The role of vagal ischemia on the destiny of Peyer’s patches: first experimental study

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Abstract

Introduction: The vagal network has a major potential role in the immune-life of Peyer’s patches, but there is no satisfying information if vagal ischemia causes Peyer’s patches (PP) disruption following subarachnoid hemorrhage (SAH).

Methods: Twenty-two rabbits were used as control (GI, n = 5), “sham” (GII, n = 5), and SAH (GIII, n = 12) groups in this experiment. 0.5 cc saline for GII and 0.5 cc autologous blood for GIII was injected into cisterna magna of the rabbits. Four weeks later, they were euthanized. Their brains, vagal nerves, nodose ganglia, Peyer’s patches, and intestines were examined, using stereological methods. The Peyer’s patches volumes (PPVs)/intestine volume per cubic millimeter was accepted as PP injury score based on a total of 10 points.

Results: The mean degenerated neuron densities of the nodose ganglia and degenerated axon densities of vagal nerves were 5 ± 2/mm² and 6 ± 2/mm² in the GI, 13 ± 4/mm² and 89 ± 16/mm² in the GI and 321 ± 83/mm² and 293 ± 88/mm² in GIII. The mean PPVs and PP score were 8 ± 1×10⁶ µm³/mm³ and 0-3 in the GI, 10 ± 3×10⁶ µm³/mm³ and 1-5 in the GI, and 21 ± 5×10⁶ µm³/mm³ and 8-10 in GIII. P < 0.0001 in PPV/PP score/degenerated axon densities of vagal nerves; P < 0.0005 in PPV/PP score and degenerated neuron densities of the nodose ganglia between GI/GII; P < 0.001 in (PPV/PP score)/degenerated axon densities of vagal nerves; P < 0.005 in PPV/PP score/degenerated neuron densities of the nodose ganglia between GII/GIII; and P > 0.05 in GI/GII were noted.

Conclusion: Vagal ischemia/insult may be responsible for PP denervation, and injury-induced dangerous intestinal immunodeficiency following SAH.

Introduction

Peyer's patches (PP) host as a security castle for lymphoid cells or a trench for pathogens. They function as intestinal microspheres. PP and the vagal network containing the neuro-immunological web are essential components of intestinal immunity and are commonly located in the jejunum and ileum. PP shape the first intestinal barrier against pathogens. Vagal fibers are found in PP, and adjacent villi. Biological, physical, and chemical homeostasis are generally maintained by the vagal and splanchnic nerves. Also, Meissner’s and Auerbach’s networks, as well as thoracic spinal and solar ganglia, modulate all intestinal lymphoid organs. The PP blood supply is maintained by the mesenteric arteries. Mesenteric artery spasm, induced by sacral parasympathetic network damage, may cause ischemic intestinal degeneration. Onuf’s nucleus originated sacral parasympathetic web ischemia can be responsible for Hirschsprung-like disease following spinal subarachnoid hemorrhage (SAH). Sym pathetic hyperactivity induced by vagal collapse may cause intestinal atrophy following SAH. Because vagal nerve insufficiency causes dangerous pathologies in the intestinal system, vagal nerve stimulation may be an alternative application for neuro-inflammatory bowel disease and ischemic intestinal disease. The electromagnetic field created by the vagal nerve is necessary for the regulation of intestinal microbiota. We showed that vagal network ischemia might cause...
denervation injuries characterized by PP enlargement and reactivity induced follicular hyperplasia in the early phase. In due course, however, vasospasm of the PP supplying arteries causes PP edema, swelling, hemorrhage, and necrosis.

Materials and Methods

Animal-study model

This experimentally induced SAH model was studied on 22 rabbits. After examination, the animals were randomly divided into the following three groups: the control group (GI, n = 5); the “sham” group (GII, n = 5), which received 0.5 cc of saline; and the study group (GIII, n = 12), which received autologous blood injections (0.5 cc) in tapering doses into their cisterna magna one time/three days for two weeks.

