Location and distribution pattern of nitric oxide synthesizing neuronal cells in the digestive tract and urinary bladder of mice

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Abstract

In order to investigate further the location and distribution pattern of NOS containing neuronal cells in the peripheral autonomic nerve system of digestive tract and urinary bladder, the digestive and bladder organs were stained by NADPH diaphorase-histochemical staining. Some groups of NADPH diaphorase positive neuronal cells were detected along all parts of the digestive tract, including esophagus, stomach, small and large intestine, as well as urinary bladder. In the digestive tract, these cells were existed in a very small number in the submucous layer (submucous plexuses of Meissner) and ubiquitous in the area between the external and internal layer of smooth muscle (myenteric plexuses of Auerbach). In the bladder, those cells existed at the bladder neck, and in the inner layer of the muscle of urinary bladder. Since the number of NADPH diaphorase containing neuronal cells per millimeter square is more numerous in stomach and large intestine, there is a possibility that the amount of nitric oxide in those organs is also higher. These facts suggest an important role for nitric oxide (NO) in those organs. Furthermore, the existence of those cells in the limited area of urinary bladder i.e. in the bladder neck may suggest the role of nitric oxide (NO) in vesico-urethral sphincter muscle function.

Keywords: NADPH diaphorase, nitric oxide synthase, submucous plexuses of Meissner, myenteric plexuses of Auerbach, bladder neck

Nitric oxide (NO) is a simple gas with free radical chemical properties, but it is often confused with the chemically distinct nitrous oxide (N₂O) which is used as an anesthetic and is chemically stable. Nitric oxide is an important messenger molecule produced by endothelial cells, neurons, skeletal muscle, neutrophils, macrophages, pancreatic islets, endometrium, epithelium of respiratory and gastrointestinal tract, and others.¹ Nitric oxide is an unusual mediator because it is a gas with unknown storage mechanism. It diffuses freely across membranes, and is extremely labile with a biological half life on the order of seconds.

No binds to and activates soluble guanylyl cyclase resulting in increased cGMP level. The exact mechanism by which cGMP exerts its biological effects are still unknown, although cGMP dependent protein kinases may be involved. Nitric oxide also modifies a critical cysteine in glyceraldehyde 3-phosphate dehydrogenase by adenosine diphosphate ribosylation,²³ or S-nitrosylation via NAD interactions.⁴
Nitric oxide also binds to a variety of iron- and sulfur-containing proteins.

Nitric oxide is synthesized via catalytic actions of nitric oxide synthase (NOS). Several homologous but separate genes code for neuronal NOS (nNOS), endothelial NOS (eNOS), and macrophage-derived NOS (also known as inducible NOS/iNOS). Neuronal NOS and endothelial NOS are produced constitutively and are calmodulin dependent; their activity is stimulated by calcium. Those enzymes also need reduced nicotinamide adenine dinucleotide phosphate (NADPH) as an electron donor in their catalytic activity. Inducible NOS are produced inducibly by glucocorticoid and cytokines like gamma interferon and lipopolysaccharide, and does not depend on calcium for its activity. All forms of NOS use arginine as their substrate, and form NO and citrulline stoichiometrically.

In neurons, NOS colocalize with NADPH diaphorase. Hope et al. presented strong evidence for the idea that NADPH diaphorase may correspond to neuronal nitric oxide synthase. This NADPH- and Ca"/ Calmodulin-dependent enzyme (NOS) forms NO by conversion of arginine to citrulline. In the presence of NADPH diaphorase, the reduced form of nitric oxide synthase can catalyze the conversion of a soluble tetrazolium to an insoluble visible formazan. This multiple step reaction used in NADPH diaphorase histochemical staining is illustrated in the schematic figure below.

This NADPH diaphorase histochemical staining is resistant to formaldehyde fixation and displays the characteristics of an enzymatic reaction.

