Early-Stage Retinal Melatonin Synthesis Impairment in Streptozotocin-Induced Diabetic Wistar Rats

Daniella do Carmo Buonfiglio,1 Rodrigo Antonio Peliciari-Garcia,1 Fernanda Gaspar do Amaral,1 Rafael Peres,1 Tatiane C. Araujo Nogueira,1 Solange Castro Afeche,2 and José Cipolla-Neto1

PURPOSE. Retinal melatonin synthesis occurs in the photoreceptor layer in a circadian manner, controlling several physiologic rhythmic phenomena, besides being the most powerful natural free radical scavenger. The purpose of the present work was to evaluate the diurnal profile of retinal melatonin content and the regulation of its synthesis in the retina of streptozotocin-induced diabetic rats.

METHODS. Diabetes was induced in male Wistar rats (12 hour–12 hour light/dark cycle) with streptozotocin. Control, diabetic, and insulin-treated diabetic animals were killed every 3 hours throughout the light–dark cycle. Retinal melatonin content was measured by high-performance liquid chromatography, arylalkylamine N-acetyltransferase (AANAT) activity was analyzed by radiometric assay, Bmal1 gene expression was determined by qPCR, and cyclic adenosine monophosphate (cAMP) content was assessed by ELISA.

RESULTS. Control animals showed a clear retinal melatonin and AANAT activity daily rhythm, with high levels in the dark. Diabetic rats had both parameters reduced, and the impairment was prevented by immediate insulin treatment. In addition, the Bmal1 expression profile was lost in the diabetic group, and the retinal cAMP level was reduced 6 hours after lights on and 3 hours after lights off.

CONCLUSIONS. The present work shows a melatonin synthesis reduction in diabetic rats retinas associated with a reduction in AANAT activity that was prevented by insulin treatment. The Bmal1-Battened gene expression and the cAMP reduction seem to be responsible for the AANAT activity decrease in diabetic animals. The melatonin synthesis reduction observed in the pineal gland of diabetic rats is also observed in a local melatonin tissue synthesizer, the retina. (Invest Ophthalmol Vis Sci. 2011;52:7416–7422) DOI:10.1167/iovs.10-6756

Melatonin (N-acetyl-5-methoxytryptamine), synthesized mainly by the pineal gland, is also produced in the retina of several vertebrate species. Retinal melatonin synthesis occurs in a subpopulation of photoreceptors and, as observed in the pineal gland, shows a clear daily variation with low levels during the light period and high levels at night. Results of previous studies have demonstrated that melatonin synthesis pathway enzymes are under circadian control, at either transcriptional or posttranscriptional levels in several species.

Arylalkylamine N-acetyltransferase (AANAT) is the main regulatory enzyme in melatonin synthesis and both its mRNA level and activity show a marked circadian variation in rat retina. Physiological target of the BMAL1:CLOCK complex with the E-box present in the AANAT promoter. The circadian clock also works as a gate, regulating rhythmic adenylyl cyclase (AC1) mRNA expression and controlling the period during the day when cAMP levels are maximally stimulated. In this way, melatonin synthesis is exclusively gated to the night, by increased Ca2+ influx and stimulation of cAMP synthesis late in the day and at night, when AC1 protein expression is high, but not during the early morning.

Several retinal physiologic phenomena were described as rhythmic, including rod disc shedding, visual sensitivity, melatonin release in vivo and in vitro, dopamine synthesis, electroretinogram b-wave amplitude, extracellular pH, and intraocular pressure. In addition, most retinal rhythms are partially modulated by rhythmic melatonin and dopamine release.

Melatonin is one of the most powerful natural free radical scavengers, preventing oxidative damage to macromolecules, including lipids, proteins, and nucleic acids. Gül et al. have shown that melatonin reduces corneal injury in a streptozotocin (STZ)-induced diabetic rat model. In an induced ischemia model, melatonin blocks apoptosis in cultured human retinal pigment epithelium (RPE) cells.

In diabetes mellitus, the hyperglycemia increases the enzymatic conversion of glucose to polyalcohol sorbitol, with concomitant nicotinamide adenine dinucleotide phosphate (NADPH) and glutathione decrease, enhancing the oxidative stress that may be involved in the development of diabetic retinopathy.

From the 1Department of Physiology and Biophysics, Institute of Biomedical Sciences, University of São Paulo, São Paulo, Brazil; and the 2Laboratory of Pharmacology, Butantan Institute, São Paulo, Brazil.

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Corresponding author: José Cipolla-Neto, Department of Physiology and Biophysics, University of São Paulo, Av. Prof. Lineu Prestes, 1524, São Paulo, SP, CEP 05508-900, Brazil; cipolla@icb.usp.br.

