Temporal changes in inflammatory mitochondria-enriched microRNAs following traumatic brain injury and effects of miR-146a nanoparticle delivery

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Abstract

MicroRNAs (miRNAs) are small non-coding RNA molecules that regulate post-transcriptional gene expression and contribute to all aspects of cellular function. We previously reported that the activities of several mitochondria-enriched miRNAs regulating inflammation (i.e., miR-142-3p, miR-142-5p, and miR-146a) are altered in the hippocampus at 3–12 hours following a severe traumatic brain injury. In the present study, we investigated the temporal expression profile of these inflammatory miRNAs in mitochondria and cytosol fractions at more chronic post-injury times following severe controlled cortical impact injury in rats. In addition, several inflammatory genes were analyzed in the cytosol fractions. The analysis showed that while elevated levels were observed in cytoplasm, the mitochondria-enriched miRNAs, miR-142-3p and miR-142-5p continued to be significantly reduced in mitochondria from injured hippocampi for at least 3 days and returned to near normal levels at 7 days post-injury. Although not statistically significant, miR-146a also remained at reduced levels for up to 3 days following controlled cortical impact injury, and recovered by 7 days. In contrast, miRNAs that are not enriched in mitochondria, including miR-124a, miR-150, miR-19b, miR-155, and miR-223 were either increased or demonstrated no change in their levels in mitochondrial fractions for 7 days. The one exception was that miR-223 levels were reduced in mitochondria at 1 day following injury. No major alterations were observed in sham operated animals. This temporal pattern was unique to mitochondria-enriched miRNAs and correlated with injury-induced changes in mitochondrial bioenergetics as well as expression levels of several inflammatory markers. These observations suggested a potential compartmental re-distribution of the mitochondria-enriched inflammatory miRNAs and may reflect an intracellular mechanism by which specific miRNAs regulate injury-induced inflammatory signaling. To test this, we utilized a novel peptide-based nanoparticle strategy for in vitro and in vivo delivery of a miR-146a mimic as a potential therapeutic strategy for targeting nuclear factor-kappaB inflammatory modulators in the injured brain. Nanoparticle delivery of miR-146a to BV-2 or SH-SY5Y cells significantly reduced expression of TNF receptor-associated factor 6 (TRAF6) and interleukin-1 receptor-associated kinase 1 (IRAK1), two important modulators of the nuclear factor-kappaB (NF-κB) pro-inflammatory pathway. Moreover, injections of miR-146a containing nanoparticles into the brain immediately following controlled cortical impact injury significantly reduced hippocampal TNF receptor-associated factor 6 and interleukin-1 receptor-associated kinase 1 levels. Taken together, our studies demonstrate the subcellular alteration of inflammatory miRNAs after traumatic brain injury and establish proof of principle that nanoparticle delivery of miR-146a has therapeutic potential for modulating pro-inflammatory effectors in the injured brain. All of the studies performed were approved by the University of Kentucky Institutional Animal Care and Usage Committee (IACUC protocol # 2014-1300) on August 17, 2017.

Key Words: cell permeable peptide-delivery; controlled cortical impact; inflammatory pathway; mitochondria-associated microRNA; nanoparticle; nuclear factor-kappaB; traumatic brain injury

Introduction

Traumatic brain injury (TBI) elicits a rapid and pronounced neuroinflammatory response propagated, in part, by resident microglia as well as circulating monocytes that differentiate into macrophages along a continuum ranging from cytotoxic pro-inflammatory to reparative anti-inflammatory response.

Funding: This work is supported by a grant (15-12A) from the Kentucky Spinal Cord and Head Injury Research Trust to JES and WWX.

How to cite this article: Wang WX, Prajapati P, Vekaria HJ, Spry M, Cloud AL, Sullivan PG, Springer JE (2021) Temporal changes in inflammatory mitochondria-enriched microRNAs following traumatic brain injury and effects of miR-146a nanoparticle delivery. Neural Regen Reg 16(3):514-522.
phenotypes. These cells have been consistently linked with pro-inflammatory signaling processes underlying chronic degenerative changes after TBI (Ziebell and Morganti-Kossman, 2010; Kumar and Loane, 2012; Woodcock and Morganti-Kossman, 2013). The expression of reparative myeloid cells peaks at approximately one week following injury and then declines, while the pro-inflammatory phenotype may persist for months to years (Ziebell and Morganti-Kossman, 2010; Kumar and Loane, 2012; Johnson et al., 2013; Woodcock and Morganti-Kossman, 2013; Kumar et al., 2015). This differential expression pattern favoring prolonged pro-inflammation limits tissue repair and is associated with poor functional outcomes. Therefore, strategies that effectively limit detrimental pro-inflammatory responses, while maintaining the reparative potential of macrophages, have clear therapeutic potential (Ziebell and Morganti-Kossman, 2010; Kumar and Loane, 2012; Woodcock and Morganti-Kossman, 2013). In support of this, experimental approaches that mitigate pro-inflammatory signaling after severe TBI can significantly reduce tissue pathology and promote functional recovery (Lloyd et al., 2008; Hsieh et al., 2014; Bachstetter et al., 2015; Morganti et al., 2015). However, identifying tools that shift these myeloid effectors towards a reparative bias with a high degree of specificity and efficacy has been very limited. One novel strategy with high translational potential is the use of nanoparticle technologies to deliver microRNAs (miRNAs) targeting genes that regulate inflammatory signaling (O’Neill et al., 2011; Contreras and Rao, 2012; O’Connell et al., 2012).

