# Melatonin And Magnetic Fields

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#### Abstract

There is public health concern raised by epidemiological studies indicating that extremely low frequency electric and magnetic fields generated by electric power distribution systems in the environment may be hazardous. Possible carcinogenic effects of magnetic field in combination with suggested oncostatic action of melatonin lead to the hypothesis that the primary effects of electric and magnetic fields exposure is a reduction of melatonin synthesis which, in turn, may promote cancer growth.

In this review the data on the influence of magnetic fields on melatonin synthesis, both in the animals and humans, are briefly presented and discussed.

#### Introduction

Extremely low frequency electric (ELF-EF) and magnetic fields (ELF-MF), e.g. generated by highvoltage transmission lines and household appliances, are present worldwide and receive increasing attention because of their potential consequences for human health [1-3], especially associated with increased risk for cancer and childhood leukemia [2]. Additionally some attention has also been paid to other possible health hazards, such as interference with cardiac pacemakers [4], Alzheimer's disease [5], and adverse pregnancy outcome [6]. Working Group organized by National Institute of Environmental Health Services concluded in the report published in 1998, on the basis of almost 900 publications, that "ELF-EMF are possibly carcinogenic to humans". This conclusion is based on limited evidence that residential exposure to ELF-MF is carcinogenic to children it terms of childhood leukemia, and occupational exposure to ELF-MF is carcinogenic to humans in terms of chronic lymphocytic leukemia [4]. It has been also concluded that there is inadequate evidence for an association between occupational exposure to EFL-MF and risk for other cancer [4]. Moreover, the same report states: "None of the evidence for adverse health effects seen after exposure to ELF EMF achieved a degree of evidence exceeding 'inadequate' (for humans) or 'weak" (for experimental animals)" [3]. In humans it concerns adverse birth outcomes, reproductive effects, Alzheimer's disease, amyotrophic lateral sclerosis, suicide or depression, cardiovascular disease [3].

Possible carcinogenic effects of EMF in combination with suggested oncostatic action of melatonin [see 7, 8] lead to the hypothesis that the primary effects of EMF exposure is a reduction of melatonin synthesis which, in turn, may promote cancer growth [9].

In this review the data on the influence of EMF on melatonin synthesis, both in the animals and humans, are briefly presented and discussed.

#### **Animal studies**

There is substantial evidence that exposure to ELF-MF may alter melatonin secretion in animals. The results of the studies on the influence of ELF-MF on melatonin concentrations in the animals are presented in Table 1.

It can be seen that in majority of studies ELF-MF decreased melatonin secretion, especially in rats and hamsters. However, large variations exist when studies are repeated [34, 35, 39]. Among other explanations [48], one reason may be the age of the animals since young rats seem to be more sensitive against magnetic fields than old ones [49]. Recently, an investigation on isolated hamster pineal organs has shown variable results when the experiments were repeated under identical conditions at 16 2/3 and 50 Hz. However, exposure was found to suppress melatonin synthesis highly significantly when the results were pooled [42].

### **Human studies**

In contrast to animal studies, the data on the influence of magnetic fields on human melatonin concentrations are scarce and contradictory. Conversely to the animal studies, although ELF-MF-induced suppression on nocturnal melatonin secretion in humans has been reported in occupational and residential studies [50-55] in the majority of laboratory-based exposure studies ELF-MF no change in melatonin or 6-hydroxymelatonin concentrations has been observed (Table 2).

It should be stressed that various parameters of magnetic fields have been employed in the experiments. Therefore, discrepancies in the results may depend on different experimental paradigms, including differences in certain characteristics of the applied magnetic fields, such as field intensity, frequency, duration of exposure, timing of exposure, applied vector, etc. The results of our studies seem to support the hypothesis that the response of the human pineal to magnetic fields may depend on the field parameters because chronic exposure to 2.9 mT, 40 Hz magnetic field caused a significant decrease in nocturnal melatonin concentrations [63], whereas chronic exposure to 25-80 µT, 200 Hz mag-

Table 1. Effects of magnetic field	s on melatonin concentrat	tions in the animals
Exposure parameters	Exposure duration	Effect

