Acute Experimental Diabetes and Aortic Depressor Nerve Ultrastructural Morphometry: Effects of Insulin Treatment

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Abstract

Previous results from our laboratory showed myelinated fiber alterations in the aortic depressor nerve (ADN) from rats rendered diabetic for 15 days (short term). The ultrastructure and the acute effect of insulin treatment, particularly in the unmyelinated fibres of this nerve, were not yet investigated in this experimental model, being the aim of this work. Wistar rats received a single intravenous injection of streptozotocin (STZ – 40 mg/Kg) 15 days (N=7) before the experiments. Control animals (N=7) received vehicle (citrate buffer) and treated animals (n=6) received a subcutaneous injection of insulin (10 IU/Kg) on a daily basis beginning 3 days after the STZ injection. Under pentobarbital anaesthesia the ADNs were isolated and had their spontaneous activity recorded. Afterwards, proximal and distal segments of the nerves were prepared for transmission electron microscopy studies. Morphometry of the myelinated and unmyelinated fibres was carried out with the aid of computer software. Nerves from diabetic animals showed some unmyelinated fibre enlargement with signs of swelling and degeneration, and the Schwann cells mitochondria presented enlargement and clustering formation. On treated animals, an important thickening of the Schwann cells' basement membrane was present and some myelinated fibers showed thin myelin sheaths. The endoneural blood vessels ultrastructure was preserved in all nerves from the three experimental groups. Myelinated axon area and diameter was significantly larger on insulin treated animals, compared to both, controls and acute diabetic groups, on both segments. No other differences were observed between groups for the myelinated or unmyelinated fibers morphometric parameters. Investigation of experimental models of diabetes in early or acute stage of the disease provide relevant information on the physiopathological mechanisms involved in the alterations present in patients with chronic diabetes and its complications. We suggest that insulin treatment alone might have a role on the morphological alterations present in diabetic neuropathy.

Keywords: Aortic depressor nerve; Acute diabetes; Insulin treatment; Morphometry; Ultrastructure

Introduction

Diabetes is a metabolic disorder of multiple etiologies caused by defects in insulin secretion, insulin action, or both, resulting in chronic hyperglycemia. Initial symptoms such as thirst, polyuria, and weight loss are unspecific and often not severe, or even absent. Thus, hyperglycemia sufficient to cause pathological associated or not to functional changes, may be present for a long time before the diagnosis is made. Neuropathy is a common complication of diabetes that accounts for a significant high morbidity [1]. Comi and Corbo [2] documented that one third of diabetic patients present some degree of neuropathy but a recent publication from the Centers for Disease Control and Prevention [3] indicated that this percent is much higher (up to 70%). Despite recent information about the mechanisms for many of the long term complications of diabetes, the exact pathogenesis of diabetic neuropathy remains unknown.

The literature is extensive on investigation of peripheral nervous system alterations in experimental models of diabetes (induced or genetic) but morphological alterations totally comparable to those that occur in the human diabetic neuropathy are not completely reproducible [4]. Nevertheless, it is expected that alterations observed in experimental models, especially in the early or acute cases, will be useful for a better understanding of the neuropathy development mechanisms [4]. Among the alterations on the peripheral nervous system due to diabetes, the involvement of the autonomic nervous system is the least understood. The autonomic diabetic neuropathy is often under diagnosed and the most common clinical manifestation is sexual dysfunction. Functional alterations of the autonomic cardiovascular reflexes have been demonstrated in clinical and animal models [5-8] and most of the reports about altered baroreflex sensitivity attribute this alteration to the diabetic vagal (afferent) neuropathy.

The aortic depressor nerve (ADN) is unique in rodents because it contains only baroreceptor afferents (sensory) fibers [9,10]. The ADN (afferent arm of the baroreflex) ultrastructure was recently explored in a long term experimental model of diabetes [11] with important

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morphological alterations being demonstrated. Nevertheless, the ultrastructural investigation of the ADN in acute diabetes was not previously reported, being the purpose of this study.

Materials and Methods

All described experimental procedures adhered to the Guide for the Care and Use of Laboratory Animals prepared by the National Academy of Sciences and published by the National Institutes of Health (Copyright©1996 by the National Academy of Sciences), and were approved by the Institutional Ethics Committee for Animal Research (CETEA–Comité de Ética em Experimentação Animal, protocol number 220/2005). All effort was made to minimize the number of animals used.

