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Short communication

## Gender- and age-specific impairment of rat performance in the Morris water maze following prenatal exposure to an MRI magnetic field

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## Abstract

We examined the effects of prenatal exposure (40 min/day, gestation days 12-18) of rats to a magnetic resonance imaging (MRI) magnetic field (MF) on their performance in the Morris water maze. At 2 months of age, female rats showed impaired performance. The animals spent longer time swimming and used inefficient strategies. However, no significant effects on maze performance were observed at 1 and 5 months of age. No evident maze performance deficit was detected in male rats prenatally exposed to the magnetic field. Thus, we conclude that prenatal exposure to MRI magnetic field induces cognitive/behavioral deficits in female rats at a specific age. © 2003 Elsevier B.V. All rights reserved.

Theme: Learning and memory Topic: Behavior

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There is increasing public concern regarding the safety of exposure to magnetic fields (MFs), including those used for magnetic resonance imaging (MRI) applications. Several lines of scientific research have clearly disclosed a number of potential harmful effects of MFs. For example, exposure to MFs causes deleterious effects on spermatogenesis and embryogenesis in animals, and abortion in both humans and animals [18].

The central nervous system (CNS) is particularly susceptible to MFs, which may markedly affect neural electrophysiological and biochemical activities [16,23]. Exposure to MFs can increase the risk of neurodegenerative diseases in humans [19] and produce deficits in attention, perception and spatial learning in rats [12]. Furthermore, the developing CNS exhibits even higher sensitivity to MFs [6]. Prenatal or perinatal exposure to MFs decreases the density of neurons in the medial preoptic nucleus, affects some sexually dimorphic structures, and impairs scent marking

and inter-male fighting behaviors during adulthood [17,20]. Therefore, elucidation of the physiological and behavioral consequences of exposure of the developing CNS to MFs requires considerable attention.

MRI is widely used to study fetal development [7,22] and assess brain injuries [1], due to its high capacity of visualizing the brain. The effects of prenatal exposure to MRI on cognitive performance at different stages of postnatal development have not been described in detail. In the present study, we investigate the effects of periodic exposure of pregnant rats to MRI fields on the offspring's performance in the Morris water maze at different postnatal ages.

Fifteen healthy pregnant female Sprague-Dawley rats were used in the present study. Animals were exposed either to MFs or sham conditions between days 12 and 18 of gestation. Eight of the pregnant rats were randomly selected and exposed to the MRI field at 7:00 p.m. for 7 days for 40 min a day. The other seven adult rats were exposed to a sham control field under similar conditions. The MRI field was a 0.35 T 1 kW power impulse magnetic field produced by a

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Diasonics MRT-35A system (Comp, USA; spin echo  $T_1$ : TR 500 ms/TE 15 ms, slice thinness 10 mm, Nex 4;  $T_2$ : TR 2000 ms/TE 120 ms, Slice thinness 10 mm, Nex 2).

After parturition, each of the 15 litters was housed in a standard plastic cage  $(46 \times 31.5 \times 20 \text{ cm})$  with food and water available ad libitum for the maternal rats. The room temperature was maintained at  $21 \pm 1$  °C. The light–dark cycle was 12:12 h (light on at 7:00 a.m.). On the second postnatal day, the number of pups in each litter was randomly reduced to 8. At 21–days of age, offspring were separated from their respective maternal rats and assigned into four general groups: (1) MRI male, (2) MRI female, (3) control male, and (4) control female. Within each group, rats were further randomly divided into three testing-age sets (designated according to their ages when tests were conducted) of 1-, 2-, and 5-month groups. In total, 114 offspring rats were assigned into 12 (4 × 3) groups, and a maximum of two

males/females per litter were distributed to each group. The MRI 5-month male group included 8 rats and the control 5-month male group comprised of 6 rats. All other groups contained 10 rats each.

Behavioral tests were conducted using a Morris water maze (120 cm in diameter and 55 cm in height) in a dim room illuminated by a carefully positioned lamp whose reflections from pool water were not perceived by the video camera. The pool was filled with water to a depth of 41 cm, with a temperature of  $23 \pm 1$  °C. The position and orientation of the pool in the testing room remained unchanged throughout the study, and thus, both geometric and landmark cues were maintained constant [21]. The pool was divided into four quadrants. A round transparent platform (8 cm in diameter) was placed at the center of a designated quadrant and submerged 1 cm beneath the water surface. The distance from the platform center to the pool edge was 35 cm.