Exposure parameters Rat	Exposure duration	Effect	[Ref.]
	15 min		
52–123 μT	5 h	<b>↓</b> (p) <b>↓</b> (s)	[10]
50 T	24 h	⊗ (p)	[44]
50 µT	<u>2 h</u> 30 min		[11]
50–100 μT	30 min	(p) = signed $\otimes$ (p) = blind	[12]
35 µT	30 min	(p) btillu	[13]
<u>2.5 μ</u> Τ	30 min	(p) ⊗ (p)	[14]
30 µT	30 min	● (p)	[15]
40 µT	60 min	⊗ (p, s)	[16]
80 μT (inter, auto)	60 min	<b>↓</b> (p)	[17]
80 µT (inter, man)		⊗ (p)	[1/]
44 μT, 33.7 Hz	2.5 h ( <i>in vitro</i> )	<b>↓</b> (p, m)	[18]
5.2 mT, 50 Hz	30 min for 15 days	<b>↓</b> (s)	[19]
<u>44 μT (inter)</u>	60 min ( <i>in vitro</i> )	<b>↓</b> (p) ⊗ (m)	[20]
<u>1, 5, 50, 250 μT, 50 Hz (CP)</u>	6 weeks	<b>↓</b> (p, pl)	[21]
44 μT (inter, man, not inverted 44 μT (inter, man, inverted)	l) 30 min, 60 min 30 min, 60 min	⊗ (p) ♦ (p)	[22]
0.03 mT, 10 Hz	10-30 min	$\otimes$ (s, night) $\clubsuit$ (s, day)	[23]
	•	(p) - nighttime exposure	
25 μT (pulsed, static)	•	$\otimes$ (p) – daytime exposure	[24]
1 µT, 50 Hz (CP)	6 weeks		[25]
1 μT, 50 Hz (HP)		⊗ (p, pl)	<u> </u>
1 µT, 50 Hz (VP)	6 weeks	⊗ (p, pl)	[26]
0.3–1 µT, 50 Hz	8–9 weeks	<b>↓</b> (s)	[27]
100 μT, 50 Hz	12 h	<b>↓</b> (s)	[28]
10–100 µT, 50 Hz	18 h for 30 days		[20]
5, 500 µT, 50 Hz	24 h	⊗ (u)	[29]
50 μT, 50 Hz	9 or 12 weeks	⊗ (s)	[30]
10 μT, 50 Hz	91 days	⊗ (p) ♥ (s)	[31]
1, 100 μT, 50 Hz	24 h	<b>★</b> (u)	[32]
900 MHz (continuous or pulsed with 217 Hz)	15 to 60 min	⊗ (s)	[33]
100 µT, 50 Hz	1 day to 13 weeks	inconsistent	[34]
50 to 500 µT	15 to 120 min		
(majority 100 µT)	(15 experiments)	inconsistent	[35]
0.06 µT, 0.05 mT	12 h ( <i>in vitro</i> )	<b>↓</b> (46%)	[36]
1 mT, 60 Hz (inter)	1 h	⊗ (u)	
1 mT, 60 Hz	20 h for 10 or 42 days	⊗ (u)	[37]
Golden hamster			
35 µT	30 min	⊗ (p)	[13]
)jungarian hamster			
100 μT, 60 Hz	15 min	<b>↓</b> (p, s)	[38]
100 µT, 60 Hz E	xp. 1–15 min for 9 days (	lp)	[20]
	xp. 1–15 min for 9 days ( xp. 2–15 min for 9 days (		[39]
 10, 100 μT, 60 Hz	xp. 2–15 min for 9 days (		
10, 100 µT, 60 Hz (inter)	15 min, 60 min	⊗ (p, s)	[40]
900 MHz (continuous or pulsed	d 15 min to 60 min	⊗ (s)	[33]
with 217 Hz)		- (-)	[]
300 μT, 50 Hz	24 h for 56 days	⊗ (s)	[41]
86 µT, 16 <sup>2/3</sup> Hz or 50 Hz	8 h (in vitro)	● (m)	[42]
Gerbil			
30 µT	30 min	$\otimes$ (p) – pigmented	[15]
	<b>↓</b> (p	, female) $\otimes$ (p, male) – alb	ino
louse			
0.5–77 (average 2.75) μT, 50 H		(pl)	[43]
	(from conception)	N 7	
'haan		@ (c)	[//]
		⊗ (s)	[44]
<b>Sheep</b> 4 μΤ, 60 Hz	24 h for 8 months	$\bigotimes(c)$	[/[]
4 μT, 60 Hz 3.77 μT, 60 Hz	24 h for 8 months 24 h for 8 months	⊗ (s)	[45]
4 μT, 60 Hz 3.77 μT, 60 Hz Baboon	24 h for 8 months	••	
4 μΤ, 60 Hz 3.77 μΤ, 60 Hz Baboon 50, 100 μΤ, 60 Hz		⊗ (s) ⊗ (s)	[45]
4 μΤ, 60 Hz 3.77 μΤ, 60 Hz <b>Baboon</b>	24 h for 8 months	••	

