cell bodies of neurons of the sympathetic nervous system. Though Salavatian et al. observed benefits in their study, additional studies are necessary to better understand the potential of spinal cord stimulation as a therapeutic intervention for cardiac diseases in humans and animals.

Conclusions

The studies highlighted in this special issue span two overlapping themes: (1) quantification of the anatomy of the peripheral autonomic nervous system at different scales, and (2) cardiovascular anatomy and neural stimulation treatments. The anatomical studies spanned functional organization of the pig vagus nerve, proximity of afferent fibers to glomeruli in mouse kidney, a review of cardiac vagal afferents, segmentation of fascicles in human cervical vagus nerve, and statistical descriptors for arrangement of axons in peripheral nerves. The latter two tools for image analyses can be applied to other parts of the peripheral nervous system. The cardiovascular studies spanned the role of kidneys in hypertension, locations of cardiac fibers in the mid-cervical pig vagus nerve, a review of cardiac vagal afferents in health and disease, as well as stimulation of either the dorsal root ganglion or spinal cord to treat cardiac sympathetic hyperactivity. While these studies advance our understanding of the autonomic nervous system, significant knowledge gaps remain in order to selectively apply neuromodulatory therapies at targeted nexus points to treat specific organ pathologies while reducing off-target effects.

Author contributions

NP: Conceptualization, Writing—original draft, Writing—review and editing. MV: Writing—original draft, Writing—review and editing. LR: Writing—original draft, Writing—review and editing. PO: Writing—original draft, Writing—review and editing. SC: Writing—original draft, Writing—review and editing.

Funding

Funding was provided by the NIH SPARC OT2OD025340, NIH SPARC 75N98022C00018, and NIH R01EB033403-02 for NP; NIH R01HL170626-01, NIH R34 HL153566, and AHA 970217 for MV; NIH R01HL152101, NIH OD026582, and Cystic Fibrosis Foundation REZNIKO20I0 for LR; NIH SPARC OT2OD023872 for PO; and Portuguese Foundation for Science and Technology EXPL/MED-NEU/0733/2021 for SC.

Conflict of interest

SC has the following patent: Conde SV, Chew DJ, Famm K, Guarino MP, Holinski B, Patel S (2015) Neuromodulation device. Patent PCT/PT2015/000047. International Bureau WO/2016/072875. SC was and is employed by NOVA Medical School.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

The author(s) declared that they were an editorial board member of Frontiers, at the time of submission. This had no impact on the peer review process and the final decision.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

References

Pelot, N. A., Goldhagen, G. B., Cariello, J. E., Musselman, E. D., Clissold, K. A., et al. (2020). Quantified morphology of the cervical and subdiaphragmatic vagus nerves of human, pig, and rat. *Front. Neurosci.* 14, 601479. doi: 10.3389/fnins.2020.601479

Ronneberger, O., Fischer, P., and Brox, T. (2015). "U-Net: convolutional networks for biomedical image segmentation," in *Medical Image Computing and Computer-Assisted Intervention-MICCAI 2015, 18th International Conference* (Munich: Springer International Publishing), 234–241.

Upadhye, A. R., Kolluru, C., Druschel, L., Al Lababidi, L., Ahmad, S. S., Menendez, D. M., et al. (2022). Fascicles split or merge every ∼560 microns within the human cervical vagus nerve. *J Neural. Eng.* 19, 054001. doi: 10.1088/1741-2552/ac9643

Veerakumar, A., Yung, A. R., Liu, Y., and Krasnow, M. A. (2022). Molecularly defined circuits for cardiovascular and cardiopulmonary control. *Nature* 606, 739–746. doi: 10.1038/s41586-022-04760-8