



HAL
open science

High-Dose Melatonin and Ethanol Excipient Combined with Therapeutic Hypothermia in a Newborn Piglet Asphyxia Model

Nicola J Robertson, Ingran Lingam, Christopher Meehan, Kathryn A Martinello, Adnan Avdic-Belltheus, Liane Stein, Mohamed Tachrount, David Price, Magdalena Sokolska, Alan Bainbridge, et al.

► **To cite this version:**

Nicola J Robertson, Ingran Lingam, Christopher Meehan, Kathryn A Martinello, Adnan Avdic-Belltheus, et al.. High-Dose Melatonin and Ethanol Excipient Combined with Therapeutic Hypothermia in a Newborn Piglet Asphyxia Model. *Scientific Reports*, 2020, 10 (1), pp.3898. 10.1038/s41598-020-60858-x . inserm-02504046

HAL Id: inserm-02504046

<https://inserm.hal.science/inserm-02504046>

Submitted on 10 Mar 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

OPEN

High-Dose Melatonin and Ethanol Excipient Combined with Therapeutic Hypothermia in a Newborn Piglet Asphyxia Model

Nicola J. Robertson^{1*}, Ingran Lingam¹, Christopher Meehan¹, Kathryn A. Martinello¹, Adnan Avdic-Belltheus¹, Liane Stein¹, Mohamed Tachrount¹, David Price², Magdalena Sokolska², Alan Bainbridge², Mariya Hristova¹, Bobbi Fleiss³, Boris W. Kramer⁴, Pierre Gressens⁵ & Xavier Golay¹

With the current practice of therapeutic hypothermia for neonatal encephalopathy, disability rates and the severity spectrum of cerebral palsy are reduced. Nevertheless, safe and effective adjunct therapies are needed to optimize outcomes. This study's objective was to assess if 18 mg/kg melatonin given rapidly over 2 h at 1 h after hypoxia-ischemia with cooling from 1–13 h was safe, achieved therapeutic levels within 3 h and augmented hypothermic neuroprotection. Following hypoxia-ischemia, 20 newborn piglets were randomized to: (i) Cooling 1–13 h (HT; n = 6); (ii) HT + 2.5% ethanol vehicle (HT+V; n = 7); (iii) HT + Melatonin (HT+M; n = 7). Intensive care was maintained for 48 h; aEEG was acquired throughout, brain MRS acquired at 24 and 48 h and cell death (TUNEL) evaluated at 48 h. There were no differences for insult severity. Core temperature was higher in HT group for first hour after HI. Comparing HT+M to HT, aEEG scores recovered more quickly by 19 h ($p < 0.05$); comparing HT+V to HT, aEEG recovered from 31 h ($p < 0.05$). Brain phosphocreatine/inorganic phosphate and NTP/exchangeable phosphate were higher at 48 h in HT+M versus HT ($p = 0.036$, $p = 0.049$ respectively). Including both 24 h and 48 h measurements, the rise in Lactate/N-acetyl aspartate was reduced in white ($p = 0.030$) and grey matter ($p = 0.038$) after HI. Reduced overall TUNEL positive cells were observed in HT+M (47.1 cells/mm²) compared to HT (123.8 cells/mm²) ($p = 0.0003$) and HT+V (97.5 cells/mm²) compared to HT ($p = 0.012$). Localized protection was seen in white matter for HT+M versus HT ($p = 0.036$) and internal capsule for HT+M compared to HT ($p = 0.001$) and HT+V versus HT ($p = 0.006$). Therapeutic melatonin levels (15–30 mg/l) were achieved at 2 h and were neuroprotective following HI, but ethanol vehicle was partially protective.

Intrapartum-related neonatal encephalopathy (NE) is a major healthcare problem. Worldwide in 2010, NE accounted for 287,000 deaths and 400,000 survivors with impairment¹. NE cannot be prevented in most cases and therapies are limited. The incidence of NE in Western Europe is 1–3/1000 term births and in low- and mid-resource settings the incidence is ~10 times higher^{1,2}. Over the last 2 decades, in settings with neonatal intensive care facilities, therapeutic hypothermia (HT) is routinely used for moderate-to-severe NE, improving survival and reducing disability³. However, although the severity of cerebral palsy has reduced with HT⁴, survivors have significantly lower cognitive scores which are on average 14 IQ points lower than matched peers even in the absence of cerebral palsy at school-age⁵. Further adjustments to HT protocols do not improve outcome^{6,7}, therefore adjunct therapies to augment HT protection are urgently needed.

