Changes in Vascular and Immune Cell Content of an LPS Treated Rat Olfactory Bulb PD Model, Measured by Fluorescence Deconvolution Microscopy

Roger J. Bick1*, Brian J. Poindexter1 and Marie-Francoise Doursout2

1Department of Pathology and Laboratory Medicine, McGovern Medical School, University of Texas Health Science Center, USA
2Department of Anesthesiology, McGovern Medical School, University of Texas Health Science Center, USA

Corresponding author: Roger J. Bick, Department of Pathology and Laboratory Medicine, McGovern Medical School, University of Texas Health Science Center, Houston, TX 77030, Tel: (713) 500-5406; E-mail: Roger.J.Bick@uth.tmc.edu

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Abstract

Objective: We have previously shown changes in protein, immune cell, nitric oxide a neurotrophic factor content and distribution in the olfactory bulb of an endotoxin-treated rat model, as all these factors have been shown to be involved in neurodegeneration. We expanded our studies by fluorescently imaging smooth muscle actin and endothelial nitric oxide synthase (eNOS), both markers of blood vessels, to investigate loss of vasculature content, as well as imaging locations and quantities of immune cells. The work was performed to shed further light on associations between vessel integrity and immune cell initiated endothelial disruption. Our goal was to demonstrate that cytokine production, NOS induction and immune cell increases, are likely part of the process that leads to a loss of olfaction and dopaminergic signaling and includes vascular perturbations.

Methods: Rats were sacrificed following lipopolysaccharide (LPS) treatment. Olfactory bulbs were harvested, sectioned from top to bottom to include the tract and sensory neurons, and probed for markers of inflammation. Inducible nitric oxide synthase (iNOS), neuronal nitric oxide synthase (nNOS), eNOS, interleukin-1 beta (IL-1β), TNF-α, interleukin-6, glial cell derived neurotrophic factor (GDNF) and circulating nitric oxide (NO) were imaged together with tagged macrophages, T-cells, B-cells and neutrophils.

Results: Serum NO levels indicated that an inflammatory episode had occurred, being significantly higher in treated animals, with tissue levels of NOS elevated for an extended period of time. Immune cell clusters were seen in a number of areas and the localization of NOS isomers suggests that they have divergent roles in neurodegeneration. For instance, eNOS was associated with blood vessels, iNOS with glial and matrix cells and nNOS with glial cells and neurons. T and B-cell numbers showed a sustained increase; neutrophil numbers rapidly increased then returned to baseline levels; macrophage numbers increased and remained high; LAMP positive cell numbers (NK-cells) increased and remained high; GDNF content increased; IL-6, TNF-α and IL-1β levels all rapidly increased, before dropping to untreated levels, while circulating, NO levels increased dramatically. Of interest, the images of vascular content, immune cell content, eNOS and smooth muscle actin, allowed us to show detrimental interactions between cells, factors and vessels. Our data show that the majority of the vessels were intact, though sections of interest were ‘extracted’ to reveal possible leaky areas. Specific sites of IL-6 positive lymphocyte clustering were noted around vessels, suggesting that interactions are occurring that lead to disruptions of blood vessel tunicae, allowing the internalization of circulating cells and subsequent cytokine-initiated endothelial cell death.

Conclusion: Our findings suggest that protective GDNF and eNOS, which maintains vascular tone, are possibly synthesized too late to combat cytokine initiated neuron damage, glial activation and chronic loss of vascular integrity.

Keywords: Neurodegeneration; Olfaction; Blood-brain-barrier; Cytokines; Nitric oxide

Introduction

Many factors play a role in chronic neurodegeneration which leads to Parkinson’s disease (PD), including inflammation [1], cytokine syntheses and release [1], protein recycling-ubiquitination problems [2], glial cell activation [3], loss of astrocyte support [4], neurotrophic factors [2], nitric oxide synthase [5] and mitochondrial dysfunction [2]. However, recent reports have shown that circulating cells [6] are also involved in disease progression, which coupled with vascular damage [7], points to both peripheral and innate factors and cell involvement in the ensuing demise of neurons.

Vascular endothelial cells are thought to be damaged via interactions with not only reactive oxygen species (ROS), but also with increased cytokine concentrations, NO and LPS [8,9]. As multiple factors have been reported associated with neurodegeneration, the suggestion is that some have a role in loss of blood vessel integrity and ensuing damage to the blood brain barrier (BBB). This ‘neurovascular unit’ (endothelial cells, astrocytes, neurons, pericytes and matrix)
[10-12]; is essential for the wellbeing of the brain and has become an area of major research regarding its role in neurodegenerative diseases, including a loss of tight junctions [13] and passage of toxic substances and circulating cells into the brain itself [14]. Thus, the disease process is multi-compartmentalized and requires therapies that do more than target the loss of dopamine [15]. For instance, therapies are needed to address functional alterations of the BBB in conjunction with elucidating the role of circulating T helper and B lymphocytes and alterations in leukocyte antigens. The immune system and associated inflammation and blood vessel integrity changes thus require focused research to elucidate new strategies [16].