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\begin{align*}
O_2 + \text{Arginine} + \text{Ca}^{2+} / \text{CaM} & \rightarrow \text{Arginine-OH} + \text{Citrulline + NO} \\
\text{NOS-flavin} & \rightarrow \text{NOS-flavinH2} \\
\text{NADPH} & \rightarrow \text{NBT} \text{formazan} \\
\text{CaM} & = \text{calmodulin} \\
\text{NADPH} & = \text{nicotinamide adenine dinucleotide phosphate} \\
\text{NO} & = \text{nitric oxide} \\
\text{NOS} & = \text{nitric oxide synthase} \\
\text{NBT} & = \text{Nitro blue tetrazolium}
\end{align*}
\]

NOS initially receives the electrons from NADPH in a calcium/calmodulin- and arginine-independent step that presumably involves reduction of one or both of the associated flavins. The reduced NOS will be rapidly re-oxidized in the presence of an appropriate electron acceptor, such as nitro blue tetrazolium (NBT). In the presence of calcium, calmodulin and molecular oxygen, the reduced NOS will hydroxylate the guanidino group of arginine in a P-450 like monoxygenase reaction. The tightly associated hydroxyarginine is then further oxidized by NOS to citrulline and NO.

In the nervous system, nNOS is located to discrete populations of neurons in the cerebellum, olfactory bulb, hippocampus, cortex, striatum, basal forebrain, and brainstem. NOS is also concentrated in the posterior pituitary gland, supra-optic and paraventricular hypothalamic nuclei, and in discrete ganglion cells of the adrenal medulla.

The localization and distribution pattern of NOS containing neuronal cells in the peripheral autonomic nervous system, especially in the digestive tract and urinary bladder are still unclear. Gabriella et al. failed to detect intramural ganglia in the urinary bladder of the mouse. However, Grozdanovic et al. claimed that they had successfully demonstrated the occurrence of intramural neurons in the lower urinary tract of the mouse. The knowledge of localization and distribution pattern of NOS containing neuronal cells in the digestive tract and urinary bladder will give implication and more stimulation to study the role of nitric oxide on the function of the digestive tract and urinary bladder.

In order to investigate further the localization and distribution pattern of NOS containing neuronal cells in the peripheral autonomic nervous system of digestive tract and urinary bladder, we stained both whole mount and sections of those organs with NADPH diaphorase histochemical staining and counted the number (density) of positive cells per millimeter square in the digestive tract and the total number of those cells in the urinary bladder. We also discussed the possible function of NO in those organs and the possible correlation between the localization and distribution pattern of those positive cells and their natural functions.

**METHODS**

**Animal**

Mice (C57BL/6xDBA/2, F1 mice) were purchased from Japan SLC (Hamamatsu, Japan). Mice were
maintained in the animal center in a specific pathogen free condition. In this experiment 3 male and 3 female mice were used in both group (the experimental and control group).

Sectioned NADPH diaphorase histochemical staining

Using a wing needle, mice were cardiachly perfused with 0.1 M phosphate buffer (pH 7.4) containing 4% paraformaldehyde. The isolated tissues were then post-fixed with 4% paraformaldehyde equilibrated with 20% sucrose, for 12 hours. Tissue blocks for cryostat sections were prepared by embedding each tissue into a tissue compound and allowed them to be hardened in the cryostat chamber. Using a cryostat, these blocks were sectioned at 10 µm, and mounted on the glass slides. The slides were then washed with 0.1 M phosphate buffer (pH 7.4) for 3-5 minutes, and put on the rack underlided with wet paper towel.

In the experimental group, after washing, the slides were incubated in 0.1 M phosphate buffer (pH 7.4) containing 1.0 mg/ml β-NADPH (Sigma, St. Louis, MO), 0.1 mg/ml nitro blue tetrazolium (NBT) and 0.3% Triton X-100, for 60 minutes, at 37°C. In the control group, the same procedure was performed, except the last step. After washing, the control slides were incubated in 0.1 M phosphate buffer (pH 7.4) containing only 0.1 mg/ml NBT and 0.3% Triton X-100, for 60 minutes, at 37°C.