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Considering the essential function of melatonin on retinal physiology and the functional interrelationship between insulin and melatonin,\textsuperscript{32–35} we investigated the diurnal profile of retinal melatonin content and AANAT activity in the retina of STZ-induced diabetic rats.

To the best of our knowledge, this is the first evidence of decreased melatonin production in retinas of STZ-induced diabetic rats.

MATERIALS AND METHODS

Animals

Male Wistar rats weighing 240 to 280 g were obtained from the Institute of Biomedical Sciences, University of São Paulo, São Paulo, Brazil. The animals were kept under a 12-hour-light/12-hour-dark cycle (lights on at 6 AM; Zeitgeber Time [ZT]0), in a temperature-controlled room (21 ± 2°C), with food and water ad libitum. Ethics approval was granted for this study by the Committee of Ethics in Animal Experimentation of the Institute of Biomedical Sciences, University of São Paulo, and it is in compliance with the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research.

Experimental Design and STZ-Induced Diabetes

Animals were divided into three groups: control, untreated diabetic, and insulin (INS)-treated diabetic group. Diabetes was induced by a single intraperitoneal injection of STZ (60 mg/kg body weight; Sigma-Aldrich, St. Louis, MO) freshly diluted in citrate buffer (10 mM, Na citrate; pH 4.5). Nondiabetic control rats were injected with citrate buffer in the same volume. All the injections were performed during the daytime. Body weight and glycemia were measured before and after the STZ intraperitoneal injection. Tail blood was collected for glucose determination using a glucometer (Optium Xceed; Medisense, Indianapolis, IN), and insulin (INS)-treated diabetic and control groups, showing that the insulin observed in the induced animals. No significant difference was found between the INS-treated and control groups, 361.7 control levels (STZ-induced diabetic 130.1 versus control 95.81, p < 0.0001, STZ-induced diabetic versus control and INS-treated). The effects of diabetes on retinal melatonin synthesis were evaluated 3 days after STZ or citrate buffer injection. Independent groups of control, STZ-induced diabetic and INS-treated diabetic animals were killed every 3 hours throughout the 24-hour light-dark cycle.

Pinealectomy

To verify whether circulating melatonin levels influence the amount of retinal melatonin we performed a pinealectomy (PINX), as previously described.\textsuperscript{37} Control and PINX animals were killed during the dark period at ZT12, -15, -18, -21, and -24, at 30 days after surgery.

Melatonin Levels Measurement

Retinal melatonin levels were measured by high-performance liquid chromatography (HPLC) with electrochemical detection (Empower software; Waters, Milford, MA).\textsuperscript{38} Melatonin was separated on a C18 column (5 μm, 150 × 3.9 mm; Resolve; Waters). The chromatographic system was isocratically operated with the following mobile phase: 0.1 M sodium acetate, 0.1 M citric acid, 0.15 mM ethylene diamine triacetic acid (EDTA), and 35% methanol (pH 3.7), at a flow rate of 1 mL/min. The electrochemical detector potential was adjusted to +900 mV. The elution time for melatonin was approximately 6 minutes. Each retina was sonicated (Microson XL 2005; Heat System, Inc., Farmingdale, NY) in a solution of 0.1 M perchloric acid (120 μL), containing 0.02% EDTA and 0.02% sodium bisulfite. After centrifugation (2 minutes, 13,000g; Eppendorf 5415C Centrifuge; Brinkman Instruments Inc., Westbury, NY), 40 μL of the supernatant was injected into the chromatographic system (model 7125 Injector, 20-μL loop; Rheodyne Inc., San Francisco, CA).

AANAT Activity Measurement

AANAT activity was measured by a radiometric assay.\textsuperscript{39,40} Briefly, 100 μL of 0.1 M sodium phosphate buffer (pH 6.8), containing 40 mM tryptamine and [3H]-acetyl coenzyme A (2 mM; final specific activity = 4 μCi/μmol; Sigma-Aldrich), was added to a microcentrifuge tube containing one retina kept at 4°C. The retinas were sonicated and then incubated at 37°C for 20 minutes. The reaction product N-[3H]-acetyltryptamine was extracted with chloroform (1 mL). Samples of 500 μL were evaporated until dry in a scintillation vial, and radioactivity was determined (Tri-Carb 2100 Packard β counter; GMI Inc., Ramsey, MN).

