INTRODUCTION
Dopamine is one of catecholamines produced in the brain by hypothalamic aminergic nuclei NRL (nucleus recessus lateralis) and NRP (nucleus recessus posterioris). It inhibits gonadotropin LH secretion by activating dopamine D2 receptors found on gonadotropic cells of the pituitary gland (Peter et al. 1986, Omeljaniuk et al. 1987). The inhibiting effect of dopamine is not limited to the pituitary gland, because studies with crucian carp proved its inhibitory effect on both LHRH synthesis and secretion (Yu and Peter 1990). This hypothesis was also confirmed by in vitro studies (Yu et al. 1991, Yu and Peter 1992). The release of LHRH by the dopamine D2 receptor is inhibited probably via neuronal projections of neighbouring dopamine fibres and LHRH fibres (Yu and Peter 1992).

Melatonin, whose concentration in the pineal gland and in the peripheral blood is highest at night and lowest during the day (Reiter 1991a), serves as an indicator of a biochemical biological clock. Melatonin synthesis and secretion, which occurs only during the night, delivers to the central nervous system an endocrine signal proportional to the length of the night (Reiter 1991b). This hormone regulates various physiological and neuroendocrinological processes that occur rhythmically, and stimulates or inhibits endocrine activity of various body glands.

This is particularly important for seasonally breeding species, in which photoperiod is the main environmental cue informing the pineal gland of annual changes in the day length. At the same time, this gland acts through its hormone melatonin on the hypothalamic-pituitary-gonadal axis by synchronizing animals with

Background. Melatonin regulates various physiological and neuroendocrinological processes that occur rhythmically, and stimulates or inhibits endocrine activity of various body glands. This acts on the hypothalamic-pituitary-gonadal axis by synchronizing animals with their reproductive cycles. The proximity of melatonin receptors and dopamine and gonadotropin production sites has led to a hypothesis that dopamine may be a link between melatonin and hypothalamic LHRH. Melatonin may have an indirect influence on animal reproduction through dopaminergic structures of the hypothalamus, but the mechanism involved remains unknown, also in fish, for this reason, the present experiment was conducted. The aim of the study was to determine the effect of melatonin on dopamine release from hypothalamic cells of mature female carp in vitro.

Material and Methods. Hypothalami were perifused with a mineral medium containing melatonin (group 1), in the presence of implanted pineal glands (group 2), and with a pure mineral medium (control). Perfusion was 180 min long and samples of the effluent perifusate were collected at 15-minute intervals. Dopamine concentration in the medium was analysed radioenzymatically. The experiment was carried out in the summer during spawning and in the winter during regression.

Results. The results indicate that melatonin inhibits the release of dopamine from hypothalamic cells. This effect was only noticeable in the experiment conducted during the spawning period.

Conclusion. The present findings show that melatonin may have a role in the hypothalamic control of hypophyseal activity during the spawning period of carp.

Key words: pineal gland, melatonin, dopamine, perifusion, carp, fish
their reproductive cycles (Reiter 1991a, b, Zachmann et al. 1991).

In looking for a link between the pineal gland and the hypothalamus, Zisapel and Laudon (1983) and Zisapel et al. (1985) showed that melatonin is able to inhibit dopamine secretion from the hypothalamic cells of the rat. The results of other studies, which showed that even picomolar concentrations of melatonin can inhibit dopamine release from the retina of birds and mammals (Cardinali et al. 1979, Dubocovich 1983, 1985), seemed to support this theory. Also in humans, the high nocturnal level of melatonin reduces dopaminergic activity in the hypothalamus (Rao and Mager 1987). In the brain of mammals, melatonin receptors are most abundant in the hypothalamus and in the anterior lobe of the pituitary gland, and much less abundant in the other parts of the brain (Weaver et al. 1991, Stankov et al. 1993). It is therefore conjectured that the hypothalamus acts as a functional link between melatonin and the endocrine system (Maywood et al. 1996). In the hypothalamus, the highest concentration of melatonin receptors is found in the medial eminence, next to the arcuate nucleus (Weaver et al. 1989). In mammals, this area also contains endings of axons from LHRH-producing cells and neurons of the tubero-infundibular system containing dopamine. Dopamine, by inhibiting LHRH release to the capillaries of the pituitary portal circulation (Wuttke et al. 1971), controls LH and FSH secretion from the pituitary gland of mammals (Gallo 1980, 1981). The proximity of melatonin receptors and dopamine and gonadotrophin production sites has led to a hypothesis that dopamine may be a link between melatonin and hypothalamic LHRH.

The studies with carp have revealed that nocturnal injections of melatonin (dark phase) into mature female carp increase the level of maturation gonadotropin (LH) in blood (Breton et al. 1993). It was also demonstrated that melatonin modulates the activity of the hypothalamic dopaminergic system rather that directly affecting gonads or the pineal gland (Popek 1991, Popek et al. 1994a, Popek and Epler 1999).