Whole mount NADPH diaphorase histo-chemical staining

Digestive and urinary bladder tissues were isolated from cardiachly perfused mice as described above, and post fixed with 4% paraformaldehyde for 12 hours. In the experimental group, after washing with PBS, whole tissues were incubated with 0.1 M phosphate buffer (pH 7.4) containing 1.0 mg/ml β-NADPH (Sigma, St. Louis, MO), 0.1 mg/ml nitro blue tetrazolium and 0.3% Triton X-100, for 90-120 minutes, at 37°C. In control group, after washing with PBS, whole tissues were incubated with 0.1 M phosphate buffer (pH 7.4) containing only 0.1 mg/ml NBT and 0.3% Triton X-100, for 90-120 minutes, at 37°C.

The Number of NADPH diaphorase positive neuronal cells

The number of NADPH diaphorase positive neuronal cells in the urinary bladder is very little (only a single cell or a cluster of ganglion cells) while they were particularly numerous in the bladder neck regions. Therefore, in the whole mount samples, total number of those cells was counted directly in all microscopic fields, under light microscope. The number per millimeter square (density) of those cells in several different parts of the digestive tract were counted by using enlargement of microphotographs of defined organs. The average and standard deviation values of the density or total number of those positive cells in the different parts of the digestive tract or urinary bladder were statistically calculated from 6 different observed animals.

RESULT

To investigate the localization and distribution pattern of nitric oxide synthesizing neuronal cells in the digestive tract and urinary bladder, histochemical staining method to detect NADPH diaphorase was performed. The diaphorase positive neuronal perikarya, neuronal fibers and epithelium cells of the mucosal layer showed blue precipitates obtained from the conversion of nitro blue tetrazolium into NBT formazan (Figure 1 and 2). These blue precipitates were clearly located only in the cytoplasm of neuronal cells and their nerve fibers.

Digestive tract

The numerous groups of NADPH diaphorase positive neuronal cells were detected along all part of the digestive tract, including esophagus, stomach, and small and large intestine (Figure1 and 2A,C). These cells existed in very limited number in the submucous plexuses of Meissner located in the submucosal layer. Furthermore, they were ubiquitous in myenteric plexuses of Auerbach located in the area between the external and internal layer, of the smooth muscle coat (Figure 1 square box areas, and Figure 2C arrow marked). The epithelial cells in the mucosal layer along all parts of digestive tract were also reactive for this NADPH diaphorase histochemical staining. No NADPH diaphorase positive cell was found in the digestive tissues of the control samples.

In the large intestine (Figure 2A and C) many diaphorase positive neuronal perikarya were located within the primary meshwork of the myenteric plexus of Auerbach, and in the area between the external and internal layer of the muscle coat. These neuronal cells
gave the innervation to the muscle layer of the digestive tract through the neurites and also to the submucous plexus of Meissner. The neuronal fibers which connected the neuronal cells one another could be detected more clearly in the large intestine than those in the other parts of the digestive tract. These NADPH diaphorase positive neuronal cells belong to bipolar and multi-polar neuronal cells (Figure 1 and 2 A, C).

Urinary bladder

Neuronal perikarya staining for NADPH diaphorase particularly existed in the urinary bladder at the bladder base and bladder neck region (around the ureteral orifices and in the floor of the bladder). These cells were located inside of the muscle coat of the bladder wall (Figure 2B and D). As in the digestive tract, these NADPH diaphorase positive cells had also fibers that connected the neuronal cells one to another. Epithelial cells of urinary bladder (urothelium) also exhibited positive reaction for NADPH diaphorase staining. All of the control samples exhibited negative reaction for NADPH diaphorase staining.

The number of NADPH diaphorase reactive cells in the digestive tract and urinary bladder

The number of NADPH diaphorase positive neuronal cells varied among parts of the digestive tract. The density of those positive cells was higher in the large intestine and was very low in the esophagus (Figure 3). Since the number of those positive cells was very little in the bladder, we counted all of the positive cells presented in the bladder directly from the whole mount samples under light microscopy. The density of those cells per-millimeter square in the different parts of digestive tract were 9 ± 0.54 (esophagus); 45 ± 3.89 (stomach); 31 ± 1.14 (small intestine); and 121 ± 12.63 (large intestine). Mean-while, the total number of those cells in the urinary bladder was 59 ± 14.35.