MiRNAs are evolutionarily conserved non-coding short RNA molecules that bind recognition elements in messenger RNA (mRNA) and fine tune post-transcriptional gene expression (Bartel, 2004; Huntzinger and Izaurrealde, 2011). MiRNAs regulate all aspects of cellular function, and disruption of miRNA activity contributes to many disease states including neurodegeneration (Calin and Croce, 2006; Nelson et al., 2008; Liu and Xu, 2011; Hata, 2013). MiRNA-based therapeutic approaches have been tested and several are currently in clinical trials targeting hepatitis C, type 2 diabetes, and several cancer types (Rupaimoole and Slack, 2017). Therefore, the ability of miRNAs to regulate expression of genes contributing to TBI-related neuroinflammatory signaling makes them attractive therapeutic candidates. Using mouse and rat models, we and others have reported that a number of miRNAs are altered in cortex and hippocampus following TBI (Redell et al., 2009; Hu et al., 2012; Liu et al., 2014; Sabirzhanov et al., 2014; Wang et al., 2015, 2017; Harrison et al., 2017; Kumar et al., 2017). The studies from our lab are the first to demonstrate that a subset of miRNAs regulating cellular inflammatory activities (e.g., miR-146a) is enriched in brain mitochondria fractions (Wang et al., 2015, 2017). We also reported that levels of these miRNAs display a compartmental shift from the mitochondria to the cytosol as early as 3–12 hours following TBI (Wang et al., 2015, 2017), which corresponds to the temporal loss in mitochondrial function after acute TBI (Sullivan et al., 1998, 2007; Singh et al., 2006). These observations suggest an interaction between mitochondrial function and miRNA-mediated inflammatory signaling at early time points following TBI. MiR-146a plays a significant role in regulating levels of inflammatory cytokines and related signaling events in a manner that favors expression of the reparative macrophage phenotype (Cui and Liu, 2015). However, the relationship between miRNA changes and inflammatory signaling in the hippocampus beyond 12 hours following TBI is not known. Moreover, strategies that modify miRNA signaling to effectively promote expression of the reparative macrophage phenotype are lacking. Therefore, the aim of this study was to quantify changes of select miRNAs and inflammatory genes isolated from the hippocampus at different time points following severe TBI. In addition, we tested the ability of miR-146a nanoparticle delivery to target the nuclear factor-kappaB (NF-κB) modulators TNF receptor-associated factor 6 (TRAF6) and interleukin-1 receptor-associated kinase 1 (IRAK1) in vitro as well as in vivo after severe TBI.

Materials and Methods

Animals and TBI procedures

All of the studies performed were approved by the University of Kentucky Institutional Animal Care and Usage Committee (IAUC protocol # 2014-1300) on August 17, 2017. Additionally, the Division of Laboratory Animal Resources at the University is accredited by the Association for the Assessment and Accreditation for Laboratory Animal Care, International (AAALAC, International) and all experiments were carried out in accordance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals (NIH Publication No. 85-23, revised 1996). Thirty young adult male (7–10 weeks old) Sprague-Dawley rats weighing ~250 g (Charles River Laboratories, Wilmington, MA, USA) were acclimated for 7 days prior to experimentation. Rats were allowed free access to food/water and housed in a temperature-controlled room on a 12-hour light/dark cycle. The surgical and brain injury procedures performed in this study followed our previously published protocols (Sullivan et al., 1999; Wang et al., 2015). The controlled cortical impact (CCI) injury is a well-established brain injury model utilized by numerous researchers and, as such, allows for comparison of outcomes across studies (Dixon et al., 1991). Briefly, experimental rats were anesthetized with a dose of 4% isoflurane (Covetech, Portland, ME, USA) via a nose cone and were positioned in a stereotaxic frame (David Kopf, Tujunga, CA, USA). The surgical and CCI procedure was conducted under sterile conditions and continued supplied dose of 2.5% isoflurane throughout the procedure. The core body temperature of the experimental rats was kept at 37°C during the procedures and the post-surgical/CCI recovery period. An incision was made along the midline in a rostro-caudal orientation to expose the skull. A 6-mm craniotomy was then created lateral to center between bregma and lambda. The brain injury was performed by driving a 5 mm diameter rod tip at 3.5 m/s velocity to compress the cortex to a depth of 2.0 mm using a pneumatically controlled impacting device (Precision Systems and Instrumentation, Fairfax Station, VA, USA). This procedure resulted in a severe injury consistent with previous studies (Baldwin et al., 1997; Davis et al., 2008; Gilmer et al., 2009) and characterized by physical damage to the cortex and the presence of neuronal damage to the underlying hippocampus. Following injury, surgical absorbable hemostat was placed in the craniotomy and a plastic disc adhered to the skull to cover and stabilize the craniotomy site. Sham animals received identical exposure to the surgical procedures, but without brain injury. The surgeries were conducted at the same time of day, 2 days apart with three CCI and two sham operations performed at each time. Animals (total n = 6 per group for CCI, n = 4 per group for sham surgery) were euthanatized with CO₂ at 1, 3, or 7 days following injury and the ipsilateral and contralateral hippocampi were rapidly removed for isolation of the mitochondria and cytosolic fractions.