The above body of evidence suggests that melatonin may have an indirect influence on animal reproduction through dopaminergic structures of the hypothalamus, but the mechanism involved remains unknown, also in fish. Therefore, the aim of the present experiment was to investigate the effects of the pineal gland and melatonin on dopamine secretion from the hypothalamus not controlled by the body, under in vitro conditions.

The effect of melatonin on the activity of hypothalamic amineergic nuclei is seasonal (Popek 1991, Popek and Epler 1999), as is variation in the rhythm of catecholamine release in the hypothalamus (Popek et al. 1994a). For this reason, the present experiment was conducted in two seasons of the year: in the spawning season of carp in June, and during gonadal regression in December. This experimental design allowed us to determine if the effect of melatonin on dopamine release from hypothalamic amineergic nuclei is season dependent.

MATERIALS AND METHODS
A total of 30 mature female common carp, Cyprinus carpio L., were used. The experiments were carried out twice a year.

Prior to each experiment, fish were decapitated and their hypothalami and pineal glands were removed as rapidly as possible (within 1 minute on average) and placed on ice. Hypothalami (with an average unit weight of 220 mg) were cut with a scalpel into 3-mm-thick slices and were individually placed on a Biogel in 15 perfusion columns of 5 mL volume. Aerated mineral medium in the form of Cortland buffered salt solution was pumped through the columns (Jalabert et al. 1973). The 15-channel peristaltic pump (Zalimp – PP 1B–05) used made it possible to maintain a continuous controlled flow (8 mL per hour) of the perfusion medium. The set contained also a luxmeter sensor placed next to the perfusion columns, whereby the light regime during the experiment could be controlled. Two 11-W Lival halogen lamps provided uniform illumination of the set, producing light of constant 2000 lx intensity (Fig. 1). The set was wrapped in aluminium foil and the light was turned off to ensure complete darkness (0 lx).

Perfusion was performed in three groups (each having 5 perfusion columns):

Group 1, with hypothalami perifused in a medium containing melatonin at a concentration of 300 pg · mL⁻¹ (MT); Group 2, with hypothalami into which three pineal glands were implanted (P); Group 3, with hypothalami perifused in a pure medium, constituting control group (C).

Melatonin (Sigma) was predissolved in 5 µL of 96% ethanol and then in perfusion medium. Temperature of the medium was 22°C.

Perfusion was 180 minutes long. The columns were illuminated (2000 lx) during the first 60 minutes of the experiment and kept in complete darkness for the next 120 minutes of perfusion. Samples of the effluent perfusate were taken using an automatic collector at 15-minute intervals, frozen at –60°C, and stored until the level of dopamine was determined.

Summer period (June, water temperature 22°C, L : D = 16 : 8). Fifteen fish weighing an average of 2.4 kg (±0.16) were investigated. Fish before the experiment, tissues in perfusion columns, and the perfusion medium were kept at 22°C throughout.

Winter period (December, water temperature 5°C, L : D = 8 : 16). Fifteen fish weighing an average of 2.8 kg (±0.24) were investigated. Division into groups and the perfusion method were the same as in the summer experiment. Only temperatures of water in the pond in which fish were kept prior to the experiment, tissues in columns and the perfusion medium were lower (5°C).

Samples of the medium were analysed radioenzymatically to determine the level of dopamine (Johnson et al. 1980). The results were analysed using one-way analysis of variance and STATISTICA procedures were used to determine statistical differences between the groups.
RESULTS

Summer period. During the summer, average dopamine concentration in the perfusion medium in group 1, in which hypothalami were perifused with a medium containing melatonin (MT), ranged from 0.431 (±0.06) to 0.983 (±0.13) pmol · mL⁻¹. The concentration of this hormone was the highest at the beginning of perfusion and decreased over the next 45 minutes to reach 0.522 (±0.06) pmol · mL⁻¹ at 60 minutes. From that time to the end of the experiment, dopamine concentration in the medium continued to be low, ranging from 0.711 (±0.12) to 0.573 (±0.05) pmol · mL⁻¹.

In group 2, in which pineal glands were implanted into hypothalami (P), average dopamine level ranged from 0.53 (±0.03) to 1.188 (±0.25) pmol · mL⁻¹. The highest concentration of this hormone in the medium was found at 30 minutes of perfusion. From 60 minutes of perfusion (perfusion in darkness), dopamine concentration in the medium gradually decreased to reach the lowest value at 135 minutes of the experiment, and was significantly lower ($P < 0.05$) than during the first hour of perfusion.

In the control group (C), average dopamine concentration in the perfusion medium was 0.616 (±0.07) at the beginning, 0.562 (±0.08) pmol · mL⁻¹ at the end, and 1.529 (±0.09) pmol · mL⁻¹ at 90 minutes of the experiment.

