

Effect of Melatonin and Tetrapeptide on Gene Expression in Mouse Brain

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A microchip technique was used to study expression of 16,897 clones from a cDNA library in the brain of mice receiving melatonin or tetrapeptide Epithalon (Ala-Glu-Asp-Gly). Expression of 53 transcripts in mouse brain underwent significant changes after treatment with the preparations. Melatonin and Epithalon modified expression of 38 and 22 transcripts, respectively. These preparations produced similar changes in the expression of 6 transcripts. Expression of 1 transcript (Rp119) was inhibited by melatonin, but induced by Epithalon. The target genes are physiologically related to the cell cycle, apoptosis, biosynthesis, processing, and transport of nucleic acids. Comparative study of gene expression in the brain and heart of CBA mice receiving melatonin and Epithalon suggest that these preparations have a tissue-specific biological effect.

Key Words: melatonin; peptide; Epithalon; gene expression; cDNA microchips

The pineal gland hormone melatonin plays an important role in physiological processes. Melatonin has *in vitro* and *in vivo* protective effects on neurons. This hormone protects various subtypes of neurons and whole brain from damages produced by ischemia, γ -radiation, or neurotoxic agents [3,9,11]. Melatonin induces expression of antioxidant enzyme RNA and proteins [3]. Melatonin serves a free radical-trapping agent and is used for the therapy of patients with Alzheimer's disease, Parkinson's disease, and consequences of stroke or exposure of nerve cells to neurotoxins and hyperbaric oxygenation [3]. Melatonin is a major mediator of endocrine function in the pineal gland. It should be emphasized that activity of the pineal gland is realized via secretion of epiphyseal peptides [4]. Epithalamin, a complex peptide bioregulator of the pineal gland, decelerates aging, increases lifespan in drosophilae, mice and rats, and inhibits spontaneous and induced carcinogenesis in rodents [4]. The study

of the amino acid composition in Epithalamin allowed us to synthesize Epithalon (Ala-Glu-Asp-Gly) with a molecular weight of 390.35 Da [4]. This peptide has high geroprotective activity and suppresses growth of spontaneous and induced tumors in mice and rats. Fine mechanisms of the action of short peptides remain unclear. The effects of melatonin and Epithalon on the expression of 15,247 transcripts in the heart of CBA mice were evaluated by means of a microchip technique with cDNA macromatrices [1,2]. The primary study of target genes for melatonin and peptides would elucidate the mechanism for their biological action in tissues. Here we studied the effects of melatonin and Epithalon on expression of 16,897 transcripts in mouse brain.

MATERIALS AND METHODS

Experiments were performed on 30 female CBA mice obtained from the Rappolovo nursery (Russian Academy of Medical Sciences). The animals received *ad libitum* food and water and were kept at a 12:12-h light/dark regimen. At the age of 6 months the mice were divided into 3 groups of 10 specimens each.

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Control animals received subcutaneous injections of 0.1 ml physiological saline. Epithalon was synthesized by E. I. Grigor'ev at the St. Petersburg Institute of Bioregulation and Gerontology. This preparation was dissolved in 0.1 ml physiological saline and injected subcutaneously in a dose of 0.1 µg. Melatonin (20 mg/liter, Sigma) was given *ad libitum* with water for 1 night (18.00-9.00). The mice were decapitated on day 6. The brain was removed, immediately frozen in liquid nitrogen, and stored at -80°C until total RNA isolation.

Experiments were performed with the library of clones NIA 17K (cDNA collection of the US National Institute of Aging, NIA). The reformatted combination of NIA 15K [7] and NIA 7.4K libraries [13] consisted of 16,897 clones (more than 12,000 unique gene products). Clones were placed on 2 nylon membranes marked by letters A and B. The distance between centers of spots was 0.85 mm. The diameter of spots was 0.4 mm. The information about nucleotide sequences of clones is available at the site of the Laboratory of Genetics (US National Institute of Aging, <http://www.igsun.grc.nia.nih.gov/>). Detailed description of the technique to obtain cDNA macromatrices is presented at the site of the Department of Microchips (Johns Hopkins University, US National Institute of Aging; <http://www.daf.jhmi.edu/microarray/index.htm>).