Cell Viability and DNA Fragmentation

Retinas obtained from nondiabetic and diabetic rats killed by decapitation were dissected, transferred to a phosphate buffer solution, and incubated for 20 minutes at 57°C with 0.25% trypsin-EDTA (Invitrogen-Gibco, Grand Island, NY). Enzymatic digestion was stopped by the addition of Dulbecco’s modified Eagle’s medium (DMEM; CultiLab, Campinas, SP, Brazil) containing 10 IU/mL penicillin, 10 μg/mL streptomycin, 10% fetal bovine serum, 2 mM glutamine, and 25 mM glucose. Tissue was mechanically dissociated with a micropipette, and cells were evaluated as described below.

Table 1. Primer Sequences for Rat Bmal1, Actb, Rp137 and Actg1, Real-Time Quantitative RT-PCR Assays

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<th>Target and Reference Genes</th>
<th>RefSeq</th>
<th>Sequence</th>
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<tr>
<td>Bmal1 (Arntl)</td>
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<td>5'-GGATGACGCACTGTGAGAAGGGACAC-3'</td>
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<td></td>
<td></td>
<td>5'-CCATGAGGAGAAGAGGGCGGACCAAG-3'</td>
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<td>5'-TCATGCGCACCTGAGGCAGAG-3'</td>
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<tr>
<td>Actb</td>
<td>NM_051144</td>
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<tr>
<td>Rp137a</td>
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<tr>
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</table>

RefSeq is provided by the National Center for Biotechnology Information, Bethesda, MD (www.ncbi.nlm.nih.gov/locuslink/refseq/).
Rat retina cells were washed and suspended in phosphate-buffered saline (PBS). Afterward, 50 μL of propidium iodide solution (50 mg/mL in PBS) was added, and the cells were analyzed with flow cytometry (FACSCalibur; BD Biosciences, San Jose, CA). Fluorescence was measured using the FL2 channel (orange-red fluorescence; 585/42 nm). Ten thousand events were analyzed per experiment. Cells with propidium iodide fluorescence were then evaluated (Cell Quest software; BD Biosciences). DNA fragmentation was calculated after DNA staining with propidium iodide according to a method previously described.41 The pellet was gently resuspended in 300 μL hypotonic solution containing 50 mg/mL propidium iodide, 0.1% sodium citrate, and 0.1% Triton X-100. The cells were then incubated for 1 hour at room temperature. Fluorescence was measured and analyzed as described above.

cAMP Immunoassay

Retinas were homogenized in 0.1 M HCl using a homogenizer (Polytron; VWR International, Lutterworth, UK). After 5 minutes of centrifugation, the supernatants were collected and assayed directly according to the kit protocol (BioVision Research Products, Mountain View, CA).

Real-Time Quantitative PCR

Retinal total RNA was isolated using a guanidine isothiocyanate-based reagent (Trizol; Invitrogen Corp., Carlsbad, CA), according to the manufacturer’s specifications. Total RNA was eluted in RNase-free water, treated with DNase genomic DNA remover (Turbo DNasefree; Ambion, Austin, TX) according to the manufacturer’s specifications. Reverse transcriptase (200 U Superscript III; Invitrogen Corp.), DTT (10 nM), dNTP (10 mM each), RNase inhibitor (40 U), and random primers (150 ng) were used to reverse transcribe 1 μg of total RNA in a final reaction volume of 20 μL. qRT-PCR was performed (7500 Real-Time PCR System; Applied Biosystems, Inc. [ABI], Foster City, CA) with 25-μL reactions containing 2 μL of cDNA (10 ng/μL), SYBR green (Power SYBR Green; ABI), and 400 nM specific intron-spanning primers (Table 1). The cycling parameters included an initial denaturation step at 95°C for 10 minutes, followed by 40 cycles of 95°C for 15 seconds and annealing and extension at 60°C for 1 minute. Melting curve analyses (Tm) and ethidium bromide (EtBr) agarose gel (2.2% wt/vol) electrophoresis of the amplified products were performed for product specificity assessment. A set of 10-fold serial dilutions of each internal standard (101–107 copies/2 μL) was used to generate a standard curve, and all qRT-PCR assays were linear within this concentration range with correlation coefficients (r2 > 0.999). Transcript numbers were determined by the system software (model 7500, ver. 2.0.3; ABI) and normalized using the geometric mean calculated from the reference genes Actb, Rpl37a, and Actg1, which were indicated as the most suitable ones by the software GeNorm.42

Statistics

All the results were plotted as the mean ± SEM. One-way ANOVA was used to evaluate the influence of the variable “time of day” on each temporal series, two-way ANOVA with Bonferroni’s posttest was used to compare groups regarding the factors time of day, treatment (control, diabetic, and INS-treated), and the interaction between them. Student’s t-test was used when appropriate (all statistical tests and graphics: Prism, ver. 5.01 for Windows; GraphPad Software, San Diego, CA). Results are representative of at least two independent experimental blocks.