Male Wistar rats, from the animal care facility of the Campus of Ribeirão Preto (University of São Paulo), were housed in plastic cages with free access to tap water and rat chow throughout the experiment. Diabetes induction was performed through a single injection of streptozotocin (STZ) (40 mg/kg, Sigma Chemical Co., St. Louis, MO, USA) into the penile vein 15 days (N=7) before the experiments. Nonfasting blood glucose (mg/dL) was determined by means of a glucose analyzer (Beckman Instruments Inc., Brea, CA, USA) on the third day after STZ injection and at the time of the experiments. Animals with glucose levels higher than 350 mg/dL, observed on the 3rd day after STZ injection, were considered diabetic and included in the study. Insulin treated animals (N=6) received a subcutaneous injection of 9 IU/kg of NPH (highly purified mixed) insulin (Biolin U-100, Biobras) in the evening, at the beginning of the dark cycle, from the 3rd day after STZ injection throughout the experiment. The animals were weighed every 3 days and, when necessary, the insulin dose was adjusted. Control animals (N=7) received a single injection of vehicle (citrate buffer, pH 5.2) into the penile vein 15 days (N=7) before the experiments.

Under pentobarbital anesthesia, the ADNs were isolated and had their spontaneous activity recorded as previously described [12]. The pulsatile arterial pressure was recorded simultaneously with the ADN activity, to ensure that the nerves studied were the ADN. Proximal and distal segments of the left ADN were prepared for light and transmission electron microscopy studies as reported previously [11,13,14]. Morphometry was performed with the aid of the KS 400 software (Kontron 2.0, Eching Bei Muchen, Germany). The investigator was blinded to group identity throughout the morphometric process. Sample morphometry was not used and 100% of the fascicle area was studied.

All nerve samples were processed at once to assure that the alterations observed were not due to artifacts introduced by the histological procedures. The light microscopy study was performed to validate the quality of the histological preparation of the nerves and to perform the light microscopy studies. Only nerves with good histological preservation were used. The fascicle images were collected from semithin transverse sections, stained with 1% toluidine blue, using a digital camera (AxioCam MR5, Carl Zeiss) connected to an IBM PC. With the aid of KS 400 software, the total number of myelinated fibers and the total number of Schwann cells nuclei present in each fascicle was identified by visual inspection and counted. The fascicular area was measured and the myelinated fiber density (fiber/mm²) and the Schwann cell nuclei density (nucleus/mm²) were calculated. Myelinated fiber morphometry was performed as described previously [15,16] and the parameters obtained were axonal area and diameter and total fiber area and diameter. Also, the ratio between the axonal diameter and total fiber diameter (which indicates the degree of myelination), described as G ratio [17,18], was obtained. Finally, myelin sheath area was calculated for each myelinated fiber measured.

For the ultrastructural studies, thin transverse sections were mounted on Formvar coated 2×1 slot grids, stained with lead citrate and uranyl acetate, and observed with a JEM-1230 transmission electron microscope (JEOL-USA, Inc., Peabody, MA, USA) equipped with a digital camera. Using the image analysis software, the total number of the unmyelinated fibers was counted, and their diameter measured as described [11,19-21]. The density of the unmyelinated fiber was calculated and the ratio between unmyelinated/myelinated fibers was determined. A histogram of the unmyelinated fiber diameter was constructed and separated into class intervals increasing by 0.1 μm. Data are presented as mean ± standard error of mean (SEM).

For the statistical analysis, the Kolmogorov-Smirnov test was applied to verify the normal distribution for weight, biochemical, physiological and morphometric data, using Sigma Stat software, version 3.01 (Jandel Scientific), followed by the Levene test of medians for variance equivalence.

Body weight and blood glucose level data were compared between the 3rd day of injection and the experimental day by the non-parametric test of Wilcoxon for paired samples and among groups by the analysis of variance on Ranks, followed by Tukey’s post hoc test, provided that these data did not show normal distribution. The mean arterial pressure (MAP) and heart rate (HR) data between groups were also compared by the analysis of variance on Ranks, followed by Tukey’s post hoc test.

Morphometric data that passed the normality and variance equivalence tests were compared by paired Student’s t-test (between the proximal and distal segments) and one way ANOVA followed by a Holm-Sidak post hoc test (for segments between groups). Otherwise, proximal and distal segments were compared using the non-parametric test of Wilcoxon for paired samples, while groups were compared by the analysis of variance on ranks, followed by Dunn’s post hoc test.