The samples were prepared, treated with isotopes, and hybridized with cDNA macromatrices as described elsewhere [1,2]. Total RNA was isolated by the guanidine isothiocyanate method [1] and used as the substrate (20 µg) to introduce a radioactive isotope of phosphorus ³²P (Amersham) during cDNA synthesis with SuperScript II reverse transcriptase (Life Technologies) on the oligo(dT)₁₂₋₁₈ primer (Amersham). Each group was exposed to 2 independent hybridizations. Experiments were performed with independently synthesized samples. After normalization by total emission, the samples were hybridized with prehybridized membranes of macromatrices using the mixture of MicroHyb (Research Genetics), Cot1 (1 µg/ml, Life Technologies), and Poly(A) (50 µg/ml, Amersham) at 42°C for 18 h. Washed membranes were exposed to a Storage Phosphor screen (Molecular Dynamics) for 5 days and scanned with a Storm 860 automatic radiation detector (resolution 50 µpixels, Amersham). The formatted image was analyzed by means of Array-Pro Analyzer 4.0.0.41 software (Media Cybernetics). Coordinate grids were superimposed on each image to estimate the strength of a signal from individual clones (arb. units). Background signals were locally normalized. The results of hybridization were normalized by the method of Z-transformation. The difference between common logarithms of signal strength and average signal strength was divided by the standard

deviation of the common logarithm of the signal. Relative expression was estimated by comparing the average results of experimental and control hybridization. The difference between Z-transformations of experimental and control groups was divided by the standard deviation to calculate the Z-ratio (Z_{rel}) [1,2]. The coefficient of variation was estimated for each membrane. We calculated the ratio between the standard deviation of the common logarithm of the signal and mean value of the common logarithm of the signal (%). The correlation coefficient was determined for each set of clones (MS Excel XP software).

RESULTS

³²P-labeled samples (2 samples of each group) were synthesized from independently isolated samples of mouse brain total RNA and hybridized with cDNA macromatrices consisting of 16,897 DNA clones (Fig. 1). Screening with 16,897 clones revealed significant expression of 65.7, 62.7, and 59.8% clones in control animals and mice receiving Epithalon and melatonin, respectively (Z -transformations > 0). In these mice the ratio of highly expressed clones was 8.2, 9.4, and 10.5%, respectively (Z -transformations > 1). The coefficients of variation and correlation confirmed high accuracy of hybridization. The correlation coefficients for hybridization of samples from control animals and mice receiving Epithalon and melatonin were 0.995, 0.991, and 0.984, respectively. The average coefficients of variation were 36.0, 33.6, and 33.8%, respectively.

The amount of gene products characterized by significant changes in the level of expression was extremely low (53 clones, 0.31% of the total number, $1.5 < Z_{rel} < 1.5$). However, a considerable number of clones exhibited high level of expression (Fig. 1). In mice receiving melatonin and Epithalon, expression of clones increased (5 and 9 clones, respectively) or decreased (33 and 13 clones, respectively, Table 1). Study of normalized Z_{rel} in animals receiving melatonin and Epithalon showed that maximum changes in the level of expression were +5.02 (RIKEN cDNA 1810049E02 gene) and -5.10 (male germ cell-associated kinase, Mnk3, respectively). Nucleotide sequences of 11 transcripts with modified expression (20.8%) corresponded to the known genes. Six clones did not exhibit similarity to the GenBank genetic information, since study of nucleotide sequences in the mouse genome was uncompleted. Nucleotide sequences of 36 transcripts corresponded to expressed gene fragments (ESTs) that were similar (7 transcripts) or dissimilar to the known genes (22 transcripts). The known genes belonged to various functional classes, including cell systems for signaling and communication (transforming growth factor β 1-induced transcript 4, guanine