RESULTS

Cell Viability and DNA Fragmentation

The flow cytometer results showed that 3 days of STZ-induced diabetes did not cause retinal cell necrosis (viability test) or retinal cell apoptosis (DNA fragmentation test) when compared to control group values (Fig. 1).

Daily Retinal Melatonin Profile after 3 Days of STZ-Induced Diabetes

As expected, retinas of the control animals showed a clear daily rhythm of melatonin synthesis, with high levels during the night and low levels during the day (Fig. 2). In the retinas of the diabetic rats, a nearly 40% reduced melatonin synthesis at ZT18 (P < 0.01) and ZT21 (P < 0.05) was observed compared with that of the control (Fig. 2). On the other hand, insulin treatment prevented melatonin reduction, since no statistical significance was found between the control and INS-treated diabetic animals after 3 days of treatment (Fig. 2).

Retinal Melatonin from PINX and Intact Animals

As can be observed in Figure 3, there was no difference in the retinal melatonin content of the PINX and intact animals. This finding leads us to conclude that pineal melatonin does not influence the amount of retinal melatonin in rats.

Daily AANAT Activity Profile after 3 Days of STZ-Induced Diabetes

In an attempt to investigate why retinal melatonin is reduced in diabetic rats, we analyzed the daily AANAT activity profile. The AANAT activity of the control group presented...
higher values in the dark period at ZT15 (P < 0.01) and lower levels in the light period at ZT6 (P < 0.01) in comparison to that in the diabetic group (Fig. 4). On the other hand, the retinal AANAT activity daily fluctuation that was abolished in rats with STZ-induced diabetes was completely restored in the INS-treated group.

**Daily Retinal Bmal1 mRNA and cAMP Content after 3 Days of STZ-Induced Diabetes**

*Bmal1* gene expression and cAMP content analysis were performed to elucidate the AANAT activity reduction observed in the diabetic group (Fig. 5).

Our data show *Bmal1* that gene expression was high in the middle of the day (ZT6) and at ZT12, -15, and -18 in the control group. In the diabetic group, this pattern of expression vanished completely (Fig. 5A). The retinal cAMP level was significantly reduced at ZT6 (P < 0.001) and at ZT15 (P < 0.001) in the diabetic group, in comparison with that in the control group (Fig. 5B).

**Retinal Melatonin and AANAT Activity at ZT18 after 15 Days of STZ-Induced Diabetes**

To verify whether the reduction of retinal melatonin content and AANAT activity persists, we analyzed both parameters at ZT18 after 15 days of STZ-induced diabetes (Fig. 6). We observed a 75% reduction in the retinal melatonin content and a decrease in AANAT activity in the diabetic animals compared with that in the control animals after 15 days of induction.

**DISCUSSION AND CONCLUSIONS**

Several studies have shown the protective properties of melatonin in retinas of STZ-induced diabetic rats. Our work is the first to show the effects of STZ-induced diabetes on retinal melatonin synthesis.

In the present work, the STZ-induced diabetic condition was established in our rats and verified by high blood glucose concentrations and hypoinsulinemia (data not shown). The insulin treatment was capable of restoring the blood glucose concentration to control levels.

Initially, the cellular viability and DNA fragmentation of retinas from control and diabetic rats did not show any significant statistical difference, validating the study of retinal melatonin synthesis in the STZ-induced diabetes model. Park et al. showed, in an STZ-induced diabetes model, degenerative changes from 1 week onward in postsynaptic processes of horizontal cells in the deep invaginations of the photoreceptor cell layer, where melatonin synthesis takes place. The same group reported photoreceptor cell apoptosis after 4 weeks of disease. Based on these studies, it is unlikely that a level of retinopathy is established after 3 days of diabetes that could affect the results that we obtained. However, we have to consider that flow cytometry does not specifically evaluate the photoreceptors, which are the melatonin-synthesizing cells.

The circadian profile of retinal melatonin content observed in the control group corroborates that found by Tosini and Menaker. On the other hand, we showed a 40% reduction on the nocturnal retinal melatonin content in the STZ-induced diabetic group in comparison to the control. Moreover, the reduction in the retinal AANAT activity corresponded to the reduced melatonin synthesis. In addition, when the animals had their diabetes treated by the administration of insulin, we found a complete reestablishment of both the melatonin synthesis profile and AANAT activity. Furthermore, we demonstrated a reduction in melatonin synthesis and AANAT activity 15 days after STZ induction.