Our studies have focused on the olfactory bulb as an early loss of olfaction in Parkinson's disease patients as a measurable change and could be the starting point to elucidate both the initiation of neurodegeneration and the subsequent chronic loss of neurons [17]. Therefore the ability of the olfactory bulb to maintain both vascular integrity and a functioning BBB becomes even more important due to intimate interactions with synaptic functions and α-synuclein [18]. As previously reported by Maon et al, α-synuclein changes have been reported as an early identifier of olfactory bulb and anterior olfactory nucleus related problems protein transmission through neurological and anatomic pathways [19].

Previous work in our lab has demonstrated the 'toxicity' of PD-CSF [20] including, the presence of non-resident cells and cytokine production in the olfactory bulb of an LPS rat model [21], specific localizations of PD-associated proteins and cytokines in LPS treated brain [22] and the induction of neurotrophins and nitric oxide [5]. We therefore continued our studies to further understand the progression of this chronic, debilitating disease, by correlating cytokine, protein and neurotrophic factor concentrations and localizations, together with immune cell numbers and a loss of vascular integrity. This focus was partly due to recent research into vascular integrity and blood vessel function during chronic inflammation [23-25]. Using fluorescent tags and deconvolution imaging we examined vascular content, vessel integrity and vessel types in images of LPS treated rat olfactory bulbs and concluded that, by and large, vessels are overall intact but with 'suspect' areas. These findings might indicate that even though protective GDNF is produced and nitric oxide is released to maintain vascular tone, subsequent inflammatory episodes are increasingly challenging due to the first assault making repair and rescue difficult due to a loss of BBB control, continuing increases in cytokine levels and reductions in normal cellular abilities [5].

Materials and Methods

i. Sectioning and staining of tissues

Adult rat brains (300 g male Sprague Dawley), from saline injected and LPS treated animals (i.e., 35 μg/kg), were removed and olfactory bulbs separated, fixed, sectioned (12 μm 3) and probed with antibodies and treated with secondary antibodies for deconvolution fluorescence microscopy. Images were acquired in a stepwise process (0.1-0.25 μm steps) following excision of the guidelines of Paxinos and Watson [26]. The protocol for this work was approved by the animal welfare committee (AWC-14-0148 Title: Long-Term Effects of LPS in Rats: A Model of Neuro-Degenerative Disease).

ii. Imaging

Sections were mounted on 25 mm glass cover slips and fluoroscically probed for the three NOS isofoms, iNOS, nNOS and eNOS. Tissue was rinsed with cold phosphate buffered saline (PBS) and fixed with 3.7% formaldehyde (1008B, Tousimis Research Corp., Rockville, MD). Fixation was halted by 3 rinses with PBS at room temperature and blocked with 10% goat serum, then incubated for 1 h at 37°C with antibodies (1:100 to 1:1000 dilutions). Sections were further incubated for 30 min at 37°C and rinsed 3 times with 0.05% Tween 20 in PBS (TBBS). Nuclear and F-actin were stained with phalloidin (Invitrogen, Grand Island, NY, USA) diluted 1:100 in 4',6-diamidino-2-phenylindole (DAPI; A-T rich region binding in DNA) solution. Specimens were covered with one drop of anti-fade (Elvanol, DuPont, Wilmington, DE, USA). Tissue sections were probed with antibodies for glial cell derived neutrophic factor (GDNF; sc-9010, Santa Cruz Biotech, CA, USA) and lysosomal associated membrane protein (LAMP-1; sc-19992; Santa Cruz Biotech, CD107a, a marker for lymphocytic cells), a CD3 antibody (sc-20047, Santa Cruz Biotech; T-cell marker), a CD20 antibody (Polyclonal B-Cell marker, sc 15361, Santa Cruz; B-lymphocyte marker), a CD44 monoclonal neutrophil marker (sc-9960, Santa Cruz Biotech) and a CD68 polyclonal macrophage marker, (sc-9139, Santa Cruz Biotech). Secondary fluorescent antibodies were purchased from Jackson ImmunoResearch (West Grove, PA, USA). Fluorescence intensity measurements were made using pixel counts of RGB deconvoluted images, as previously described [22,27,28] and 3D reconstructions and rotations were made [28]. NOS antibodies were from Abcam (Cambridge, MA, USA; nNOS-ab5586, iNOS-ab3523; eNOS-M221).

