Extrahypothalamic Projection of Immunoreactive Vasotocin Fibers in the Brain of the Toad, *Bufo japonicus*

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ABSTRACT — Extrahypothalamic projection of vasotocin (AVT) fibers in the brain of the toad (*Bufo japonicus*) was examined immunohistochemically by the avidin-biotin-peroxidase complex (ABC) method. Immunoreactive AVT perikarya are localized in the nucleus preopticus pars magnocellularis. The AVT neurons send their immunoreactive varicose fibers to many discrete brain regions, such as the limbic cortex, the thalamus, the optic tectum and the lower brain stem, in addition to the neurohypophysis. A dense network of AVT fibers was found in the septal nuclei and the anterior part of the preoptic nucleus. AVT fibers which run postero-dorsad project to the nucleus posterocentralis thalami, the nucleus posterodorsalis tegmenti mesencephali, and the nucleus isthmi. Meanwhile, AVT fibers which run through in the dorsal infundibular region and then the mesencephalic reticular formation are distributed in the medulla oblongata. These findings suggest that AVT acts as a neuromodulator or a local hormone in the toad brain.

INTRODUCTION

It is well established that, in anuran amphibians, vasotocin (AVT) has both antidiuretic and vaso­pressor effects. In addition, AVT shows pronounced effects on reproductive behavior in *Rana pipiens* [1] and *Taricha granulosa* [2]. In *T. granulosa*, AVT may be involved in control of sexual behavior by acting neurons in the central nervous system [3]. We have shown in the toad brain that vasotocin neurons project their varicose axons into the anterior part of the preoptic nucleus (APON) which is considered to be the triggering center for male mate calling behavior in anuran amphibians [4]. These results suggest that AVT neurons transmit APON neurons peptidergic information concerned with initiation of sexual behavior.

In mammalian brains, the distributions of various neurohormones including arginine-vasopressin (AVP) are not confined only in the hypothalamo-neurohypophyseal system. They are widely distributed throughout in discrete brain loci [5]. Ultrastructural studies showed that axon endings of immunoreactive (ir) luteinizing hormone-releasing hormone (LHRH) fibers [6] and ir-AVP ones [7] form synapses and synaptoid contacts with other neurons. Further, varicose LHRH fibers form en passant synapses in the preoptic area of the guinea pig [6]. Therefore, it is probable in amphibian brains that, in addition to the APON and the neurohypophysis, AVT neurons send their fibers to many extrahypothalamic regions.

In this study, we examined immunohistochemically the extrahypothalamic distribution of AVT fibers in the toad brain to learn whether AVT neurons project their axons to the loci which are related to control of mating behavior. Further, we have tried to elucidate phylogenetically fundamental distributional pattern of AVT in the vertebrate brain, since the amphibian brain is considered to show fundamental structural organization of the vertebrate brain [8].

MATERIALS AND METHODS

Adult toads (*Bufo japonicus*) of both sexes, body weight ranging from 117 to 303 g, and body
length (snout to vent) from 11.9 to 14.1 cm, were used. These animals were either obtained from an animal supplier in late September or were captured at the breeding season. Three animals, which were obtained from an animal supplier in late November, were used for immunohistochemical staining of thick frozen sections.

**Immunohistochemical procedure**

Distribution of AVT was immunohistochemically localized in paraffin sections and thick frozen sections that were cut either transversally, sagittally or horizontally to determine exact loci where ir-A VT fibers were found. As for immunohistochemical staining of vasotocin in the paraffin sections of the toad brain, details of fixation, tissue preparation and immunohistochemical procedure, in which the Vectastain ABC kit (Vector) was used, have been described previously [9, 10].

For the staining of thick frozen sections, the brains were fixed by transcardial perfusion with a fixative solution containing 1% glutaraldehyde, 2% paraformaldehyde and 4% sucrose in 0.1 M phosphate buffer (pH 7.4). The brains were removed, postfixed in the same fixative at 4°C overnight, and were washed in 0.1 M phosphate buffer. Frozen sections were cut at 50 μm, and were washed in 0.1 M phosphate buffer. Frozen sections were cut at 50 μm, and were washed in 0.1 M phosphate buffered saline (pH 7.4) at room temperature for 30 min. Then, they were stained immunohistochemically by use of the Vectastain ABC kit. The sections were first incubated in the avidin-biotin-peroxidase complex in PBS-T for 30 min. After a brief incubation in a blocking medium at 4°C for 12–16 hr, the solution for incubation contained anti-arginine-vasopressin serum (1: 8000 dilution, Bioproducts), which cross-reacts completely with vasotocin but not with neurotocin (cross-reactivity, 0.3%), in 0.1% Triton X-100 dissolved in 0.1 M phosphate buffered saline (PBS-T). After the incubation with the primary antiserum, the sections were washed in PBS-T at room temperature for 30 min, incubated with biotinylated anti-rabbit Ig-G for 1 hr, and were washed in PBS-T. Afterward, the tissue sections were incubated in the avidin-biotin-peroxidase complex in PBS-T for 30 min. After a few rinses, they were incubated in DAB solution including 0.05% 3,3'-diaminobenzidine (Sigma) and 0.01% hydrogen peroxide for 10 min, washed briefly in phosphate buffer, and were mounted on slide glasses with 40% ethanol containing 0.75% gelatin. They were then dehydrated, and were cover-slipped with Permount (Fisher). The tests for specificity of immunohistochemical staining followed the previous study [4, 9].