Differences were considered significant when p<0.05.

Results

Biochemical and physiological data

At the beginning of the experiments, there was no difference in body weight between al groups. At the experimental day, all groups showed significant increase in body weight comparing to the beginning of the experiments but STZ animals gained less weight compared to control and compared to insulin treated groups. Blood glucose level was significantly high on both STZ groups (treated and untreated) three days after injection compared to controls. This difference was also observed on the experimental day, but treated animals showed a significant reduction on the blood glucose level compared to the beginning of treatment. Mean arterial pressure was significantly reduced on STZ animals compared to controls and treated animals, with no difference on the heart rate between groups. Numerical data is presented on Table 1.

Morphology

No evident morphological differences were observed between the experimental groups in the light microscopy level, except that the insulin treated nerves seemed to be smaller compared to nerves from the other two experimental groups. The ultrastructural qualitative analysis showed that control nerves present morphological...
characteristics similar to the normal ADN described for rats [12]. Nerves from diabetic animals showed some unmyelinated fiber enlargement with signs of swelling and degeneration, but the most striking alteration was observed on the Schwann cells mitochondria, that presented enlargement and clustering formation (Figure 1). For the insulin treated animals, an important thickening of the Schwann cells’ basement membrane was present and some myelinated fibers showed thin myelin sheaths (Figure 1). The endoneural blood vessels ultrastructure was preserved in all nerves from the three experimental groups.

**Fascicle morphometry**

Table 2 shows the average values for fascicular morphometric parameters of the ADN from control, acute diabetic and treated acute diabetic rats. There were no differences between proximal and distal segments from nerves in the same experimental group. Despite the morphological observation that nerves from treated animals tended to be smaller compared to controls, no differences on the fascicular area were found between groups. Also, there were no differences between groups on the Schwann cells nuclei number and density. Myelinated fiber number was significantly smaller on insulin treated groups, on both segments, compared to controls and to diabetic animals, with no corresponding differences on the myelinated fiber density.

**Fiber morphometry**

The average area and diameter of the myelinated fibers and respective axons as well as the average myelin sheath area and G ratio for the ADN form the three experimental groups are shown in Table 3. Values from proximal and distal segments are shown. Myelinated axon area and diameter was significantly larger on treated animals, compared to both, controls and acute diabetic groups, on both segments. No other differences were observed between groups for the myelinated fibers morphometric parameters. The ADN average values for unmyelinated fiber number, density, diameter and the unmyelinated/myelinated fiber ratio are shown in Table 4 for proximal and distal segments of nerves from the three experimental groups. No significant differences were found for all parameters between groups. Also, either for myelinated or unmyelinated fibers morphometric parameters, no differences between proximal and distal segments were present in nerves from the same experimental group.

**Distribution histograms**

Histograms of size distribution of the unmyelinated fibers showed that fibers diameter ranged from 0.1 to 1.2 µm, with a peak at 0.4 µm on proximal segments for the three experimental groups (Figure 2). On the distal segments, fibers also ranged from 0.1 to 1.2 µm but while controls and insulin treated groups showed peaks at 0.4 µm, diabetic animals presented the distribution peak at 0.5 µm with a shift of the distribution to the right.

**Discussion**

Few literature reports deal with transmission electron microscopy studies of autonomic nerves in diabetes, particularly in early onset of the disease. We investigated the ADN, an important nerve related to cardiovascular reflexes known to be altered in diabetes. Important results from this study include early Schwann cell involvement in diabetes with severe mitochondrial damage. Also, insulin treatment resulted in thickening of the Schwann cells basement membrane.

Alterations on the nerve extracellular matrix in diabetic patients have been investigated and include thickening of the microvasculature.
basement membrane as well as thickening of the perineurial cell basement membrane [22-25]. Thickening of the Schwann cells basement membrane was less reported, either in humans [25] or in experimental diabetes [21]. The extracellular matrix in peripheral nerves gives mechanical support to the surrounding cells. It also participates on the regulation mechanisms and interactions mediated by molecules on the surface of the cells [25]. In this way, any structural change on the basement membranes might lead to cellular dysfunction in multiple ways. We are showing that thickening of the Schwann cells basement membrane occurs associated with the onset of insulin treatment. One possible interpretation for this finding is that the animals might have undergone through periods of hypoglycaemia during the treatment. The effects of insulin-induced hypoglycaemia on peripheral nerves were recently revised [26] and no similar findings to those in our study were described. Thus, it is more likely that the insulin is causing a direct effect on the Schwann cells metabolism and synthesis of surface molecules.