Figure 1. Microphotographs of NADPH diaphorase positive neuronal cells in the esophagus (A and D), stomach (B and E), and small intestine (C and F). (A, B, and C) are whole mount staining; (D, E, and F) are sectioned-NADPH diaphorase histochemical staining. The areas in the small square-boxes (D, E, and F) indicate the location of neuronal cells with positive NADPH diaphorase histochemical staining. MCS, SbMCS, and MUSC indicate the mucous, submucous, and muscle layers. Magnification: 60x (A, B, and C); 20x (D, E, and F).
Figure 2. Microphotographs of NADPH diaphorase positive neuronal cells in the large intestine (A and C) and urinary bladder (B and D). (A and B) are whole mount staining; (C and D) are sectioned-NADPH diaphorase histochemical staining. Arrow marks indicate the location of the NADPH diaphorase positive fibers and neuronal cells in large intestine (A and B), and urinary bladder (B and D). MCS, and MUSC indicate the mucous and muscle layer. Magnification: 100x (A and B); 20x (C), and 50x (D).

Figure 3. Bar-graphic diagram of the number of neuronal cells with NADPH diaphorase positive staining per-millimeter square in several different parts of digestive tract. The density of neuronal cells per-millimeter square is highest in the colon (large intestine) and lowest in the esophagus.
DISCUSSION

The location of the blue precipitates in the cytoplasm and nerve fibers indicate that nitric oxide synthase (NOS) and NADPH diaphorase belongs to cytoplasmic proteins. However, not only the neuronal cells, the epithelial cells of mucosal layer in the digestive tract and urinary bladder were also positively stained. These results were similar to the other previous reports. So far, there is no report describing whether these cells really contain nitric oxide synthase or not. The correlation between the existence of positive NADPH diaphorase staining in the epithelium layer and their function is still poorly understood. In their histochemical studies, Bredt et al could not find nitric oxide synthase in parenchymal or stromal cells of the liver, heart, lung, spleen, kidney, thymus, testes, or salivary glands. Despite these observations, it can not be excluded that mucosal NADPH diaphorase (that represents an isofrm of nitric oxide synthase) is not readily detectable with the antisera raised against the neuronal nitric oxide synthase. In fact, these antisera did not react with nitric oxide synthase of activated macrophages. Furthermore, if the epithelial cells really contain nitric oxide, it is supposed that NO is involved in the cellular process of epithelium cells.

Although NADPH diaphorase activity was detected in certain nerve cells and neuronal processes along the digestive tract (Figure 1), the greatest density of diaphorase positive neuronal cell, which was indicated as the number of NADPH diaphorase positive neuronal cells per-millimeter square (Figure 3), was found in the large intestine, followed by that in stomach. The NADPH diaphorase positive enteric neuronal cells in the sub-mucous layer were also easier to be identified in the large intestine (Figure 2 and C). Since the number of NADPH diaphorase positive neuronal cell was higher in stomach and large intestine than in other parts of the digestive tract (Figure 3), it is supposed that nitric oxide (NO) as a neurotransmitter is required in a larger amount by the smooth muscle of the stomach and large intestine for their adequate movement or peristalsis during the digestion process of food in the stomach and the excretion process of feces from the large intestine.

Several studies revealed that nitric oxide, which was released from inhibitory non-adrenergic non-cholinergic nerve, served as an inhibitory non-adrenergic non-cholinergic (NANC) neurotransmitter in several autonomic functions. Nitric oxide is involved in mediating relaxation of smooth muscles of the digestive tract, which is necessary for facilitating the passage of material, changing food to a bolus and generating peristalsis in the digestive tract. Mice lacking the nitric oxide synthase (NOS) gene exhibited grossly enlarged stomachs, with hypertrophy of the pyloric sphincter and the circular muscle layer. Meanwhile, mice with hyperinnervation of NADPH diaphorase positive enteric neuronal cells exhibited megacolon phenomena with prolonged transit time of barium throughout the gastrointestinal tract during barium enema (our unpublished data).