Fig. 1. Average escape latency during the nine testing sessions for sham-exposed (open squares) and magnetic field-exposed (filled squares) offspring. The error bars indicate the standard errors of the mean.

Tests on each rat were conducted for 5 successive days. Two test sessions were performed in each of the first 4 days, and one session was conducted on the fifth (last) day. Each session comprised four trials, with an inter-trial interval of 60 s. The inter-session interval within a single day was 2 h.

In each trial, the rat was gently placed into the pool at the middle site of the circular edge in a randomly selected quadrant, with the head facing the pool edge. The rat was allowed to find the hidden platform within 120 s and the minimal time for staying on the platform was 3 s. If a rat failed to stay on the platform for at least 3 s within the 120 s of testing time, it was placed on the platform for 10 s by the experimenter, and its performance score (latency) was marked as "120 s". After each trial, rats were removed from the pool, dried with a towel, and placed in a heated and dry cage. Tests were conducted between 8:00 and 12:00 a.m., and between 2:00 and 6:00 p.m.

The final test was performed on the afternoon of the fifth training day and comprised only a single trial lasting 120 s. The platform was removed from the pool, and each rat entered the pool from the quadrant opposite to the one containing the platform (the "platform" quadrant) in previous testing sessions.

Rat positions in the pool were automatically tracked by a commercial video/computer system (Beijing Logon Science and Technology) with a video camera mounted above the center of the pool. The latency for finding the platform and the following four swimming strategies were recorded: (1) marginal: swimming along the pool edge, (2) random: swimming randomly, (3) tendency: swimming around towards the platform area, and (4) straight: swimming straight-



Fig. 2. Average incidence of each of the four strategies used by the sham-exposed (open bars) and magnetic field-exposed (solid bars) offspring. The error bars indicate the standard errors of the mean. \*Significant differences between control and MF groups (p < 0.05).

ly towards the platform. In the last test, the time that the rat spent within the "platform" quadrant was additionally recorded.

Repeated-measure analysis of variance was applied across sessions. Treatment, age, and sex were the three between-subject factors. Swimming strategies were analyzed with the Mann–Whitney non-parametric test. The time spent on the "platform" quadrant in the last test for each rat was analyzed with one-way analysis of variance.

Latencies across sessions for all the animal groups are shown in Fig. 1. All rats improved in performance during the tests, as indicated by the progressive reduction in escaping latencies over the sessions ( $F_{8, 816}=244.84$ , p < 0.05). For the three between-subject factors (treatment, age, and sex), the interaction of treatment × age was significant ( $F_{2,102}=7.501$ ; p < 0.05). However, interactions of treatment × sex, sex × age, and treatment × age × sex were not significant.

To analyze the treatment main effects, we examined the differences between control and treatment groups for each age and sex combination. Interactions between testing sessions and treatment were not significant. The treatment main effect was significant for the 2-month female group ( $F_{1,18}$ =17.35, p<0.05), but not for the other groups.

Examination of the strategy profiles used by each animal group revealed that the 2-month-old MRI female rats used marginal strategy more frequently but less tendency and straight strategies than the 2-month-old control female rats. No significant treatment effects were observed in the other groups (Fig. 2).

In the last test, no differences in time spent in the "platform" quadrant were observed between groups (Table 1).

The present data show that prenatal exposure to the MRI field resulted in impairment of the Morris water maze performance in 2-month-old female rats only. These females used inefficient strategies and spent longer times to find the platform, compared to their control counterparts. No significant effects were observed in the male

Table 1								
Average time	spent in	the platform	quadrant in	the	Morris	maze	probe	tests