Pre-clinical studies suggest that melatonin (N-acetyl-5-methoxytryptamine) in pharmacologic levels is safe and neuroprotective for hypoxic-ischemic injury in the adult⁸ and neonatal⁹ brain, mediated by anti-oxidant, anti-apoptotic and anti-inflammatory properties^{10,11}. Extrapolating from *in vitro*¹² and pre-clinical piglet studies^{13,14} showing melatonin reduces cell death in a concentration-dependent manner, we deduce that a plasma

¹University College London, London, WC1E 6HX, UK. ²University College London Hospitals NHS Trust, London, UK. ³RMIT, Melbourne, Australia. ⁴University of Maastricht, Maastricht, The Netherlands. ⁵INSERM, Paris, France. *email: n.robertson@ucl.ac.uk

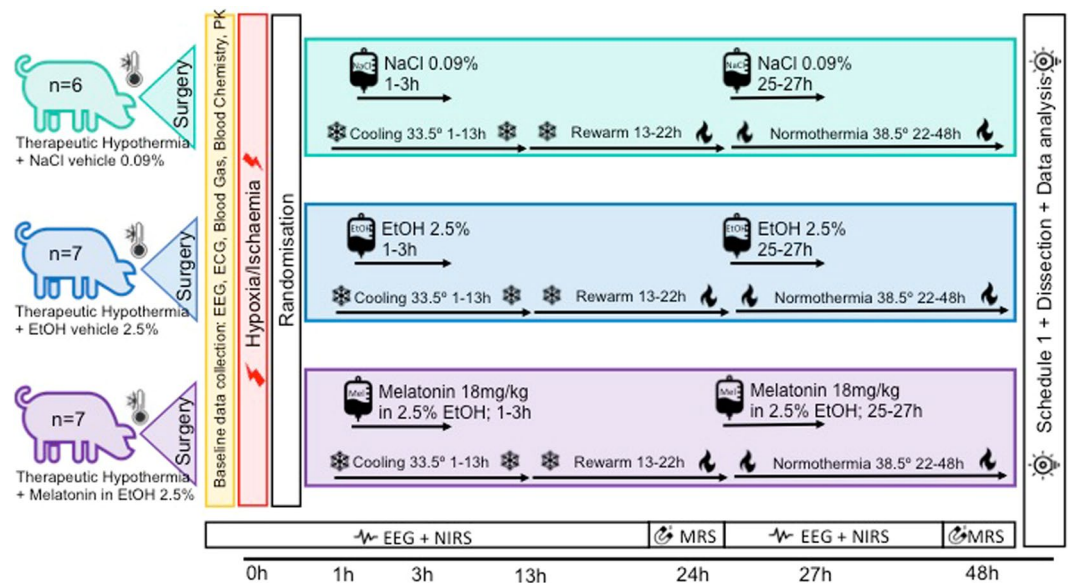


Figure 1. Study time-line. Following baseline data acquisition, piglets underwent cerebral HI. At the end of HI (time $t = 0$), piglets were randomized to (i) HT (33.5°C) from 1–13 h with saline bolus at 1–3 and 25–27 h or (ii) HT+vehicle (HT+V; 2.5% ethanol); (iii) HT+melatonin (HT+M; 18 mg/kg dissolved in 2.5% ethanol) at 1–3 and 25–27 h. Intensive care support was given for 48 h following HI. aEEG/continuous video EEG was acquired. MRI/MRS was performed at 3 T at 24 and 48 h. The experiment was terminated at 48 h and brain immunohistochemistry analyzed.

level 15–30 mg/l is required for optimal protection. As melatonin is sparingly soluble, solubility enhancers, such as ethanol, are used. Ethanol could confound previous studies of melatonin protection^{13,15–19}, as low-dose ethanol ~4 h after HI is protective in adult models of stroke^{20–22}. In a previous piglet study, we observed augmentation of HT protection with 30 mg/kg/24 h melatonin with ethanol excipient when started 10 mins after hypoxia-ischemia (HI), reaching blood levels >15 mg/l within 1 h¹³. In a subsequent study using a proprietary melatonin formulation (15 mg/kg), without ethanol excipient, given 2–8 h after HI, there was less clear protection with therapeutic levels reached only at 8 h¹⁴. In this same study, no protection was seen with 5 mg/kg melatonin started at 2 h (blood melatonin <4 mg/l)¹⁴. Taken together, these studies suggest melatonin protection is dose-dependent, time critical and influenced by excipient. With the aim to achieve therapeutic levels within 3 h of HI, we subsequently performed pharmacokinetic (PK) modelling, suggesting optimised melatonin dosing would be 18 mg/kg/24 h over 2 h started 1 h after HI¹⁴.