The NADPH diaphorase positive neuronal cells in the urinary bladder consisted of single cell or group of cells, which were particularly numerous in the region of bladder base or bladder neck (around the ureteral orifices and in the floor of the bladder). Those cells, which are located in the wall of urinary bladder are considered to be the source of the parasympathetic post-ganglionic nerve supply to this organ. Since the vesico-urethral sphincter muscle is also located almost in the same area as NADPH diaphorase positive neuronarl cells, nitric oxide is thought to be a neurotransmitter necessary for facilitating the relaxation of vesico-urethral sphincter. However, its exact function in vivo remains ill-defined. Studies in fetal sheep demonstrated that nitric oxide inhibition caused bladder hyperactivity and increased bladder capacity, perhaps by preventing sphincter relaxation. In rat, inhibitors of the NO system caused bladder hyperactivity and decreased bladder capacity. In our observations, mice with hyper-innervation of NADPH diaphorase positive neuronal cells showed decreased bladder capacity and decreased threshold pressure, perhaps by hyperrelaxation of vesico-urethral sphincter muscle (our unpublished data).

The exact function and role of nitric oxide (NO) as a nonadrenergic noncholinergic (NANC) neurotransmitter in various tissues and organs of the body is still far from understood. The pharmacological effects of nitrates are used in clinical conditions, e.g. ischemic heart diseases, hypertension, and motor disorders of esophagus. Furthermore, nitroglycerin is used in Germany (and possibly also in other places) as an antispasmodic agent in patient complaining of biliary colic. These observations are interesting because recent animal studies suggest that the choleystokinin-induced relaxation of the sphincter of Oddi is mediated by the NANC neural pathway operated by an unidentified transmitter substance. Burnett et al reported that NO was a physiologic mediator of erectile function. Neuronal NOS (nNOS) was located...
in rat penile neurons innervating the corpora cavernosa and in neuronal plexuses in the adventitial layer of penile arteries. Small dose of NO synthetase inhibitors abolished electrophysiologically induced penile erections. Ferrante et al. reported that NADPH diaphorase positive neurons were selectively spared in neurodegenerative diseases such as Huntington’s diseases. Huang et al. reported that mice lacking nitric oxide synthase (NOS) gene developed grossly enlarged stomachs with hypertrophy of the pyloric sphincter and the circular muscle layer, resembling infantile pyloric stenosis in human, in which gastric outlet obstruction was associated with the lack of NADPH diaphorase neurons in the pylorus. Hatano et al. and Shirasawa et al. reported that mice lacking Ncx/Hox 11L1 gene developed megacolon with hyperinnervated enteric ganglia. Those mice can be used as a model for human neuronal intestinal dysplasia (NID) diseases, in which myenteric neuronal hyperplasia and megacolon are seen. Finally the impact of the results of researches in nitric oxide on clinical aspect of diseases, in which nitric oxide (NO) may play a special role as the cause, is very important and might be useful to find the therapy for such diseases.

CONCLUSION

The numerous groups of NADPH diaphorase positive neuronal cells were detected along all parts of the digestive tract, including esophagus, stomach, small and large intestine as well as urinary bladder. These cells existed in a very small number in the submucous plexuses of Meissner located in the submucous layer, and ubiquitous in myenteric plexuses of Auerbach located in the area between the external and internal layer of the smooth muscle of the digestive tract. In the bladder, those cells existed at the bladder neck, inside of the muscle coat of urinary bladder.

Since the number of NADPH diaphorase positive neuronal cells per millimeter square is more numerous in stomach and large intestine, there is a possibility that the amount of nitric oxide in those organs is also higher. This fact suggests the important role of nitric oxide (NO) in those organs. Furthermore, the existence of those cells in the limited area of urinary bladder i.e. in the bladder neck may be correlated with the function of nitric oxide (NO) for vesico-urethral sphincter muscle.

Acknowledgements

The author would like to express the appreciation and special thank to Dr. Masahiko Hatano MD, Division of Developmental Genetics, Chiba University, Graduate School of Medicine, for his kind technical guidance, and the supply of animals and materials; and to Mrs. W. Indarty Aulia for her help in preparing and typing this manuscript.

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