Age	Sex	Group	N	Time (s)
1 month	М	CON	10	$38.51 \pm 2.75$
		MRI	10	$37.10 \pm 1.94$
	F	CON	10	$39.36 \pm 3.52$
		MRI	10	$38.57 \pm 2.73$
2 month	М	CON	10	$35.24\pm3.55$
		MRI	10	$36.96 \pm 3.34$
	F	CON	10	$34.39\pm5.19$
		MRI	10	$40.13 \pm 3.96$
5 month	М	CON	6	$37.65 \pm 3.78$
		MRI	8	$43.93 \pm 3.06$
	F	CON	10	$39.88 \pm 2.02$
		MRI	10	$40.88\pm3.09$

Means  $\pm$  S.E.M. are presented for each group.

groups. These results are similar to previous reports that prenatal exposure to alcohol, toluene, or lead caused poor performance in the water maze only in female offspring of rats [4,10,11], indicating an important sexual difference in response to harmful prenatal environments. It is proposed that males generally use a single type of cues (geometric) in spatial learning, while females depend on multiple cue types [21]. Thus, strategies used by females are considered more complex than the one used by males and therefore may demand more on various signaling processes, causing the females' water maze performance more susceptible to prenatal harmful influences. The present finding, that prenatal MRI exposure has a particularly marked effect on the cognitive performance of 2-monthold female offspring, is in agreement with this theory. This disruptive effect is not apparent in younger or adult females. In view of the previous findings that perinatal exposure to MFs affects sexually dimorphic structures [17], it is of interest to determine whether this sex/agespecific effect is related to the brain sexual differentiation. A number of studies have showed that the sex difference in the midsagittal area of the adult rat corpus callosum is mediated, in part, by gonadal steroids in early development, with the sensitive period of hormone action in the female between 25 and 78 postnatal days [3,14]. Additionally, sex differences are evident in spatial learning during early development [5]. The prolonged sexual maturation in females may lead to higher sensitivity to prenatal MFs exposure, and the consequences are more obvious when females have just become sexually mature. Accordingly, the influence of prenatal exposure to MFs on sexual hormonal systems, such as those related to estrogen, should be thoroughly investigated. It is established that estrogen is closely associated with the neural plasticity that occurs during learning [15].

Interestingly, prenatal exposure to MFs is accompanied by an increase in dopamine levels in the offspring's striatum between postnatal weeks 4 and 12 [13]. Estrogen modulates dopamine activity differentially in the striatum of females and males [2], and dopamine activity in the striatum is closely related to the water maze performance [8,9]. Thus, impaired water maze performance in the 2month-old female offspring suggests an altered estrogendopamine interaction pattern in the striatum, following prenatal exposure to the MRI field.

On the other hand, prenatal exposure to the MRI field does not affect the birth weight and litter size of the offspring (data not shown). In our experiments, prenatal exposure was not too severe, and the observed effects in the 2-month female group were not due to a change in birth weight.

The results of the present study strongly call for serious caution in regard to prolonged exposure of pregnant humans to medical MFs. The effects of prenatal exposure to MFs on the interplay between brain sexual differentiation, sexual hormonal modulation, and dopamine activity require further investigation.

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## References

- S. Akoka, P. Descamps, C. Genberg, F. Franconi, B. Arbeille, R. Laurini, A. Locatelli, L.D. Platt, P. Arbeille, Cerebral MRI on fetuses submitted to repeated cocaine administration during the gestation: an ovine model, Eur. J. Obstet. Gynecol. Reprod. Bio. 85 (1999) 185–190.
- [2] J.B. Becker, Gender differences in dopaminergic function in striatum and nucleus accumbens, Pharmacol. Biochem. Behav. 64 (1999) 803-812.
- [3] H.A. Bimonte, R.H. Fitch, V.H. Denenberg, Neonatal estrogen blockade prevents normal callosal responsiveness to estradiol in adulthood, Dev. Brain Res. 122 (2000) 149–155.
- [4] B.A. Blanchard, E.P. Riley, J.H. Hannigan, Deficits on a spatial navigation task following prenatal exposure to ethanol, Neurotoxicol. Teratol. 9 (1987) 253–258.
- [5] J.M. Cimadevilla, H. González-Pardo, L. López, F. Díaz, E.G. Cueto, L.M. García-Moreno, J.L. Arias, Sex-related differences in spatial learning during the early postnatal development of the rat, Behav. Processes 46 (1999) 159–171.
- [6] L.G. Cockerham, G.D. Prell, Prenatal radiation risk to the brain, Neurotoxicology 10 (1989) 467–474.
- [7] N. Girard, C. Raybaud, D. Gambarelli, D. Figarella Branger, Fetal brain MR imaging, Magn. Reson. Imaging Clin. N. Am. 9 (2001) 19–56.
- [8] J. Hagan, J. Alpert, R. Morris, S. Iversen, Effects of neostriatal and mesocorticolimbic dopamine depletions on learning in a water maze, Behav. Brain Res. 5 (1982) 103.
- [9] C. Heim, W. Kolasiewicz, T. Kurz, K.H. Sontag, Behavioral alterations after unilateral 6-hydroxydopamine lesions of the striatum, effect of alpha-tocopherol, Pol. J. Pharmacol. 53 (2001) 435–448.
- [10] K.S. Hougaard, U. Hass, S.P. Lund, L. Simonsen, Effects of prenatal exposure to Toluene on postnatal development and behavior in rats, Neurotoxicol. Teratol. 21 (1999) 241–250.
- [11] D.A. Jett, A.C. Kuhlmann, S.J. Farmer, T.R. Guilarte, Age-dependent