netic field did not influence melatonin levels [64].

### **Concluding remarks**

Summarizing, it seems that there is substantial evidence that exposure to ELF-MF decreases melatonin secretion in some animals. However, presently there are no convincing data showing a distinct effect of magnetic fields on melatonin secretion in humans, and more studies are needed to identify the specific circumstances under which such effects may occur.

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Table 2. Effects of magnetic fields on serum (SE), plasma (PL), salivary (SA) and urinary (U) melatonin concentrations or 6-hydroxymelatonin sulfate (6-OHMS) urinary concentrations in humans

Exposure parameters	Exposure duration	Outcome	[Ref.]
150 mT (MRI)	40.5 min	no effect (SE)	[56]
0.057 μT, 60 Hz	4 weeks		
0.66 µT, 60 Hz	4 weeks	no effect (6-0HMS)	[57]
0.46 μT, 60 Hz	7 weeks		
1.5 T (MRI)	60 min	no effect (PL)	[58]
10 μT, 50 Hz	9 h (continuous)	no effect (SE, 6-0HMS)	[59]
10 μT, 50 Hz	9 h (intermittent)		
1 μT, 60 Hz	8 h (intermittent)	no effect (PL) (reduction in	[60] n)
20 μT, 60 Hz	8 h (continuous)	men with low baseline melatonin)	
20 μT, 50 Hz	8 h (continuous)	no effect (PL)	[61]
20 µT, 50 Hz	1.5 to 4 h	no effect (PL)	[62]
		(delayed rise)	[02]
2.9 mT, 40 Hz	3 weeks (20 min per day, 5 days a weeks)	decrease (SE)	[63]
25–80 μT, 200 Hz	3 weeks (16 min per day, 5 days per weeks)	no effect (SE)	[64]
2–7 mT	<u> </u>	no effect (6-0HMS)	[65]
28.3 µT, 60 Hz	8 h	no effect (U, 6-0HMS)	[66]
100 µT, 50 Hz continuous or intermittent	30 min	no effect (PL, 6-OHMS)	[67]
0.7–9.1 μT, 50 Hz	overnight for 11 weeks	no effect (6-0HMS)	[68]
GSM-standard: 900 MHz, pulsed with	8 h	no effect (SE)	[69]
217 Hz, pulse width of 577 μs			
GSM-standard: 900 MHz, pulsed with	4 weeks (2 h per day,	no effect (SE)	[70]
217 Hz, pulse width of 576 μs	5 days a week)	no enect (SE)	
DCS-standard: 1800 MHz, pulsed with	4 weeks (2 h per day,	no effect (SE)	[70]
217 Hz, pulse width of 576 μs	5 days a week)	no enece (SE)	[/0]
GSM-standard: 900 MHz, pulsed with	20 randomly allotted	no effect (SA)	[71]
217 Hz, pulse width of 577 μs	4 h sessions		[, 1]
GSM-standard: 900 MHz, pulsed with 217 Hz, pulse width of 576 µs	60 min	no effect (6-0HMS)	[72]

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