**Table 1:** Body weight and blood glucose level on the 3rd day after streptozotocin (STZ) or vehicle injection, and on the experimental day (15 days after injection), together with average mean arterial pressure (MAP) and heart rate (HR) on the experimental day. Data are expressed as mean ± standard error of mean (SEM).

|                        | Control - 3 days | STZ - 3 days | STZ+Ins - 3 days |
|------------------------|-----------------|--------------|-----------------|
| Body Weight (g)        | 203 ± 14        | 201 ± 10     | 205 ± 11        |
| Blood Glucose (mg/dL)  | 85 ± 6"         | 533 ± 27"    | 538 ± 52"       |
| MAP (mmHg)             | 538 ± 10        | 96 ± 8       | 315 ± 32        |
| HR (bpm)               | 291 ± 63        | 291 ± 63     | 301 ± 15        |

STZ = diabetic group; STZ+Ins = Diabetic group treated with insulin. * significant difference compared to control group at the same experimental time; † significant difference compared to STZ group at the same experimental time; ‡ significant difference compared to the same group at 3 days.

|                        | Proximal Segments | Distal Segments |
|------------------------|-------------------|-----------------|
| Fascicular Area (µm²)  | 869 ± 93          | 915 ± 144       |
| Myelinated Fiber Number| 74 ± 8            | 8 ± 1           |
| Myelinated Fiber Density (fiber/mm²) | 89790 ± 8763 | 75462 ± 9057 |
| Schwann Cell Nuclei Number | 8 ± 1      | 7 ± 1           |
| Schwann Cell Nuclei Density (nucleus/mm²) | 49558 ± 16999 | 7158 ± 500     |

|                        | Proximal Segments | Distal Segments |
|------------------------|-------------------|-----------------|
| Myelinated Fiber Area (µm²) | 4.24 ± 0.32      | 4.27 ± 0.20     |
| Myelinated Fiber Diameter (µm) | 1.92 ± 0.09     | 1.95 ± 0.06    |
| Myelinated Axon Area (µm²) | 1.40 ± 0.08      | 1.44 ± 0.10    |
| Myelinated Axon Diameter (µm) | 1.05 ± 0.04     | 1.08 ± 0.05    |
| Myelin Sheath Area (µm²) | 2.83 ± 0.26      | 2.93 ± 0.33    |
| G Ratio                | 0.55 ± 0.01      | 0.60 ± 0.02    |

* significant difference compared to control group; † significant difference compared to STZ group.

**Table 2:** Morphometric parameters of the proximal and distal fascicles of the aortic depressor nerve (ADN) in control, diabetic (STZ) and diabetic treated with insulin (STZ+Ins) groups. Data are expressed as mean ± standard error of mean (SEM).

|                        | Control | STZ | STZ+Ins |
|------------------------|---------|-----|---------|
| Fascicular Area (µm²)  | 871 ± 59 | 816 ± 158 | 868 ± 261 |
| Myelinated Fiber Number| 79 ± 8  | 61 ± 9  | 32 ± 12" |
| Myelinated Fiber Density (fiber/mm²) | 86473 ± 6897 | 77819 ± 5787 | 55277 ± 17385 |
| Schwann Cell Nuclei Number | 10 ± 1   | 9 ± 2   | 7 ± 1   |
| Schwann Cell Nuclei Density (nucleus/mm²) | 11302 ± 1302 | 11575 ± 2145 | 14435 ± 2784 |

* significant difference compared to control group; † significant difference compared to STZ group.

|                        | Proximal Segments | Distal Segments |
|------------------------|-------------------|-----------------|
| Myelinated Fiber Area (µm²) | 4.24 ± 0.32      | 4.27 ± 0.20     |
| Myelinated Fiber Diameter (µm) | 1.92 ± 0.09     | 1.95 ± 0.06    |
| Myelinated Axon Area (µm²) | 1.40 ± 0.08      | 1.44 ± 0.10    |
| Myelinated Axon Diameter (µm) | 1.05 ± 0.04     | 1.08 ± 0.05    |
| Myelin Sheath Area (µm²) | 2.83 ± 0.26      | 2.93 ± 0.33    |
| G Ratio                | 0.55 ± 0.01      | 0.60 ± 0.02    |

* significant difference compared to control group; † significant difference compared to STZ group.