Cytosolic and mitochondrial fractionation

For mitochondrial bioenergetics and the miRNA and mRNA
expression studies, the mitochondria and cytosol fractions were obtained from the hippocampus following a previously described Ficoll density gradient procedure (Pandya et al., 2009). Briefly, harvested hippocampal tissue was homogenized in pre-cooled mitochondrial isolation buffer (215 mM mannitol, 75 mM sucrose, 0.1% bovine serum albumin (BSA), 20 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES), 1 mM ethylene glycol-bis(β-aminoethly) ether)-N,N,N',N'-tetraacetic acid (EGTA); pH adjusted to 7.2 with KOH). The homogenate was then centrifuged at 1300 x g for 3 minutes to remove tissue debris and nuclei. The resulting supernatant was then subjected to a centrifugation at 13,000 x g for 10 minutes to obtain crude mitochondrial pellets and the supernatant was saved as cytosol (CT) fraction. The resuspended crude mitochondria sample was then placed into a nitrogen cell disruption chamber for 10 minutes (1200 psi) to rupture and release synaptosomal mitochondria (Brown et al., 2004). The crude mitochondria fraction was then laid over a discontinuous Ficoll gradient (7.5% + 10%) and further separated by centrifugation using a SW 55 Ti rotors at 100,000 x g for 30 minutes. The mitochondria pellet (MT) was rinsed in mitochondrial isolation buffer in the absence of EGTA and centrifuged at 10,000 x g at 4°C for 10 minutes and resuspended in the same buffer to achieve a concentration of ~1.0 mg/mL. The protein content of the various fractions was analyzed using BCA protein assay kit (ThermoFisher, Waltham, MA USA).

Mitochondrial bioenergetics
Mitochondrial bioenergetics were assessed using the Seahorse Biosciences XFe24 Flux Analyzer (Agilent Technologies [Santa Clara, CA]; RRID:SCR_013575) using a procedure developed in our lab (Sauerbeck et al., 2011; Hubbard et al., 2016; Pandya et al., 2016). In these experiments, rotenone, oligomycin, and trifluoroethoxy carbonyl cyanide phenylhydrazone, 2-[4-[(trifluoromethoxy)phenyl] hydrazinylidene]-propanedioinitril (FFCP) were purchased from Biomol (Plymouth Meeting, PA, USA). All other reagents were purchased from Sigma-Aldrich (St. Louis, MO, USA). Briefly, 3.5 µg protein equivalent of isolated mitochondria were suspended in respiration buffer (RB) (125 mM KCl, 2 mM MgCl₂, 0.1% BSA, 20 mM HEPES and 2.5 mM KH₂PO₄, adjusted pH 7.2). The combination of substrates/inhibitors/uncoupler were prepared in RB and 75 µL of each injected as pyruvate + malate + adenosine diphosphate (ADP), oligomycin, FCCP, and rotenone + succinate sequentially through ports A→D at the final concentrations of 5 mM pyruvate, 2.5 mM malate, 1 mM ADP, 1 µg/mL oligomycin, 4 µM FCCP, 0.1 µM rotenone, and 10 mM succinate. ADP driven complex I mediated coupled respiration (State III), ATPase locked leaky respiration (State IV), complex I driven maximum uncoupled respiration (State Vₒ) and complex II-driven maximum respiration (State Vₒₒ) were determined with each subsequent injection. The results are presented as oxygen consumption rate (OCR) and expressed as pmol of oxygen consumed per minute.

TaqMan single-tube miRNA RT-qPCR
Isolation of total hippocampal tissue RNA from the cytosolic and mitochondrial fractions and TaqMan single-tube RT-qPCR (ThermoFisher) of select miRNA were performed as previously reported (Wang et al., 2008, 2015). Selected miRNAs for analysis included miR-146a (Assay ID: 000468), miR-142-3p (Assay ID: 000464), miR-142-5p (Assay ID: 002248), miR-155 (Assay ID: 002571), miR-150 (Assay ID: 000473), miR-223 (Assay ID: 002248), miR-146a (Assay ID: 001182), and miR-19b (Assay ID: 000396). Each miRNA analysis was performed in technical triplicates of either contralateral or ipsilateral hippocampi for a total of 6 TBI and 4 sham biological replicates from each time point (n = 6 TBI, n = 4 sham). The raw cycle threshold (CT) values were determined using QuantStudio™ Real-Time PCR Software with automatic baseline and threshold (ThermoFisher). Overall miRNA levels following TBI or sham surgery in the cytoplasmic and mitochondrial fractions at each time point were calculated using formula RQ = 2 –([Ct(ipsi)–Ct(Contra)] / ([Ct(Contra)] – [Ct(Contra)])) (Table 1) comparing the ipsilateral injured side or sham surgery side relative to the uninjured contralateral side or no surgery side (RQ: relative quantity). To evaluate the net change of miRNA enrichment in mitochondria, the following formula was used: RQ = 2^[ΔCt(Mito)–ΔCt(Cyto)] comparing the mitochondria fraction to cytosolic fraction of the same brain side. The calculation method followed the instructions detailed in the “Guide to Performing Relative Quantitation of Gene Expression Using Real-Time Quantitative PCR” (Applied Biosystems).

Gene expression analysis using TaqMan low density array panel
A customized TaqMan low density array (TLDA) qRT-PCR panel in an 8 x 48 gene array format (ThermoFisher) was employed to analyze gene expression in cytosolic fractions from the injured ipsilateral hippocampi relative to the uninjured contralateral side. Forty-seven genes including a group of inflammatory-related marker genes (Gensel et al., 2017) and an internal control ‘house-keeping’ gene (18S) were selected for the custom array. The list of genes and their assay IDs are provided in Additional Table 1. Total cytosolic RNA (500 ng) was subjected to first-strand cDNA synthesis using SuperScript™ III First-Strand Synthesis System (ThermoFisher). The resulting cDNA from each sample was then combined with TaqMan Universal PCR Master Mix (No AmpErase UNG, ThermoFisher) and loaded into each port of the TLDA cards following manufacturer’s instructions. Quantitative real-time PCR was performed with a QuantStudio™ 7 Real-Time PCR System (ThermoFisher) using the manufacturer’s standard program (hold 2 minutes at 50°C, followed by 10 minutes at 95°C, then 40 cycles of 15 seconds at 95°C and 1 minute at 60°C).