Following the head anteflexion position for 10 minutes in foramen magnum, 0.5 cc of autologous blood was injected over about one minute via 22-gauge needle after 0.5 cc cerebrospinal fluid aspiration.

Isoflurane was administered by a face mask, and 0.2 cc/kg of the anesthetic combination (ketamine HCl, 150 mg/1.5 cc; Xylazine HCl, 30 mg/1.5 cc; and distilled water, 1 cc) was subcutaneously injected.

Clinical-experimental data collection and results

Heart rate, respiration rate, and blood pressure values were recorded for ten days. The rabbits were fasted for six hours before surgical intervention and decapitated after balanced injectable anesthesia was induced. Their brains, vagal complexes, and intestines were extracted just after intracardiac formalin injection and then fixed in 10% formalin solution. The intestines were examined with anatomical microscopy. For histopathological analysis, microsections were taken from the whole brains, vagal complexes, and intestines cross-cutting PP. Tissue sections (5 µm) were stained with hematoxylin-eosin (H&E), S-100, GFAP, and TUNEL methods. The advantages of these methods are as follows: They allow particle numbers to be counted accurately, and they have high specificity.

Histopathological procedures

To detect ischemic vagal lesions, brain materials were sectioned crossing the vagal motor nucleus. Vagal nerve axons and nodose ganglia were embedded in the same paraffin block and sectioned horizontally. Their numbers were estimated using the stereological method. The ileum segments were cut into 5 µm paired sections per 20 µm every 20th and 21st sections to calculate the PPV. All sections were stained with H&E, S-100, GFAP, and TUNEL methods and examined with light microscopy. The total PP numbers of the intestines were estimated with the fractionation method.

To estimate the total PP numbers and volume values, the sections were embedded in paraffin blocks. They were stained with H&E and immunohistochemical (S-100, GFAP, and TUNEL) methods. The advantages of these methods are as follows: They allow particle numbers to be counted accurately, and they have high specificity.

Table 1. Histopathological Scoring System of Peyer's Patches

| Score |
|-----------------|
| lymphoid aggregates without dendritic network under the surface epithelium | 0 |
| primary follicles with the dendritic network were observed under the surface epithelium | 1 |
| secondary follicular structures with prominent germinal centers under the surface epithelium | 2 |
| enlarged, interconnected, abundant secondary follicles detected in PP | 3 |
| lymphoid follicular atrophy or necrosis | 4 |
| Total Score | 10 |

Score <2: considered as normal, score 2-4: considered as hyperplasia, score 4-7: considered as gross enlargement PP, score >7; considered as invaded intestine by PP.
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be estimated easily. They are readily performed. They are intuitively simple. They are free from assumptions about particle shape, size, and orientation: and they are unaffected by overprotection and truncation.

**Stereological analysis**

Stereological methods provide reliable results to estimate particle number, material volume, and number. For this article, the stereological and Cavalieri methods have been explained in our previous studies.9,13,16

The total PPV was estimated by taking the sum of all PPVs. Each PP was considered an ellipsoid structure and its volume was calculated with the following formula:

\[ V_{pp} = \frac{4}{3}\pi \left( \frac{x + y + z}{3} \right)^3 \]

The x, y, and z are half of the radius values of the ellipsoid-shaped PP, which are coordinated on the x, y, and z apsis included in the analytical space. The total PPV was estimated using the following formula:

\[ \sum V_{pp} = \sum_{i=1}^{n} nxVn \]

The physical dissector method was preferred to evaluate the numbers of living and degenerated follicle cells of PP, nodose ganglia, vagal axons, and neurons of the vagal motor nuclei. The used analytical and geometrical methods have been clearly explained in our previous studies.9,13,15

The data obtained were analyzed with the SPSS software package (SPSS® for Windows v. 12.0, Chicago, USA). The one-way ANOVA test was used and the differences were considered to be significant at P < 0.05.