Nissl stained tissue sections were referred to for describing precise localization of ir-A VT. Nomenclatorial usage in this paper is basically those in *Rana pipiens* [11] and *Biuro japonica* [9].

**RESULTS**

**Distribution of ir-A VT perikarya and fibers**

As was described previously [4, 9], ir-A VT perikarya are localized in the ventral (VMC) and dorsal (DMC) magnocellular parts of the preoptic nucleus (Figs. 1 and 5). Beaded or varicose ir-A VT fibers were widely distributed among the discrete extrahypothalamic loci in the limbic system and the brain stem (Figs. 2–4). They were not found in the dorsal and anteroventral regions of the telencephalon. The extrahypothalamic ir-A VT projections can be classified roughly into three groups according to their destinations (see Fig. 5). Neither notable seasonal variation nor sexual difference was found in the distribution of ir-A VT fibers in this study.

**Projection to the telencephalon** (Figs. 5–8)

A part of ir-A VT fibers emanating from the VMC project to the postero medial region of the telencephalon, principally to the nuclei medialis septi and lateralis septi. Ir-A VT fibers are sent out to these loci mainly through the white matter including the medial forebrain bundle which surrounds the neuronal cell mass of the APON. A
found in the telencephalon have a varicose form, however, they rarely show Herring bodies which are frequently observed in the magnocellular preopticohypophyseal neurosecretory system.

**Projection to the thalamus and the tectum (Figs. 5, 8-9).**

Ir-A VT fibers which project to the brain stem arise from ir-A VT neurons in the VMC. They initially proceed laterad into the white matter in the preoptic region. Then, they turn their destination caudal to the direction of the lower brain stem with many ir-A VT fibers that project to the neurohypophysis. Thereafter, the fibers projecting to the brain stem diverge from the preoptico-neurohypophyseal tract around the dorsal infundibular region. A few beaded ir-A VT fibers are localized in the nucleus infundibularis dorsalis (Fig. 2). Many ir-A VT fibers run down to the mesencephalic tegmentum. They project to the nuclei posteriorodorsalsis tegmenti mesencephali and isthmi, and further to the griseum centrale rhombencephali. A considerable number of fine beaded fibers gather together to form a plexus in the region anterior to the nucleus isthmi (Fig. 3).

**Projection to the brain stem (Fig. 5, 10-11).**

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**DISCUSSION**

The present study showed that ir-A VT fibers are distributed widely among many extrahypothalamic loci in the toad brain. Such regions are the limbic cortex, the thalamus, the optic tectum, the isthmic region, and the lower brain stem.

The distributional pattern of extrahypothalamic ir-A VT fibers in the toad brain seems to be homologous to those described in the brains of other vertebrate classes. In the rat and the monkey, vasopressin neurons, the mammalian counterpart of AVT neurons, project their immunoreactive fibers to the hippocampus, the septum, the amygdala, and the preoptic area. They
project further to the thalamus, the superior nuclei [12]. A similar distributional pattern of ir-A VT fibers was observed in the brain of the lizard Gekko geko [13] and the eel Anguilla japonica (Fujiwara et al., unpublished).

A radioimmunoassay study of microdissected brain areas of rough-skinned newts also showed a similar distributional pattern of AVT fibers in the brain of the lizard Gekko geko [14]. These results indicate that the patterns of extrahypothalamic projections of AVT and vasopressin are fundamentally homologous in all vertebrate classes.

Immunoelectron microscopic studies demonstrated the presence of synapses containing neurohypophysial hormones in the rat brain [7, 15]. In the previous study, we showed that LHRH and AVT fibers may contact synaptically with APON neurons [4]. It is therefore highly probable that, in the toad brain, the extrahypothalamic AVT fibers form ordinary and/or en passant synapses with neurons in the lobi where AVT fibers were localized. As was discussed in our previous paper [4], beaded or varicose ir-A VT fibers traveling through the white matter may form vesicular or varicose ir-A VT fibers located in the cerebral cortex, the preoptic area, the optic tectum and the central gray. In the limbic cortex, a considerable number of ir-A VT fibers were found in the septal nuclei, and also in the nucleus of the diagonal band of Broca. These regions contain the majority of ir-LHRH neurons in the toad brain [4, 10]. A similar projection of vasopressin fibers was found in the organum vasculosum laminae terminalis in the mammalian brain where many ir-LHRH perikarya are localized [12]. In the eel, vasotocin fibers were found in the preoptic region (Fujiwara et al., unpublished). Meanwhile, ir-LHRH fibers project to the VMH in the toad brain [20]. These observations suggest that the LHRH-ergic and vasotocinergic neurosecretory systems are mutually connected, and that they interact on each other for controlling sexual behavior.

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REFERENCES