TLDA data processing
Raw qPCR CT (cycle threshold) values from the TLDA card were first determined using QuantStudio™ Real-Time PCR Software (ThermoFisher) with automatic baseline and threshold. PCR data that failed software-defined QC and those CT values equal to or more than 35 were considered as undetectable and were given a value of 35. The CT value of 18S ribosomal RNA was used to obtain ∆CT values of a given sample and the whole dataset was then further normalised using the Global Mean Normalization method (Mestdagh et al., 2009) to minimize the technical variability among array sets. The comparative ∆∆CT method was used to calculate relative gene expression levels. Mean ∆∆CT values and standard deviations were calculated following the “Guide to Performing Relative Quantitation of Gene Expression Using Real-Time Quantitative PCR” (Applied Biosystems (2008)). The effect of TBI on each gene in the TLDA card was expressed as fold change in cytosolic fractions from the ipsilateral hippocampus relative to uninjured contralateral side.

Cell permeable peptide-delivery of miRNA mimic in cell cultures
Cell culture experiments were conducted to document the ability of a mellitin-derived cell permeable peptide (p5RHH) to introduce Cy5-labeled miR-146a or scrambled control (Integrated DNA Technologies, Skokie, IL, USA) to BV-2 (kind gift from Dr. Linda van Eldik of University of Kentucky) and SH-SY5Y (Cat# CRL-2266, ATCC, Manassas, VA, USA, RRID:CVCL_0019) cells. The p5RHH peptide...
Levels of miRNAs were calculated using the formula: fold change = 2–(Ct(Ipsi)–Ct(Contra)) comparing the ipsilateral injured/sham-operated side of the brain relative to that of the uninjured/no operated contralateral side of the same animal. Data represent the mean value of the six biological replicates for traumatic brain injury (TBI) and four biological replicates for sham, and the P-values were generated using two-tailed Student’s t-test (GraphPad 8.0.2). The fold change range is shown in parentheses. Significantly different expressed miRNAs are highlighted in red (P < 0.05). 1, 3, or 7 days indicates the tissue harvesting time following controlled cortical impact or sham surgery.

(VLTLGLPALISWIRRRHRRHC) was synthesized by NeoScientific (Neobioslabs, Cambridge, MA, USA) and a stock of 10 mM was prepared in phosphate and nuclease free water, aliquoted and stored at –80°C. A total of 3 × 10⁵ cells were prepared using protease and nuclease free water, and the craniotomy site and overlying skin treated and closed with sutures. The animals were then euthanatized with CO₂. After 45 minutes, the brain was left in place for an additional 2 minutes, slowly withdrawn, and a WPI Ultramicropump III connected to a WPI MICRO2T SMARTouch controller (WPI, Sarasota, FL, USA). The needle was directed to the ventral hippocampus ipsilateral to the injury side using stereotaxic coordinates (Paxinos and Watson, 2015) and western blotting for TRAF6.

Protein isolation and western blot analysis

Protein isolation and Western blot analysis were performed using standard techniques routinely performed in our lab (Wang et al., 2015). Briefly, RIPA buffer (Cat# R0278; Sigma) was used to lyse proteins of rat hippocampal tissue, as well as total cellular protein in transfected cells. Protein concentrations were determined using BCA protein assay (Cath 23227, Pierce Biotechnology, Rockford, IL, USA). For Western blotting, equal amounts of protein (10–20 µg) were loaded on 4–16% Mini-PROTEAN® TGXTM gels (Cat# 456-1106, Bio-Rad, Hercules, CA, USA) and run at 0.03 A. The gels were then transblotted onto polyvinylidene fluoride membranes (Cat# 1620177, Bio-Rad) and non-specific binding sites blocked by incubating the membrane with 5% nonfat milk for 1–2 hours. The blots were then incubated overnight with primary antibody at 4°C. Primary antibodies used in this study included anti-TRAF6 (1:1000 dilution; rabbit, Cat# ab33915; Abcam, Cambridge, MA, USA) and anti-actin (1:2000, rabbit, Cat# A5060; Sigma). After 3–4 rinses following primary antibody incubation, blots were incubated with goat anti-rabbit peroxidase-conjugated secondary antibody (Jackson ImmunoResearch Laboratories, Inc. West Grove, Pennsylvania, USA) or IRDye® 800 goat anti-rabbit IgG (1:20,000, Cat# 925-68070, Li-COR, Lincoln, Nebraska, USA) for 1 hour at room temperature. The immunoreactive signals were visualized using SuperSignal™ West Pico PLUS Chemiluminescent Substrate (Pierce Biotechnology) or using Odyssey CLx (Li-COR, Lincoln).

Table 1  Changes in miRNA levels in cytoplasmic fractions of hippocampal tissue following TBI and in sham-operated control animals