iii. Deconvolution fluorescence microscopy

3D image reconstructions of NOS isofoms, actin and nuclei were generated [5,21,29] and images were saved as TIFF files. Comparative statistical analyses (one way ANOVA) of fluorescence densities (pixel numbers) were made using Corel (Ottawa, Ontario, Canada) and SigmaStat software (SPSS, Chicago, IL, USA).

iv. Circulating cytokine and nitric oxide level determinations

Blood samples from LPS treated and control animals were assayed for plasma cytokines using a commercial, enzyme-linked immunosorbent assay ELISA kit (R&D Systems, MN, USA). Plasma NO levels were quantified by a modified chemiluminescence method [30] using a Sievers NO analyzer 270B (GE Analytical, CO, USA). Nitrite concentrations were quantified by comparing measured versus sodium nitrate standards (20-400 pmol).

Results

Some of the results discussed are from previous reports, but are crucial to constructing a global picture of what we believe is occurring in the early stages of neurodegeneration and with vascular integrity.

Cytokine and immune cell content in olfactory bulb

Measurements of cytokine and immune cell content in the olfactory bulb are shown in Figure 1. A rapid, and sustained, rise in T-cell, B-cell and macrophage content, as well as small increases in the number of neutrophils, were observed. IL-6 and TNF-α, which are frequently associated with inflammation and cell death pathways, were increased over the first 12 h post treatment and remained elevated for at least 48
The data indicate that an inflammatory episode has taken place, and justifies this model for our studies and identifies cytokines and cells involved in the process.

Figure 1: Changes in olfactory bulb levels of cytokines, inflammation-associated cells and circulating nitric oxide levels, during the first 12 h post-LPS treatment: Blue stars reveal that there are significant increases (p<0.01) in cytokine levels and cell number(s) when compared to time zero, with red stars denoting a significant decrease (p<0.05) compared to zero time levels for neutrophils and TNF-α (n=8; 8 controls and 8 treated; samples were frozen as time dictated). Maximum level is the highest level measured during the first 12 h post-treatment.

Imaging of immune cell content in olfactory bulb

Increases in immune cell numbers were noted as shown in low magnification images (Figure 2), visualizing T cells, B cells and macrophage numbers over the first 6 h. These findings suggest the peak response time of the acute phase of the inflammation, when enough GDNF is present to overcome at least some of the endotoxin assault, cytokine-initiated cell death and glial cell activation [5]. Of note, is the reduction in the number of macrophages and neutrophils after an initial increase, while B-cell numbers remain high at 6 h and T-cell content continues to increase. Figure 2 shows immune system cell content in the olfactory bulb illustrated in the colored panels (all the probes are visualized) and in black and white split screen images of fluorescence of only the immune cell tag. The colored images contain green f-actin, blue nuclei and red for the designated cell type (B-cells, Macrophages, T-cells, and neutrophils, top to bottom).

Localization of immune cells using fluorescence imaging

Figure 3 shows representative images from olfactory bulb sections. Saline-treated (Panels A-D) is compared to LPS-treated tissues (Panels E-H), and focuses on the localization of cell types. Note in Panels A and E the vessel-associated ‘cuff’ of T-cells, with increased numbers in the treated sample (E). While in panels B and F, the white arrows point to B-cells and neutrophils (Panels C and G) revealing a specificity of being located along nuclei tracts. Macrophages are obviously increased in treated v control (H v D) at 6 h, but do have a homogenous distribution. The findings strongly implicate the interaction of T-cells and vascular endothelium, as well as some the occurrence of some cell death and/or the presence of non-resident as supported by the large number of macrophages in the tissue.

Images to detail comparison and quantification of vascular content

Figure 4 compares two random sections (A-saline-treated, C-LPS treated) to estimate vascular content by fluorescence pixel numbers. Values show content using single channels from RGB images following channel splitting. A count of 3 areas from each of 16 treated animal olfactory bulb sections revealed no significant difference in vessel content, specifically with eNOS (red tag; green=f-actin, blue=nuclei). Similar results when smooth muscle actin was imaged. As an example
of fluorescence imaging. Figure 5 shows an arteriole, once again with factin (green), nuclei (blue) and smooth muscle actin (red). The yellow and light greens are due to red and blue tags overlapping/merging, while the small black and white images show the channels individually. Note the capillary plexus formation above the main vessel. There is thus a suggestion that angiogenesis has occurred, probably not due to LPS treatment, and that loss of vasculature can be overcome for a short period of time.

Figure 4: Fluorescent imaging of blood vessels: Comparison of two sections (A-untreated, C-treated) probed for vascular content. Images A and C show probes for f-actin (green), nuclei (blue) and smooth muscle actin (red). Values are the content of smooth muscle in split channel renditions. Pixel densities in white boxes reveal that there are no significant difference in smooth muscle content in olfactory bulb sections following LPS treatment (57 measurements total were performed; Mag × 900).