There is no information in the literature about retinal melatonin synthesis in diabetic rats; the available data concern pineal melatonin synthesis in different diabetes models. In accordance with our results in the retina, some reports described a significant reduction of pineal melatonin synthesis in diabetic rats induced by alloxan or STZ (Amaral FG, personal communication, 2010) as well as in hamsters. Decreased plasma melatonin levels were found in type 1 diabetic patients, in type 2 diabetic patients, and in type 2 diabetic rats.

In an attempt to elucidate whether the reduced plasma melatonin content observed in diabetic rats would explain the reduced levels of retinal melatonin in our diabetic animals, we analyzed the retinal melatonin content in pinealectomized rats and compared it with that in control animals. No difference was found between the groups, so it is possible to conclude that the melatonin measured in the retina is exclusively of retinal origin. In this regard, the melatonin reduction found in the diabetic group retinas is most probably due to the impaired intrinsic mechanism of melatonin synthesis in the retina that led us to proceed with the analysis of retinal AANAT activity.

Considering the AANAT activity profile in the retinas, our results are in agreement with those of Niki et al. who demonstrated the circadian pattern of this enzyme activity. STZ-induced diabetic rats showed a loss of AANAT activity rhythmicity, which was partially restored in INS-treated animals. Although insulin treatment did not totally recover the control AANAT activity pattern, the melatonin content results show that insulin reposition was able to recover not only the retinal melatonin level, but also the rhythmic pattern that is observed.
Bmal1 expression contributing to the arrhythmic AANAT activity and reduction observed (Fig. 7).

Recent data from the literature suggest that both Clock and Bmal1 participate in the regulation of energy metabolism. Young et al. showed that STZ-induced diabetes is associated with alterations in the circadian clock of the rat heart. Further experiments are under way to evaluate CAMP decline and how the retinal circadian clock is affected by diabetes.

There are no data in the literature that show the effects of diabetes on the retinal melatonin synthesis pathway. On the other hand, it has been reported that retinal insulin signaling is in a constitutively higher steady state and that the basal phosphorylation and activity of the insulin receptor and serine/threonine protein kinase (Akt) maintain a tonic cell-survival signal in the metabolically active retina. Diabetes induced by STZ treatment gives rise to late neural cell death in the retina, but it has not been established yet whether this effect is directly related to reduced circulating insulin, or the reduction of insulin increases apoptosis by an indirect mechanism such as hyperglycemia. Considering our results, we do not know whether there is a direct action of insulin in the retinal melatonin synthesis pathway, or the effect is caused by hyperglycemia, or both factors contribute. Insulin treatment prevented a reduction in melatonin synthesis, but it also controlled the hyperglycemia.

Although additional studies are necessary to clarify the mechanisms underlying the decreased insulin and consequent hyperglycemia involvement in the retinal melatonin synthesis impairment, it is important to note that the melatonin synthesis reduction already reported in the pineal gland of STZ-induced diabetic rats is also observed in a local melatonin tissue synthesizer, the retina.

In this way, in the presence of diabetes, the reduction of local melatonin synthesis, a powerful natural antioxidant, may contribute, in addition to the well-known impairment in control rats. Thus, the AANAT activity peak found at ZT15 supports the melatonin synthesis peaks observed at ZT18 and -21 in INS-treated animals. Reinforcing our data, Lynch et al. demonstrated, in vitro, that insulin is able to potentiate the norepinephrine-induced melatonin synthesis in the rat pineal gland via phosphatidylinositol-3 kinase (PI3K) signaling. 

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It is known that the BMAL1:CLOCK complex is involved in AC1 and Aanat gene expression regulation and the cAMP signaling pathway is crucial for AANAT stability. Our data show high Bmal1 gene expression in the middle of the day (ZT6) and at ZT12, -15, and -18 in the control group. This multipeaked control profile is probably due to the measurement of Bmal1 mRNA levels in the whole retina, not just in the melatonin-synthesizing cells. The retina is a multilayered tissue that contains different cell types, and Bmal1 expression has been detected in the photoreceptors, the inner nuclear layer, and the ganglion cell layer.

In the diabetic retinas, the variation in the daily expression of Bmal1 was lost in comparison with that in the control retinas, leading us to suppose that this flattened Bmal1 expression could impair the AC1 levels, resulting in the observed decrease in cAMP level and consequent AANAT activity reduction at ZT15. Moreover, Aanat gene expression may also be affected by the flattened Bmal1 expression contributing to the arrhythmic AANAT activity and reduction observed (Fig. 7).

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in retinal antioxidant defense, to the development of diabetic retinopathy, a frequent complication of this disease.

Acknowledgments

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