| miRNA    | 1 d Fold (range) | 1 d P-value | 3 d Fold (range) | 3 d P-value | 7 d Fold (range) | 7 d P-value |
|----------|-----------------|-------------|-----------------|-------------|-----------------|-------------|
| TBI      | 0.93 (0.67–1.27) | 0.625       | 0.99 (0.87–1.13) | 0.919       | 1.03 (0.73–1.45) | 0.885       |
| miR-124  | 2.24 (1.61–3.11) | 0.001       | 3.49 (1.97–6.17) | 0.001       | 0.93 (0.70–1.24) | 0.886       |
| miR-142-3p | 2.62 (2.05–3.35) | 0.0001      | 3.41 (2.47–4.70) | 0.0002      | 0.94 (0.72–1.22) | 0.878       |
| miR-142-5p | 1.55 (0.98–2.47) | 0.148       | 1.93 (1.09–3.42) | 0.078       | 0.93 (0.54–1.58) | 0.852       |
| miR-150  | 0.87 (0.59–1.29) | 0.541       | 0.88 (0.66–1.19) | 0.678       | 0.96 (0.87–1.06) | 0.548       |
| miR-155  | 11.54 (6.3–21.3) | 0.00017     | 6.18 (3.92–9.72) | 0.020       | 1.15 (0.73–1.81) | 0.828       |
| miR-223  | 15.0 (9.9–22.8)  | 0.000002    | 3.92 (2.07–7.42) | 0.003       | 0.66 (0.32–1.35) | 0.538       |
| miR-19b  | 1.31 (1.06–1.63) | 0.072       | 1.33 (1.04–1.72) | 0.028       | 0.83 (0.70–0.97) | 0.318       |
| Sham     | 0.90 (0.70–1.17) | 0.711       | 0.86 (0.7–1.05) | 0.213       | 0.88 (0.27–1.77) | 0.482       |
| miR-142-3p | 0.89 (0.82–1.29) | 0.897       | 1.23 (0.86–1.76) | 0.529       | 1.50 (0.83–2.71) | 0.573       |
| miR-142-5p | 0.87 (0.70–1.07) | 0.485       | 1.28 (0.68–2.42) | 0.520       | 1.36 (0.67–2.76) | 0.701       |
| miR-146a | 1.05 (0.74–1.48) | 0.865       | 0.96 (0.47–1.98) | 0.941       | 1.76 (1.25–2.47) | 0.495       |
| miR-150  | 0.87 (0.73–1.04) | 0.581       | 0.91 (0.60–1.37) | 0.719       | 1.07 (0.96–1.19) | 0.740       |
| miR-155  | 1.89 (1.27–2.81) | 0.202       | 1.22 (0.44–3.40) | 0.735       | 3.04 (1.58–5.84) | 0.355       |
| miR-223  | 1.67 (1.15–2.43) | 0.095       | 1.61 (0.71–1.82) | 0.494       | 2.55 (1.82–3.57) | 0.357       |
| miR-19b  | 1.00 (0.87–1.15) | 0.999       | 0.85 (0.61–1.17) | 0.379       | 1.39 (1.00–1.94) | 0.372       |

Changes in miRNA levels in cytoplasmic fractions of hippocampal tissue following TBI and in sham-operated control animals
Results
Mitochondria bioenergetics following TBI
The post-injury bioenergetic profile of total mitochondria isolated from the hippocampus revealed significant changes in state-dependent oxygen consumption rates (OCR) over time (Figure 1). The OCR significantly decreased in all four tested respiration states ($P < 0.01$, for State III, State IVo, and State Vco, respectively; $P < 0.05$ for State IV) in mitochondria isolated from the ipsilateral hippocampus at 1 day following CCI relative to intact contralateral side (Figure 1A). At 3 days post CCI, OCR in mitochondria from the ipsilateral hippocampus continued to be significantly decreased in State IV and State Vco ($P < 0.05$), with near significant decreased levels in State III ($P = 0.058$) compared to the contralateral side. By 1 week post-injury, there were no significant differences in any respiration state in mitochondria from the injured hippocampus relative to contralateral side. Sham-surgery did not result in any significant changes in bioenergetics across all three time points (Figure 1B).

Alteration of the association of mitochondria-enriched miRNAs following TBI
We reported previously that levels of miR-146a, miR-142-3p, and miR-142-5p are highly enriched in the mitochondria compared to cytosol (Wang et al., 2015). We also found that the levels of these mitochondria-enriched miRNAs are decreased in the mitochondrial fraction at early times following TBI (3 and 12 hours), and that all three miRNAs increased in cytoplasmic fractions at the same time points (Wang et al., 2015, 2017). In the present study, we further investigated levels of these miRNAs at extended times up to 7 days following TBI (Figure 2). When miRNA levels in the mitochondrial fraction are normalized to that of cytoplasmic fraction, it provides a relative measure of mitochondrial enrichment. The enrichment of miR-142-3p and miR-142-5p significantly decreased ($P = 0.050$ and 0.002, respectively) at 1 day post TBI in mitochondria from the injured ipsilateral hippocampus compared to the uninjured contralateral hippocampus (Figure 2A). The mitochondrial enrichment of these two miRNAs remained significantly lower for up to 3 days ($P < 0.01$) and returned to near normal levels at 7 days post-injury. The levels of another mitochondria-enriched miRNA, miR-146a trended lower at 1 day in the mitochondrial fraction following TBI with a similar pattern being observed at 3 days. However, by 7 days post-injury, miR-146a levels in the mitochondrial fraction were found to be significantly elevated ($P = 0.033$; Figure 2A).

The levels of several non-mitochondria-enriched miRNAs also exhibited changes over time following TBI with different trends (Figure 2B). miR-124a, miR-150, miR19b and a known TBI-responsive miRNA miR-155 were either increased (such as miR-124a, miR-155, miR-19b) or not significantly altered (miR-150) in mitochondria at any time examined following TBI. One exception was miR-223 (Figure 2B), which is also a TBI-responsive miRNA that showed significantly decreased levels in mitochondria at 1 day after TBI ($P < 0.01$). However, by 7 days post-injury, the levels of miR-223 were significantly elevated ($P < 0.05$) in mitochondria isolated from the injured hippocampus. No major changes in any of the above mentioned miRNAs were observed between the ipsilateral and contralateral hippocampal mitochondria isolated from the group of sham operated animals (Additional Figure 1).

Interestingly, while levels of the mitochondria-enriched miRNAs miR-142-3p and miR-142-5p were reduced in mitochondria at 1 and 3 days following TBI, their levels in cytoplasmic fractions significantly increased ($P < 0.01$; Table 1). By 7 days following TBI, the levels of both miR-142-3p and miR-142-5p in the cytoplasmic fraction had returned to normal. A similar trending pattern was observed for miR-146a. Finally, levels of two TBI-responsive miRNAs, miRNA miR-155 and miR-223 were significantly elevated in the cytoplasm at 1 and 3 days post-injury and returned to normal levels by 7 days (see levels and $P$ values in Table 1).