nucleotide-releasing factor RAN (Mog1), and ESTs similar to isoform 1 of human caldesmon gene), expression of genes and proteins (ribosomal protein L19 (Rp19), ribonucleic acid-binding protein S1, and interferon regulatory factor 3 (Irf3)) and metabolism (Tousled-like kinase 2 (Tlk2), dihydrolipoamide branched chain transacylase E2 (Dbt), and guanosine monophosphate reductase isologue (Gmpr2)). Screening procedure did not reveal significant changes in expression of genes for glutathione peroxidases (Gpx1,2,3,4) and superoxide dismutases (Sod1,2). Maximum values of Z_{ran} for these genes in rats receiving melatonin and Epithalon were +0.33 (Gpx2) and +0.27 (Sod1), respectively. The databank of the US National Institute of Health for detection, visualization, and complex study (<http://apps1.niaid.nih.gov/david/>) was used for further identification of transcripts similar to nucleotide sequences of clusters in the UniGene databank (46 transcripts). According to the classification of biological processes, 5 and 3 transcripts corresponded to categories of metabolism, cell growth, and/or maintenance of viability, respectively. Study with classification of molecular function revealed a correspondence of transcripts to categories of nucleic acid binding ($n=4$) and transferase activity ($n=3$).

Published data show that melatonin and Epithalon possess protective properties in the brain [4,6]. It can be hypothesized that several transcripts with modified expression are involved in the mechanism of action of these substances. Previous studies showed that melatonin inhibits expression of Tousled-like kinase 2 (Tlk2). This serine/threonine kinase is regulated by the cell cycle and functionally coupled to DNA replication [10]. Human Tik genes affect chromatin assembly during DNA replication. Their activity is maximum in S-

phase of the cell cycle and decreases under conditions of DNA damage and treatment with replication inhibitors [8]. Mutation-produced changes in Tik activity are followed by arrest of cell division (interphase) and apoptosis. However, overexpression of Tik impairs chromatin structure [8]. Melatonin inhibited expression of a guanine nucleotide-releasing factor RAN (Mog1). The mean values of Z_{ran} for Tlk2 and Mog1 were -1.55 and -1.65, respectively. The factor performs similar physiological functions and is required for nucleic protein import [3]. This factor is associated with RAN during DNA replication, regulation of the cell cycle, biosynthesis, processing, and transport [3]. The mean value of Z_{ran} for transforming growth factor β 1-induced transcript 4 (Tsc22) in the brain of mice receiving melatonin was -1.60. This factor plays an important role in the regulation of signals for cell death after DNA damage [5]. Overexpression of the factor initiates apoptosis in human tumor cells [12]. None of these genes were described in studying the biological effect of melatonin or epiphyseal peptides.

Melatonin significantly inhibited expression (average Z_{ran} -2.12) of the gene for interferon regulatory factor 3 (Irf3), which was probably related to the existence of functional relationship between melatonin, CREB-binding protein (CREBBP), and Irf3. The balance between phosphorylation and dephosphorylation of pCREB/CREB is mediated by serine/threonine phosphatase, modulates expression of the gene for arylalkylamine-N-acetyltransferase (Aanat), and regulates melatonin biosynthesis [14]. CREBBP and Irf3 contribute to coactivation of the mechanism mediated by double-stranded RNA-activated transcription factor 1 (DRAF1). Nuclear translocation of IRF3 is accompanied by the increase in phosphorylation of serine and

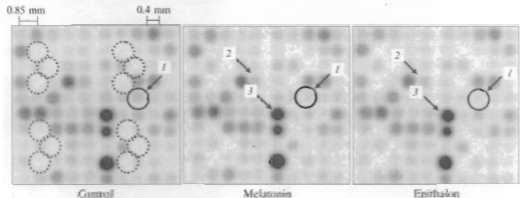


Fig. 1. Hybridization of brain samples from mice receiving physiological saline (control), melatonin, or Epithalon. Complete circle: clone with variants in expression, $Z_{\text{ran}} < -1.5$ (melatonin, -1.69; Epithalon, -0.93). Dotted circles: non-cDNA-containing spots. 1) expressed gene transcripts (ESTs), Nm_261776 cluster of UniGene databank; 2) guanosine triphosphate-binding protein 3 (Gltpt3) (pending); 3) subunit 2 of splicing factor 'sa' (Ssk2). Scales: diameter of membrane spots with applied cDNA (0.4 mm) and distance between adjacent spots (0.85 mm).