**Results**

**Clinical findings**

Meningeal irritation signs, such as neck stiffness, decreased coma score, convulsion, and fever, as well as cardiorespiratory disturbances, were reported.

**Anatomical and pathological findings**

Blurred-hemorrhagic brain surface, cortical swelling, sulcal narrowing with adhesion, bloody basal cistern, thrombosed blood vessels, gyral bombing, and less extend tentorial herniations were observed in some cases (Figure 1/Base).

**Histopathological findings**

Microscopic views showed constructed vasa vasorum, deformed vagal axons, apoptotic axons, and apoptotic nodose ganglia neurons in the rabbits with SAH (Figure 1). Brain stem sections included ischemic vagal motor nuclei just behind the aqueduct. Apoptotic neurons were also seen (Figure 2). Normal PP and villi with abundant myenteric plexuses were observed in normal rabbits (Figure 3). Minimally enlarged atrophic/degenerated villi-PP with reactivated focuses, and follicular dendritic cells were seen in the sham group (Figure 4). Enlarged inflamed pathologic PP with importantly atrophic/degenerated villi representing denervated/atrophic Meissner’s networks and reactivated focuses of PP with normal and deformed follicular dendritic cells were observed in the study group (Figures 5 and 6). An apoptotic neural network and stromal cells were observed in the rabbits with developed vagal ischemia (Figure 7). The PPHi calculation method

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**Figure 1.** Macroscopic view of the basal side of the brain with SAH (Yellow arrow), constructed vasa-vasorum (arrow) and deformed axons of the vagal nerve within jugular foramen (LM, H&E, x40/A); a cross-sectional area in a normal rabbit (LM, H&E, x40/B) and apoptotic axons (LM, Tunnel, x40/C) and apoptotic nodose ganglia neurons (LM, Tunnel, x10/D) are seen with a SAH created rabbit.

**Figure 2.** Histopathological view of the brain stem section including vagal motor nucleus (VMN) just behind of aqueduct (Aq) (LM, H&E, x4/Base; GFAP, x4/A); and apoptotic neurons (LM, Tunnel, x20/B) are seen in vagal ischemia developed rabbit.
that was used was designed by the authors (Figure 8). More ischemic vagal injury resulted in more intestinal injury such as myenteric network deformation, intestinal glandular atrophy, and regional necrosis.

**Numerical results**
The rabbits with SAH showed an unconscious state at the beginning of the SAH, and two of them were dead. The mean degenerated neuron density of the nodose ganglia, and axon densities were estimated, respectively, as $5\pm 2/mm^3$ and $6\pm 2/mm^2$ in the control group, $13\pm 4/mm^3$ and $89\pm 16/mm^2$ in the sham group, and $321\pm 83/mm^3$ and $293\pm 88/mm^2$ in the study group. The mean PPV and histopathological scores were $8\pm 3\times 10^6 \mu m^3/mm^3$ and 0-3 in the control group, $10\pm 3\times 10^6 \mu m^3/mm^3$ and 4-7 in the sham group, and $21\pm 5\times 10^6 \mu m^3/mm^3$ and 8-10 in the study group. Results and p values are summarized in Table 2.

**Discussion**
The immunomodulatory role of vagus on PP is the critical component of intestinal immunity commonly located in the jejunum and ileum. Vagal fibers are found in PP and adjacent villi. PP form the first intestinal barrier against pathogens. Physical and chemical homeostasis of the alimentary tract is sensed by the nodose ganglia and regulated by parasympathetic vagal fibers. The vagal and splanchnic nerves convey information to the

**Figure 3.** Histopathological view of the ileal section with normal Peyer’s patches (PPh) and Willi (LM, H&E, x4/Base) and abundant myenteric plexuses (Red arrow) (LM, S-100, x10/A) are seen in a normal rabbit.