**Table 3:** Morphometric parameters of the proximal and distal segments of the aortic depressor nerve (ADN) myelinated fibers in control, diabetic (STZ) and diabetic treated with insulin (STZ+Ins) groups. Data are expressed as mean ± standard error of mean (SEM).
Table 4: Morphometric parameters of the proximal and distal segments of the aortic depressor nerve (ADN) unmyelinated fibers in control, diabetic (STZ) and diabetic treated with insulin (STZ+Ins) groups. Data are expressed as mean ± standard error of mean (SEM).

|                           | Control          | STZ              | STZ+Ins         |
|---------------------------|------------------|------------------|-----------------|
| **Proximal Segments**     |                  |                  |                 |
| Unmyelinated Fiber Number | 269 ± 51         | 330 ± 97         | 110 ± 63        |
| Unmyelinated Fiber Density (Fiber/mm²) | 301x10³ ± 49x10³ | 389x10³ ± 112x10³ | 183x10³ ± 88x10³ |
| Unmyelinated Fiber Diameter (µm) | 0.44 ± 0.04     | 0.45 ± 0.03      | 0.43 ± 0.01     |
| Unmyelinated/Myelinated Fiber Ratio | 4 ± 1           | 7 ± 3            | 7 ± 3           |
| **Distal Segments**       |                  |                  |                 |
| Unmyelinated Fiber Number | 213 ± 41         | 261 ± 79         | 144 ± 65        |
| Unmyelinated Fiber Density (Fiber/mm²) | 227x10³ ± 55x10³ | 334x10³ ± 86x10³ | 196x10³ ± 87x10³ |
| Unmyelinated Fiber Diameter (µm) | 0.43 ± 0.03     | 0.50 ± 0.03      | 0.41 ± 0.04     |
| Unmyelinated/Myelinated Fiber Ratio | 3 ± 1           | 4 ± 1            | 9 ± 5           |

* significant difference compared to control group; # significant difference compared to STZ group

Small fiber neuropathy in diabetic patients is commonly assessed by intraepidermal fiber quantification [30] while transmission electron microscopy studies are rarely performed. In experimental models, a reduction of the myelinated fiber and axon sizes is often described for somatic nerves either by light [31,32] or transmission electron microscopy studies [33]. For the autonomic nerves, small myelinated fibers loss was shown in STZ-model of diabetes [21], suggesting that somatic and autonomic nerves might react in distinct ways to diabetes. We showed that the onset of insulin treatment caused a reduction of the small myelinated fiber number. Since the axon size was significantly larger on insulin treated animals, it is suggestive that the remaining myelinated fibers are the larger ones that shifted the average size to higher values.

The ultrastructure of the phrenic nerve in chronic experimental diabetes [20] showed no differences in number or density of the unmyelinated fibers, similar to our study, despite the important alterations on the endoneural blood vessels described by those authors. It has been suggested that small fiber alterations could be due to a poor vascularity of the endoneural space [20,34]. In our study, no morphological alterations were present on the endoneural blood vessels, pointing to the fact that the small myelinated fiber loss was not related to endoneural ischemia. Also, it is suggestive that Schwann cells are more sensitive to the insulin treatment and respond readily than the blood vessels cells, in thickening the basement membrane.

Figure 2: Size distribution histograms of the unmyelinated fibers in the aortic depressor nerve (ADN) from proximal (upper panel) and distal (lower panel) segments of the aortic depressor nerve (ADN) in control (black circle), diabetic (STZ; white circle) and diabetic treated with insulin (STZ + Ins; black triangle) groups. Insulin treated animals’ histograms present a narrow base and a higher peak compared to control and STZ animals. Also, STZ animals’ distribution was slightly shifted towards large fibers. No significant differences were observed between groups. Bars represent mean ± SEM.
Conclusion

Investigation of experimental models of diabetes in early or acute stage of the disease provide relevant information on the physiopathological mechanisms involved in the alterations present in patients with chronic diabetes and its complications. This information may lead to a better understanding of the disease itself and provide background for the development of new approaches to therapeutics. We suggest that insulin treatment alone might have a role on the morphological alterations present in diabetic neuropathy.

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