TABLE 1. Effects of Melatonin and Epithalon on Gene Expression in Mouse Brain

No.	Microchip		Z-ratio	Database		Gene
	A/B	position		mele-tonin	Epithalon	
1.	B	44-2-2	5.02	3.96	*BG085452	RIKEN cDNA 1810049E02 gene
2.	B	175-3-1	1.90	2.31	*AW651351	Mitochondrial gene for NADH dehydrogenase 4 (mt-Nd4)
3.	B	72-2-3	2.12	1.18	BG0730021	Expressed AA408740 sequence
4.	A	196-3-4	2.13	0.57	BG066802	ESTs weakly associated with retroviral polypeptide POL (POL2)
5.	B	117-2-1	0.26	2.03	BG072397	No data
6.	B	36-2-5	1.51	0.63	BG073331	Ribonucleic acid-binding protein S1
7.	B	69-2-3	-0.31	7.81	BG073008	ESTs weakly associated with RIKEN cDNA 5730493B19 gene
8.	B	19-2-2	-1.62	2.55	*BG085419	Ribosomal protein L19 (Rp19)
9.	B	154-1-4	0.35	-1.74	BG071739	ESTs
10.	B	95-1-2	-1.51	0.04	BG071020	ESTs
11.	B	1-3-5	-1.55	0.05	BG087221	Touled-like kinase 2 (Tlk2)
12.	A	185-1-2	1.10	-2.61	BG063396	RIKEN cDNA B230219D22 gene
13.	A	57-4-5	-1.55	-0.01	AL022853	ESTs
14.	B	117-4-4	1.25	-2.86	AW556794	Human Gmp2 gene for guanosine monophosphate reductase (LOC51292)
15.	B	247-4-4	-1.52	-0.17	AW557098	No data
16.	B	30-1-5	-0.09	-1.64	BG071980	ESTs moderately associated with reverse transcriptase
17.	B	25-1-2	-1.62	-0.10	BG070878	EST, chromosome 2
18.	B	35-1-3	-1.60	-0.13	BG084173	Transforming growth factor β 1-induced transcript 4
19.	A	246-1-2	-1.58	-0.16	BG063470	ESTs
20.	A	132-5-3	0.15	-1.90	BG082634	No data
21.	B	356-1-3	-2.39	0.57	BG071576	ESTs
22.	B	287-5-1	-2.06	0.15	—	No data
23.	A	378-4-5	-0.03	-1.94	BG068750	ESTs
24.	B	161-2-1	-1.91	-0.07	BG072440	RIKEN cDNA 1700027N10 gene
25.	B	328-1-4	-0.36	-1.82	BG071888	ESTs moderately associated with COI protein
26.	B	80-5-5	-1.74	-0.32	BG052303	ESTs strongly associated with cerebral cavernous malformation gene 1 (CCm1)
27.	B	303-3-1	-1.64	-0.48	BG066337	RIKEN cDNA 2810031L11 gene
28.	B	205-5-4	-1.58	-0.53	BQ551867	ESTs
29.	A	55-5-1	-1.85	-0.29	BG068305	RIKEN cDNA 4900595265 gene
30.	B	303-5-3	1.66	-3.69	BQ513566	ESTs
31.	B	237-5-3	-1.52	-0.71	BQ551391	ESTs

TABLE 1.