Figure 5: Fluorescent image or arteriole and associated capillary plexus: High power image to highlight an arteriole with f-actin (green), nuclei (blue) and smooth muscle actin (red and yellow). The yellow and light greens are due to red and blue tags together (A). The three, smaller black and white images (a, b, c) show the channels individually. Note the appearance of a capillary plexus formation above the main vessel (Mag. × 1200).

High Power images to indicate possible endothelial discontinuity

Figure 6 is an image of an arteriole from an LPS-treated olfactory bulb, tagged with a Texas Red antibody to eNOS. A section has been cut out to detail possible areas of loss of endothelial integrity and likely structural damage. Image magnifications are shown in individual panels (Mag. × 900).

Discussion

Studies in our lab with cultured microglia, cultured astrocytes and LPS-treated rats, have suggested that the chronic nature of idiopathic PD might be due to a gradual process associated with protein folding, increasing exposure to high cytokine levels, perivascular immune cell invasion, a breached blood barrier and a loss of astrocyte adhesion, is eventually overwhelmed. The LPS model has been used by us and other researchers [31-33] being a naturally occurring substance, as opposed to MPTP and rotenone for example, and has produced a robust inflammatory response in the olfactory pathways. The endotoxin model has also been shown to elicit effects in the olfactory bulb, with subsequent loss of neuronal function leading to full-blown degradation, which includes detrimental astrocyte function, reduced phagocytosis and reduced neurotrophic factor (NGF, GDNF, BDNF) synthesis and release.

While it is now accepted that PD is a results of an inflammatory episode and subsequent microglia activation, increasing loss of functioning dopaminergic neurons [34], synthesis and release of multiple cytokines, accumulation of toxic substances, subcellular organelle dysfunction and increased effects of invading cells, a loss of BBB integrity, vascular endothelium integrity and astrocyte damage, are strongly implicated in a continuing loss of protective abilities [7,35-39].

Fluorescence imaging has yielded a more detailed insight into explaining the unremitting progression to loss of function and quality of life for the patient as well as for loved ones. Overall, our findings
suggest the recognition of a potential time point from our work with the olfactory bulb and the idea that loss of olfaction is an early-enough marker in neurodegeneration to recognize that the initiation of strategies which will overcome a continuing loss of cognitive abilities will need to be considered and implemented [40-42].

Multiple ‘non-resident’ cell types are present in the olfactory bulb and their number increases following LPS treatment, with co-localization to specific areas and structures. These increases become important in understanding the rapid events leading to cell destruction and loss of normal function. For example, neutrophils which are present in increasing numbers in the first 6 h post LPS treatment in our experiments, are known to be both protective and damaging [36]. The same can be said for astrocytes, with ‘misplacement’ of the cells in PD and Alzheimer’s disease and their release of deadly molecules [39]. Thus, protective cells can become harmful during inflammatory episodes and, coupled with an influx of immune cells, causes disruption of the BBB which is exacerbated by usually beneficial astrocytes that are mechanistically altered and interact with invading cells [37-40]. Indeed, our previous research has shown cuffs of T-cells around vessels after endotoxin treatment, as well as ubiquitin and TNFα in vascular endothelial cells, suggesting that perivascular disruption occurs and that cells respond in both self-preservation and destructive manners [5,43]. Previous results also indicated that astrocytes, which are multi-functional glia cells and an integral part of the BBB, synthesize peripherally localized α-synuclein, suggesting that the protein has adhesive properties, and the adhesion is disrupted via glial activation causing a loss of astrocytic support of vascular endothelial cells and an ensuing, leaky BBB [20]. Imaging the localizations of diverse immune cells as there is increasing evidence for their involvement in neurodegeneration, loss of blood vessel integrity and age-related problems [44,45], suggests roles for both circulating-invading cells as well as resident cells.

In these experiments in our study, we did not find a decrease in vasculature content after LPS treatment although there was some evidence for loss of vessel integrity suggested by a lack of continuity of the tagged smooth muscle and associated endothelium. Increases in NOS isoforms (not shown; Ref 5) suggested that they might each have a different role in neurodegeneration. NOS induction, increased levels of protective cytokines and GDNF synthesis, might be an adequate, initial, group of protective mechanisms. However, with chronic inflammation due to either a massive single event or repetitive/additive episodes, the mechanisms are overcome, with ensuing, sustained microglial activation, prolonged and continuous protein misfolding, the tagged smooth muscle and associated endothelium. Increases in evidence for loss of vessel integrity suggested by a lack of continuity of the vasculature content around vessels [30].