Statistical analysis showed that dopamine level at 90 minutes of perfusion in groups 1 (MT) and 2 (P) was highly significantly lower ($P < 0.01$) than in the control group (C). Significant differences ($P < 0.05$) were also noticed at 30, 60, 75, 105, 120, 135, 150, and 165 minutes of perfusion in group 1.

The course of changes in dopamine concentration in the perfusion medium flowing through the columns with hypothalami of mature female carp during the summer is given according to groups in Fig. 2.

Winter period. Average DA concentration at the start of the experiment was similar in all the groups, ranging from 0.263 (±0.02) to 0.367 (±0.03) pmol · mL⁻¹. In the next samples taken during the next minutes of perfusion, DA concentration increased gradually in all the groups and at 105 minutes it reached 0.62 (±0.07) pmol · mL⁻¹ in group 1, where the medium was supplemented with melatonin, 0.469 (±0.05) in group 2, in which hypothalami were perifused in the presence of pineal glands (P), and 0.72 (±0.08) pmol · mL⁻¹ in the control group (C).

Starting from 120 minutes of perfusion, DA concentration in the medium in all the groups slightly decreased, ranging at 180 minutes of perfusion from 0.38 (±0.04) to 0.51 (±0.06) pmol · mL⁻¹.

Statistical analysis did not show any highly significant ($P < 0.01$) differences in dopamine concentration in the medium between the groups. Significant differences ($P < 0.05$) were noticed between summer and winter periods in groups 2 (P) and 3 (C).

The course of changes in dopamine concentration in the perfusion medium flowing through the columns with hypothalami of mature female carp during the winter is shown according to group in Fig. 3.

DISCUSSION

The way in which melatonin affects reproductive processes in animals has puzzled scientists for a long time. Many hypotheses were put forward, some indicating that
Melatonin may modulate the activity of the hypothalamic-pituitary-gonadal axis (Rollag et al. 1978, Lincoln and Short 1980, Arendt et al. 1981, Kennaway et al. 1983, Karsch et al. 1986). More accurate indications were provided by Zisapel and Laudon (1983), Zisapel et al. (1985) and Rao and Mager (1987), who proved that melatonin can inhibit dopamine secretion from hypothalamic cells in the rat. The way in which melatonin affects reproductive.
processes has also been explored in fish. It was shown that melatonin affects changes in the circadian rhythm of LH release from the pituitary gland (Popek 1994b) but in an indirect way, because melatonin shows no influence on pituitary gonadotropes in carp (Popek et al. 2000). Other studies have excluded its direct effect on gonads and oocyte maturation (Popek et al. 1996), although it was shown that melatonin affects changes in the seasonal release of estradiol (Popek et al. 1997a) and it was even suggested that through steroids, it can affect the reproductive processes. However, these and other literature data fail to demonstrate the exact mechanism of melatonin influence on the hypothalamic-pituitary-gonadal axis in fish.

Considering the seasonality of the reproductive cycle in carp and the evident contribution of melatonin to circadian and seasonal synchronization of animal reproduction, the present experiment was performed in two diametrically opposed seasons of the year: in the summer, when sexual activity of the carp is at its highest, and in winter.

In in vivo experiments, where melatonin is injected into the brain of live fish, the interaction of other neurohormones, systems or feedbacks in the body cannot be ruled out. The observed effects of such experiments in the form of melatonin-induced changes in the dopaminergic activity of the hypothalamus can vary. However, to ensure that melatonin has a direct effect on dopamine secretion from the hypothalamus, in vitro studies were conducted outside the organism. The object of the studies were dissected hypothalami placed in perfusion columns filled with a flowing medium (Fig. 1). In the control group, hypothalami were only perfused with a physiological salt solution (pure medium). In the experimental groups, synthetic melatonin (MT) was dissolved in the medium and, for the sake of comparison; live pineal glands taken during the same time from the fish were implanted into the hypothalami (P). During perfusion, medium temperature was constant in keeping with temperature of the water in which fish were kept, and varying photoperiod (light–dark) was applied. We used the observations of other authors (Gern and Greenhouse 1988, Kezuka et al. 1989), who showed that thanks to photoreceptive cells, fish pineal gland also reacts in vitro to changes in light intensity by changes in melatonin secretion, in exactly the same manner as in physiological conditions.

The results obtained in the summer have fully confirmed our hypotheses because in the hypothalamic-pituitary-gonadal axis during spawning period of carp. Probably this effect is stimulatory and involves inhibiting the hypothalamic-pituitary-gonadal axis during spawning (Popek 1991, Popek et al. 1991, 1997a, b, Popek and Epler 1999).

In conclusion, it is hypothesized that melatonin can be one of the major factors controlling LH release from the pituitary gland during the spawning period of carp. Probably this effect is stimulatory and involves inhibiting the hypothalamic secretion of dopamine.

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