Inflammatory gene expression at different time points following TBI
A customized TLDA was used to analyze the expression of selected inflammatory-related genes in hippocampal...
cytoplasm. In general, alterations in gene expression were found to be most active at 1 day following TBI (Table 2). By 3 days following TBI, the number and levels of genes significantly altered were less than that at 1 day, especially with respect to anti-inflammatory genes (Table 2). Among the tested genes, there was only one anti-inflammatory gene, Tgfb1, which remained elevated at 3 days post-injury. Among the pro-inflammatory genes examined, the chemokine (C-C motif) ligand 2 (CCL2, also called monocyte chemoattractant protein 1, MCP1) appeared to have the greatest change in expression with respect to anti-inflammatory genes (Table 2). Among the tested genes, there was only one anti-inflammatory gene, Tgfb1, which remained elevated at 3 days post-injury. Among the pro-inflammatory genes examined, the chemokine (C-C motif) ligand 2 (CCL2, also called monocyte chemoattractant protein 1, MCP1) appeared to have the greatest change in expression with respect to anti-inflammatory genes (Table 2). Among the tested genes, there was only one anti-inflammatory gene, Tgfb1, which remained elevated at 3 days post-injury. Among the pro-inflammatory genes examined, the chemokine (C-C motif) ligand 2 (CCL2, also called monocyte chemoattractant protein 1, MCP1) appeared to have the greatest change in expression with respect to anti-inflammatory genes (Table 2).

**Effect of miR-146a nanoparticle delivery to BV-2 and SH-SY5Y cells**

miR-146a targets TRAF6 and IRAK1, two effectors that are important regulators on the NF-κB pathway (Taganov et al., 2006). Given our previous observations demonstrating a differential expression of miR-146a at early hours following TBI (Wang et al., 2015, 2017), we selected this anti-inflammatory miRNA for the peptide-based nanoparticle delivery studies. We first tested the effects of the peptide/miR-146a nanoparticle delivery system using the mouse microglial BV-2 and human neuroblastoma SH-SY5Y cell line. These two cell lines were incubated with p5RHH + miR-146a-Cy5 mimic or scrambled control nanoparticles for 48 hours. Additional cell culture wells were incubated with miR-146a-Cy5 mimic (or scrambled control) without any carrier and we did not observe any significant differences in any of the other genes in the panel at 7 days between the ipsilateral injured and contralateral sides.

**Effect of miR-146a delivery to the injured rat hippocampus**

The p5RHH peptide was then used to deliver miR-146a mimic or scrambled control to the rat hippocampus following severe TBI. The miR-146a mimic-nanoparticle complex was injected into the dorsal hippocampus immediately following a severe CCI and animals sacrificed 48 hours later. Hippocampi were then harvested for analysis of TRAF6 and IRAK1 gene and protein (TRAF6) expression. RT-qPCR revealed a significant reduction in IRAK1 (P = 0.048) and TRAF6 (P = 0.036) mRNA levels following miR-146a delivery in comparison to scrambled control demonstrating the efficacy of the nanoparticle delivery system (Figure 4A). Importantly, Western blot analysis demonstrated that TRAF6 protein levels were also decreased in the hippocampus of animals receiving p5RHH + miR-146a nanoparticles (Figure 4B).

**Discussion**

Our previous studies reported that several mitochondria-enriched miRNAs regulating inflammation including miR-142-

| Table 2 | Response of inflammatory-related genes in hippocampus following TBI |
|---|---|---|---|
| Gene symbol | 1 d | 3 d | 7 d |
| **Anti-inflammatory** | Fold (range) | P-value | Fold (range) | P-value | Fold (range) | P-value |
| Arg1 | 0.37 (0.26–0.53) | 0.005 | 0.54 (0.39–0.75) | 0.025 | 1.29 (0.81–2.04) | 0.340 |
| Ccl2 | 0.69 (0.3–1.62) | 0.550 | 0.35 (0.30–0.42) | 0.151 | 1.08 (0.64–1.82) | 0.799 |
| Eps1 | 0.61 (0.47–0.78) | 0.001 | 0.85 (0.63–1.14) | 0.301 | 1.21 (0.88–1.66) | 0.271 |
| Il4 | 0.28 (0.12–0.61) | 0.072 | 0.57 (0.27–1.23) | 0.285 | 1.71 (0.73–4.01) | 0.351 |
| PparG | 0.61 (0.44–0.83) | 0.032 | 0.52 (0.24–1.11) | 0.178 | 1.01 (0.75–1.37) | 0.952 |
| Sphk1 | 6.10 (3.4–10.9) | 0.004 | 1.49 (1.03–2.16) | 0.390 | 0.83 (0.69–1.01) | 0.275 |
| Stat6 | 0.66 (0.51–0.86) | 0.013 | 0.92 (0.78–1.09) | 0.514 | 1.22 (0.95–1.56) | 0.155 |
| Tgfb1 | 1.53 (1.2–1.94) | 0.006 | 2.28 (1.85–2.82) | 0 | 0.60 (0.34–1.07) | 0.282 |
| Tgm2 | 2.40 (1.78–3.24) | 0.011 | 1.17 (0.99–1.39) | 0.220 | 1.31 (0.81–2.11) | 0.442 |
| **Pro-inflammatory** | Fold (range) | P-value | Fold (range) | P-value | Fold (range) | P-value |
| Ccr2 | 10.67 (6.3–18.0) | 0.004 | 4.93 (3.54–6.88) | 0.007 | 0.65 (0.21–2.00) | 0.505 |
| Ccl2 | 160 (85.5–299.8) | 0.002 | 15.43 (8.16–29.2) | 0.001 | 0.47 (0.27–0.94) | 0.290 |
| Ccr7 | 0.47 (0.3–0.74) | 0.079 | 0.74 (0.45–1.21) | 0.361 | 1.54 (1.14–2.07) | 0.242 |
| Cd86 | 0.87 (0.47–1.59) | 0.646 | 2.51 (1.91–3.31) | 0.007 | 0.50 (0.28–0.92) | 0.290 |
| Il12a | 0.30 (0.21–0.42) | 0.001 | 0.55 (0.45–0.67) | 0 | 1.26 (0.94–1.67) | 0.396 |
| Il12b | 0.36 (0.24–0.54) | 0.010 | 0.44 (0.30–0.64) | 0.057 | 0.76 (0.51–1.14) | 0.235 |
| Il1b | 9.46 (6.55–13.7) | 0.001 | 1.58 (1.23–2.04) | 0.053 | 0.50 (0.27–0.94) | 0.458 |
| Il6 | 0.78 (0.51–1.18) | 0.392 | 0.36 (0.18–0.72) | 0.512 | 1.13 (0.77–1.68) | 0.737 |
| Nos2 | 2.12 (0.47–2.64) | 0.009 | 0.51 (0.36–0.60) | 0.098 | 0.47 (0.12–1.80) | 0.702 |
| Stat1 | 0.89 (0.71–1.4) | 0.377 | 0.98 (0.81–1.18) | 0.720 | 1.15 (0.87–1.52) | 0.314 |
| Ilf1 | 1.48 (0.9–1.45) | 0.010 | 3.13 (2.11–4.63) | 0.004 | 0.54 (0.24–1.19) | 0.313 |
| Nlrp3 | 1.04 (0.77–1.4) | 0.801 | 2.17 (1.82–2.59) | 0.001 | 0.85 (0.54–1.32) | 0.590 |
| Igar | 1.56 (1.23–1.98) | 0.018 | 3.07 (2.52–3.74) | 0.001 | 0.71 (0.41–2.11) | 0.419 |