effects of developmental lead exposure on performance in the Morris water maze, Pharmacol. Biochem. Behav. 57 (1997) 271–279.

- [12] H. Lai, M.A. Carino, I. Ushijima, Acute exposure to a 60 Hz magnetic field affects rats' water-maze performance, Bioelectromagnetics 19 (1998) 117–122.
- [13] B.C. Lee, G.Y. Bing, W.K. Jhoo, J.M. Yoon, K.S. Kang, E.J. Shin, W.K. Kim, K.K. Ho, H.C. Kim, Prenatal exposure to magnetic field increases dopamine levels in the striatum of offspring, Clin. Exp. Pharmacol. Physiol. 28 (2001) 884–886.
- [14] C.M. Mack, R.H. Fitch, L.A. Hyde, A.J. Seaman, H.A. Bimonte, W. Wei, V.H. Denenberg, Lack of activational influence of ovarian hormones on the size of the female rat's *corpus callosum*, Physiol. Behav. 60 (1996) 431–434.
- [15] B. McEwen, Estrogen actions throughout the brain, Recent Prog. Horm. Res. 57 (2002) 357–384.
- [16] J. Miyakoshi, T. Tsukada, S. Tachiiri, S. Bandoh, K. Yamaguchi, H. Takebe, Enhanced NOR-1 gene expression by exposure of Chinese hamster cells to high-density 50 Hz magnetic fields, Mol. Cell. Biochem. 181 (1998) 191–195.
- [17] S. Muligan, M.A. Persinger, Perinatal exposures to rotating magnetic fields 'demasculinize' neuronal density in the medial preoptic nucleus of male rats, Neurosci. Lett. 253 (1998) 29–32.
- [18] V.R. Narra, R.W. Howell, S.M. Goddu, D.V. Rao, Effects of a 1.5tesla static magnetic field on spermatogenesis and embryogenesis in mice, Invest. Radiol. 31 (1996) 586–590.
- [19] C.W. Noonan, J.S. Reif, M. Yost, J. Touchstone, Occupational exposure to magnetic fields in case-referent studies of neurodegenerative diseases, Scand. J. Work, Environ. Health 28 (2002) 42–48.
- [20] M.A. Persinger, S. Mulligan, Decreased density of neurons in the medial preoptic nucleus and increased testicular weights for rats exposed perinatally to an 0.5 Hz rotating magnetic field, Int. J. Neurosci. 108 (2001) 99–107.
- [21] R.L. Roof, D.G. Stein, Gender differences in Morris water maze performance depend on task parameters, Physiol. Behav. 68 (1999) 81–86.
- [22] I. Trop, D. Levine, Normal fetal anatomy as visualized with fast magnetic resonance imaging, Top. Magn. Reson. Imaging 12 (2001) 3-17.
- [23] A.B. Uzdensky, O.Y. Kutko, Effect of weak extremely low frequency magnetic field on isolated crayfish stretch receptor neuron: Nonlinear dependence on field amplitude and frequency, Electro- Magnetobiol. 16 (1997) 267–279.