No.	Microchip		Z-ratio		Database		Gene
	A/B	position	mela- tonin	Epi- thalon	GenBank	UniGene	
32.	B	192-4.4	-0.69	-1.55	BG088274	Mm.30077	ESTs weakly associated with isoform 1 of human caldesmon gene (L-CAD, H-CAD, NAG22 protein, L-caldesmon II)
33.	A	348-4.1	-1.53	-0.77	BG067645	Mm.221064	EST
34.	A	40-2.5	-1.67	-0.68	BG065627	Mm.24899	ESTs
35.	B	133-4.4	-1.58	-0.78	BG088220	Mm.18561	Gene for cysteine- and histidine-rich protein 1 (Cyhr1)
36.	B	301-5.4	-1.68	-0.67	BQ552034	Mm.221668	ESTs
37.	A	45-1.5	-1.65	-0.73	BG077598	Mm.143774	Guanine nucleotide-releasing factor RAN (Mog1)
38.	B	329-1.3	-0.61	-1.81	AU043423	No data	No data
39.	A	275-5.3	-2.12	-0.34	BG069754	Mm.3960	Interferon regulatory factor (Irf3)
40.	B	25-3.5	-1.88	-0.63	BG074613	Mm.77786	ESTs
41.	B	174-4.4	-2.03	-0.52	BG075723	Mm.103003	Expressed sequence AU015603
42.	A	28-5.3	-1.69	-0.93	BG069529	Mm.26176	ESTs
43.	B	227-4.3	-1.64	-1.05	BG075441	Mm.103168	ESTs
44.	A	48-1.4	-1.35	-1.85	BG063926	Mm.197220	ESTs
45.	A	141-2.5	-2.20	-1.04	BG065708	Mm.27654	ESTs weakly associated with the gene for Diaphanous protein homologous 1 (Dm1, DRF1, p140mDia)
46.	A	5-5.2	-1.85	-1.49	BG069123	Mm.173776	ESTs
47.	B	48-2.1	-1.03	-2.40	BG072943	Mm.11227	Expressed sequence AU017169
48.	A	207-5.3	-1.92	-1.61	*BG069687	Mm.32532	EST, chromosome 16
49.	B	211-1.2	-3.28	-0.61	BG071133	No data	No data
50.	A	277-3.4	-1.91	-2.40	*BG080013	Mm.34798	RIKEN cDNA 493050K01 gene
51.	B	151-4.4	-2.59	-1.99	*BI076814	Mm.9626	Dihydropteroate branched chain transacylase E2 (Dbt)
52.	B	270-4.4	-2.74	-2.45	*AW557182	Mm.43213	RIKEN cDNA 9030402K04 gene
53.	A	234-5.2	-0.77	-5.10	BG082365	Mm.8149	Male germ cell-associated kinase (Mak)

Note. Transcripts with significant changes in expression (1.5Z₂-5.1Z₃). Transcripts were classified (M) by mean values of Z₂ in experimental groups, A/B: membrane of microchip (A or B). Position: position of a clone in the membrane (grid). Z₂-ratios: mean values in experimental groups (1.5Z₂-5.1Z₃, printed in italic type). GenBank: index of GenBank database (NCBI). UniGene: number of a cluster in UniGene database (NCBI). *Statistically significant changes in expression of transcripts in both experimental groups (melatonin and Epithalon). EST: expressed sequence; Mog1: expressed gene fragment.

threonine, activation of DRAFI, and specific changes in gene expression produced by viral infections [15].

Expression of 6 transcripts (dihydroliipoamide branched chain transacylase E2 (Dbt), 4 unidentified genes and expressed gene fragments (ESTs, expressed sequence tags), and mitochondrial gene for NADH dehydrogenase 4; mt-Nd4) underwent similar changes in mice receiving melatonin and Epithalon. Expression of 1 transcript (ribosomal L19 (Rpl19)) was inhibited by melatonin, but induced by Epithalon. Mutations in the only described gene of this group (Dbt) are followed by the development of type 2 maple syrup urine disease (MSUD2). The molecular mechanisms underlying suppression of gene expression in the brain of mice receiving melatonin and Epithalon remain unclear.

Comparative study revealed qualitatively different changes in gene expression in the brain and heart of mice receiving melatonin and Epithalon [1,2]. The transcript corresponding to mouse mitochondrial gene for NADH dehydrogenase 4 (mt-Nd4) was the only transcript whose expression in the brain and heart of CBA mice underwent similar changes after treatment with Epithalon (average Z_{rel} 2.31 and 2.03, respectively) and melatonin (average Z_{rel} 1.71 and 1.90, respectively). Transcription of other transcripts in tissues did not significantly change. Our results confirm the hypothesis that Epithalon [4] and melatonin [3] have a tissue-specific biological effect. The action of melatonin can be mediated by specific receptors and various transcription factors. A screening procedure was performed with 16,897 and 15,247 transcripts in the brain and heart of mice, respectively. The number of highly expressed transcripts in the brain was much lower than in the heart. These differences can be related to the above mentioned reason. A larger sample of tissues should be studied to derive a final conclusion about the tissue-specific effect of Epithalon and melatonin.

A large-scale study of the influence of melatonin and Epithalon on gene expression in the brain of mice revealed their specific and general effects. Target ge-

nes are physiologically related to the cell cycle, apoptosis, biosynthesis, processing, and transport of nucleic acids.

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