A TLDA card consisting of selected inflammatory responsive genes was used to assess mRNA expression levels in hippocampal cytoplasm obtained from each time points following traumatic brain injury (TBI). Gene expression levels were calculated using formula: fold change = 2-ΔΔCt(control sample - experimental sample) comparing the ipsilateral injured side of the brain relative to the uninjured contralateral side. Data represent the mean value of 4 biological replicates, and the P values were generated using two-tailed Student’s t-tests. The fold change range is shown in parentheses. Significantly different (P < 0.05) expression genes are highlighted in red for increased expression and in green for decreased expression. The 1, 3, and 7 days indicate the tissue harvesting time following controlled cortical impact.
delivery of miR-146a for targeting modulators of NF-κB pro-inflammatory responses that maintain expression of the reparative phase of injury (Morganti-Kossmann et al., 2002; Semple et al., 2010; Ziebell and Morganti-Kossmann, 2010; Morganti et al., 2016). Therefore, sustaining a balance of inflammatory and reparative factors is a key factor in determining cell survival and recovery (Sullivan et al., 1998; Singh et al., 2006). In our current study, mitochondrial bioenergetics in the hippocampus were significantly compromised for at least 3 days following TBI and recovered by 7 days post injury. Interestingly, we observed that changes in the expression patterns of the mitochondria-enriched, inflammatory miRNAs (miR-142-3p, miR-142-5p, and miR-146a) were highly correlated with the loss in mitochondrial bioenergetics. Mitochondria can be considered as “first-responders” to cellular stressors and can trigger various signaling pathways involving gene regulation, in which miRNAs play a critical role. Therefore, we postulate that mitochondrial function may determine the subcellular localization and function of these mitochondria-enriched miRNAs. Future studies are needed to understand the correlation of translocation of these mitochondria-enriched, inflammatory miRNAs with mitochondrial bioenergetics and its functional impact on the inflammatory response following TBI.

Our miRNA and gene expression studies indicate that alterations of selected inflammatory related miRNAs and mRNAs peaked at 1 day and their activities remained elevated at 3 days. The activities were then reduced to control levels at 1 week following TBI. The inflammatory gene expression activity observed in this study is consistent with previous reports (Morganti-Kossman et al., 2002; Semple et al., 2010; Ziebell and Morganti-Kossman, 2010; Morganti et al., 2016) and may provide the basis for designing miRNA-based interventional strategies targeting acute times following TBI. For example, the neuroinflammatory response following TBI is characterized by a prolonged period of pro-inflammation persisting for months to years after the injury (Johnson et al., 2013). However, there also exists a reparative anti-inflammatory phase that is only present for acute time points following TBI (Jin et al., 2012; Hsieh et al., 2013; Kumar et al., 2016). Therefore, sustaining a balance of inflammatory responses that maintains expression of the reparative phase

translocation of mitochondria-enriched miRNAs in response to cellular stressors (e.g., TBI) that impact mitochondrial function. We now show that this compartmental shift is still evident up to 3 days following TBI for miR-142-3p and miR-142-5p in hippocampal tissue, and returns to normal levels by 7 days. Although not statistically significant, alterations in miR-146a showed a trend similar to miR-142-3p and miR-142-5p.