**Figure 4.** Histopathological view of the ileal section with a minimally enlarged pathologic Peyer’s patches (PPi) with partially atrophic/degenerated Willi (LM, H&E, x4/Base; LM, S-100, x10/A); a magnified form of reactivated focus (LM, GFAP, x20/B) and follicular dendritic cells are seen in a rabbit from sham group.

**Figure 5.** Histopathological view of the ileal section with an enlarged, inflamed pathologic Peyer’s patches (PPi) with importantly atrophic/degenerated Willi (LM, H&E, x4/Base); representation of denervated/atrophic Meissner network (red arrow) and PP magnified form of reactivated focus with follicular dendritic cells (LM, H&E, x20) are seen in a study rabbit.

**Figure 6.** Histopathological view of the ileal section with an enlarged, inflamed pathologic Peyer’s patches (PPi) with importantly atrophic/degenerated Willi (LM, H&E, x4/A); representation of denervated/atrophic Meissner network and PP magnified form of the reactivated focus of PP (LM, H&E, x10/Base), normal (N) and deformed (D) follicular dendritic cells (LM, GFAP, x40/B).
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Central nervous system from mechanosensory in the intestinal tract regarding the feeding of nutrients and specific endocrine-motor patterns. PP can be considered intestinal microspleens. PP cells modulate antibiotic-bacteria interactions. Each PP has several elongated dome regions flanked by intestinal villi formed from lymphoid follicles covered with enterocytes. The interfollicular zone has high endothelial venules. Phagocytes include dendritic cells, which play a significant role in mucosal homeostasis. Dendritic cells are sensitive to the intestinal microenvironment and are at the front line in bacterial invasion. Dendritic cells have an essential role in the first adaptive immune responses. Mucosal layers of PP regulate microbiota colonization. PP have selective sensitivity to digested materials. Nonbiological particles undergo phagocytosis by PP cells. PP synthesize most of the immunoglobulins, especially IgA. Immunoglobulin synthesis primarily occurs in intestinal PP.

There is mounting evidence for interactions between the PP and the vagal system. Lymphoid tissues are well supplied by somatic and autonomous nerves. Oral microbiota has great importance and is regulated by the olfactory, trigeminal, and lower cranial nerves. All intestinal lymphoid organs are innervated by the myenteric and solar plexuses as well as the thoracic spinal ganglia because immune modulation of the enteric nervous system is required for a dense neural network. To demonstrate the enteric nervous system and PP, specific markers, such as the S-100 protein, GFAP protein, and TUNEL methods, were used. This has been mentioned by Krammer et al in their research.

Dangerous histomorphological changes occur in denervated PP, especially with vagal ischemia, because vagal network ischemia results in intestinal atrophy. Sacral parasympathetic network ischemia causes mesenteric artery spasm related intestinal degeneration. Onuf’s nucleus originated from sacral parasympathetic network ischemia, can be responsible for Hirschsprung-like disease following spinal SAH. Olfactory bulb lesions

Table 2. Numerical results of the study

|                      | Group I (n = 5) | Group II (n = 5) | Group III (n = 12) |
|----------------------|----------------|-----------------|-------------------|
| Degenerated neuron density of nodose ganglia (mm$^3$) | 5 ± 2          | 13 ± 4          | 321 ± 83$^a$      |
| Degenerated axon densities (mm$^3$)                  | 6 ± 2          | 89 ± 16$^b$    | 293 ± 88$^a$      |
| Mean volumes of Peyer’s patches (x10$^6$µm$^3$/mm$^3$) | 8 ± 1          | 10 ± 3$^c$     | 21 ± 5            |
| Histopathological score                                | 0-3            | 4-7$^c$        | 8-10$^c$          |

The values are given as mean ± standard deviation or median (minimum-maximum), Group I: Control Group, group II: SHAM group, group III: study group.

$^a$ $P<0.0001$ Group III vs. I and II one-way ANOVA test.

$^b$ $P<0.0001$ Group I vs. II one-way ANOVA test.

$^c$ $P<0.05$ Group I vs. II one-way ANOVA test.

Figure 7. Histopathological view of the ileal section with apoptotic neural network and stromal cells (arrow, L&M, Tunnel, x10) in vagal ischemia developed rabbit.

Figure 8. Peyer’s patches hyperplasia index (PPHi) calculation method designed by ourselves shown as an equation $PPHi = \frac{\sum PP}{\sum x_n}$. Each PP is like a cylinder or sypher, and their volumes may be estimated as the following common formula: $V_{PP} = \pi \left( \frac{4}{3} \pi \left( x + y + z \right)^3 \right)$; total PP volumes were estimated as a formula $V = \pi \sum \frac{4}{3} \pi \left( x + y + z \right)^3$; a cylindrical segment volume estimated as a formula $V_{SI} = \pi \left( R^2 - r^2 \right)$. Because PP become enlarged owing to various immunological phenomena following vagal ischemia, PP volume values will increase, and also $PPHi$ values will be increased. Consequently, the value of the equation will be greater than zero. According to our standardization the equation should be under 0.01. If the equation is over 0.2, there is an enormous PP hyperplasia; it is equal to zero, necrosis will be started. If it approaches infinity owing to extremely PP hyperplasia, the intestine is dead. Constructed mesenteric arteries aggravate PP and intestinal necrosis.
could rely on denervation injuries of PP affected by vagal software. Gut microbiota change in a dangerous manner following PP-vagal network lesions. PP is the most affected part of the intestine following radiotherapy; probably due to an increased intestinal electromagnetic field with the summation of magnetic fields. This theory of Wang, who justified Nikola Tesla's theory, supports our theory.

Vagal nerve stimulation may be an alternative application for immune suppression in neuroinflammatory bowel disease, a condition in which PP are the essential mediators. A century ago, the great genius Nicola Tesla declared that the value of electromagnetic fields is the most crucial factor for all living things. Based on his theorem, the electromagnetic field created by the vagal nerve is an indispensable necessity for the presence of intestinal microbiota.

PP regulate the energy production, proliferation, effector functions, and metabolism of immune cells in response to the immunologic stimulus. Intestinal ischemia is a necessary pathophysiologic process for PP dysfunction. Maternal colostrum, and milk immunoglobulins are essential for the development of strong immunity in jejunal and ileal PP. Also, Liu et al. declared that ischemic stroke damages the intestinal epithelium as well as immunity. Invasion by parasites, bacteria, or dangerous particles is frequently seen following ischemic intestinal disease. Vagal nerve-breast network is essential for lactation, breastfeeding and the development of baby immunity.

Limitations Better results could be obtained by conducting bacteriologic, electrophysiologic, radiologic, and biochemical examinations.

Conclusion The PP-vagal network is an excellent barrier against internal and external hazardous agents with its anti-infective, anti-tumoral, and anti-poisoning effects through its neuro-immunomodulatory functions. Although the mentioned functions of the vagal network are well known, there is no satisfying information as to whether or not vagal ischemia results in intestinal immune system hardware/software disruptions following SAH. This article demonstrates that SAH-induced vagal ischemia may be responsible for intestinal immunodeficiency by way of PP denervation injury.

Future insight Vagal nerve stimulation or intestinal transplantation would be used for intestinal immune deficiency situations in the future.

Conflict of Interest Authors declare no conflict of interest in this study.

Ethical Approval Ethical approval was obtained from the ethical committee of Ataturk University Faculty of Medicine (19.09.2017/1700254064).

Authors' contributions NA, EK, and MDA carried out the design, coordinated the study, and participated in the experiments. EOA participated in anesthesia and follow-ups. SO, OC, ED, TD participated in the examination of histopathological data. MDA, aided in statistical analysis and manuscript preparation. IM and MDA assisted in data gathering and participated in manuscript editing.

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