It has been well described that TBI elicits a rapid and significant loss of mitochondrial function in the hippocampus and is a key factor in determining cell survival and recovery (Wang et al., 2015, 2017). Specifically, the levels of these miRNAs were reduced significantly in mitochondria at 3–12 hours following TBI and remained significantly compromised for at least 3 days following TBI and recovered by 7 days post injury. Interestingly, we observed that changes in the expression patterns of the mitochondria-enriched, inflammatory miRNAs (miR-142-3p, miR-142-5p, and miR-146a) were highly correlated with the loss in mitochondria bioenergetics. Mitochondria can be considered as “first-responders” to cellular demands and stressors and can trigger various signaling pathways involving gene regulation, in which miRNAs play a critical role. Therefore, we postulate that mitochondrial function may determine the subcellular localization and function of these mitochondria-enriched miRNAs. Future studies are needed to understand the correlation of translocation of these mitochondria-enriched, inflammatory miRNAs with mitochondrial bioenergetics and its functional impact on the inflammatory response following TBI.

We suggested previously that mitochondria and miRNA interactions may play a role in miRNA cellular function related to inflammation in the injured brain (Wang et al., 2015, 2017). This was based on our observations that levels of miR-146a, miR-142-3p and miR-142-5p were significantly decreased in the mitochondrial fraction and increased in the cytoplasmic fraction at early time points post TBI (Wang et al., 2015). We hypothesized that this compartmental shift may represent a novel cell-penetrating peptide nanoparticle system for bioenergetics. We further demonstrate the utilization of miRNAs that is related to TBI-induced loss of mitochondrial compartmental re-distribution of mitochondria-enriched, inflammatory miRNAs. Future studies are needed to understand the correlation of translocation of these mitochondria-enriched, inflammatory miRNAs with mitochondrial bioenergetics and its functional impact on the inflammatory response following TBI.
will be essential to promoting tissue survival and recovery.

We found that miR-146a levels increased in cytoplasm initially, but then returned to control levels by 7 days post-injury. These observations are intriguing for at least two reasons. First, miR-146a plays a significant role in regulating the production of certain pro-inflammatory cytokines by targeting two adaptor proteins TRAF6 and IRAK1 that function upstream of the NF-κB signaling pathway (Liu et al., 2017). Second, miR-146a expression is actually regulated by activation of the NF-κB pathway (Taganov 2006). Thus, miR-146a is activated by and also acts as a negative feedback regulator of NF-κB signaling (Figure 5), which may enable a fine-tune control of the inflammatory response. Taken together, it is postulated that the transient expression of anti-inflammatory reparative signaling may be related to the downregulation of miR-146a through NF-κB. While the mechanisms by which miR-146a modulates inflammatory signaling in the injured brain are currently not known, these observations prompted us to test whether nanoparticle delivery of miR-146a can serve as a therapeutic strategy for modulating the neuroinflammatory response following TBI. Specifically, we employed a cell-permeable peptide p5RHH to deliver miR-146a mimic to cultured cell lines and also following TBI. The p5RHH cell-permeable peptide forms a non-immunogenic nanoparticle complex with small inhibitory RNA molecules and has been shown to be an effective delivery platform (Hou et al., 2013a; Hou et al., 2013b). Our data demonstrated that p5RHH effectively delivered miR-146a mimic and reduced the expression levels of TRAF6 and IRAK1 in vitro and in vivo. The results of these studies are encouraging as they establish a proof of principle that this novel nanoparticle delivery approach has therapeutic potential for modulating pro-inflammatory effectors in the injured brain.

Author contributions: Study concept and design: WXW, JES, PGS; Experimental implementations: WXW, PP, HIV, MS, ALC; Data acquisition and analysis: WXW, PP, HIV; Statistical analysis: WXW, PP, HIV; Manuscript preparation: WXW, JES, PP, HIV; Manuscript editing and review: JES, WXW, PGS, PP, HIV, MS, ALC. All authors approved the final version of the manuscript.

Conflicts of interest: All authors report no conflicts of interests.

Financial support: his work is supported by a grant (15-12A) from the Kentucky Spinal Cord and Head Injury Research Trust to JES and WXW.

Institutional review board statement: All of the studies performed were approved by the University of Kentucky Institutional Animal Care and Usage Committee (IACUC protocol # 2014-1300) on August 17, 2017.

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Data sharing statement: Datasets analyzed during the current study are available from the corresponding author on reasonable request.

Plagiarism check: Checked twice by iThenticate.

Peer review: Externally peer reviewed.

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Additional files:
Additional Figure 1: Profile of mitochondria-enrichment of miRNAs in sham operated animals.
Additional Figure 2: In vitro delivery of p5RHH peptide-based miR-146a nanoparticles to cultured SH-SYSY cells.
Additional Table 1: Customized TaqMan low density array (TLDA) Genes and the assay ID.

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**Research Article**

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C-Editors: Zhao M, Li CH; T-Editor: Jia Y
Additional Figure 1 Profile of mitochondria-enrichment of miRNAs in sham operated animals.

Sham operated animals (total n = 4 per group) were euthanatized at 1, 3, or 7 days following surgery and the ipsilateral and contralateral hippocampi were rapidly removed for isolation of the mitochondria and cytosolic fractions. The fractions were then subjected to RNA isolation and miRNA analysis. The level of mitochondria-enrichment of a miRNA expressed as relative quantity (RQ) was calculated using the formula

$$RQ = 2^{(Ct(Mito) - Ct(Cyto))}$$

where mitochondrial and cytoplasmic fractions were isolated from the same tissue. Green and orange bars correspond to the contralateral and ipsilateral hippocampi, respectively, of the sham-surgery animals. Data are expressed as the mean ± SD.
Additional Figure 2 \textit{In vitro} delivery of p5RHH peptide-based miR-146a nanoparticles to cultured SH-SY5Y cells.

Confocal microscopy graph of miR-146a and scramble control transfections (Scale bars:10 \( \mu \)m). Pre-seeded SH-SY5Y cells were incubated either with miR-146a or scramble mimic/p5RHH-nanoparticle complexes or miR-146a or scramble mimic/Lipofectamine, or without carrier for 24 hours and the cells were visualized and photographed using a Nikon C2+ confocal microscope.