We used a piglet model with similar protocols to our neonatal intensive care unit and similar sequences for magnetic resonance spectroscopy (MRS) as NE babies. Our objective was to assess safety, efficacy and PK of an optimized melatonin-dosing regimen based on prior PK-modelling. We hypothesized that 18 mg/kg melatonin started 1 h after HI, infused over 2 h would reach therapeutic levels within 3 h and augment HT. To clarify any influence of ethanol vehicle on neuroprotection, we studied ethanol with and without melatonin. Primary outcome measures were: (i) Cerebral MRS biomarkers (proton (1H) and phosphorus (³¹P) MRS). Thalamic lactate/N-acetyl aspartate (Lac/NAA) is the most accurate outcome biomarker at 2-years^{23,24}, used in clinical neuroprotection trials²⁵, with clear superiority over other MR methods²⁴. ³¹P MRS is less accessible on MRI systems but has defined secondary energy failure in NE²⁶ and its relation with 1-year brain growth and outcome²⁷; (ii) aEEG recovery; aEEG is used in NE babies during cooling and recovery predicts outcome²⁸; (iii) Quantitative cell death in 8 brain regions (TUNEL-positive cells) at 48 h.

Materials and Methods

Animal experiments, surgical preparation and randomization. All animal experiments were approved by the UCL Ethics Committee and performed according to UK Home Office Guidelines [Animals (Scientific Procedures) Act, 1986]. The study complies with ARRIVE guidelines. Large White male piglets were anesthetized, surgically prepared and intensive care maintained as described previously²⁹. Criteria for study entry were: (i) normal aEEG/EEG at baseline after surgery; (ii) no pyrexia; (iii) no aEEG recovery within 1 h of HI. The experimental plan is shown in Fig. 1.

Following HI and resuscitation, animals were randomized to (i) Therapeutic hypothermia 1–13 h (HT) (0.9% saline (7.2 ml/kg/h) at 1–3 h and 25–27 h); (ii) HT+Vehicle (HT+V) (2.5% ethanol (7.2 ml/kg/h) at 1–3 h and 25–27 h); (iii) HT+ Melatonin (HT+M) (18 mg/kg (7.2 ml/kg/h) at 1–3 h and 25–27 h). Intensive care support for the animal throughout 48 h and complications were managed in accordance with local neonatal intensive care guidelines.

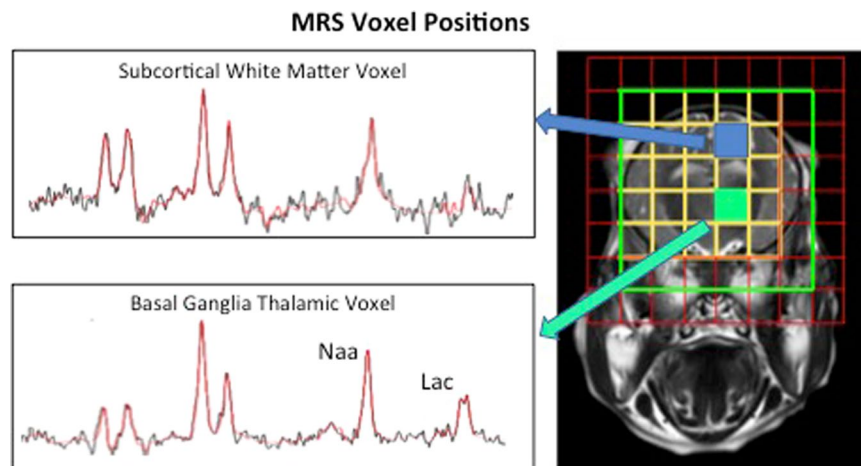


Figure 2. Representative spectra and voxel positions for ^1H MRS. Spectroscopy data were collected from voxels located in the dorsal right subcortical white matter at the centrum semiovale level and from two voxels in the deep grey matter centred on both thalami.

Transient cerebral hypoxia ischemia. Compared to the original studies based on assessment of NTP reduction during HI on ^{31}P MRS²⁹, the monitoring and titration of the HI insult were modified recently in this and other studies³⁰. Baseline physiological observations and aEEG were monitored prior to HI. Carotid occluders were inflated and fraction of inspired oxygen (FiO_2) reduced simultaneously at the start of the insult. FiO_2 was decreased to 6% over the first 3 min and titrated to mean blood pressure and EEG. Oxygen delivery was increased in the event of a mean BP <27 mmHg and restricted further if recovery of EEG activity was observed during the insult. Blood gas analysis was performed at 5 min intervals during HI. Total duration of HI was anticipated to be 20–25 min, depending on the duration of isoelectric EEG, hypotension (mean BP <30 and <25 mmHg), total reduction in FiO_2 (AUC FiO_2) and severity of acidosis on blood gas analysis. At the end of the insult, the animal was resuscitated, occluders deflated and FiO_2 increased to air.

Melatonin administration. *Melatonin preparation and delivery.* Melatonin (Sigma-Aldrich) was dissolved in ethanol and 0.9% NaCl (2.5% v/v vehicle) in the dark shortly before administration. 18 mg/kg melatonin (7.2 ml/kg/h) was infused intravenously over 2 h starting at 1 h after HI and at 25 h after HI. Blood was sampled at baseline and 2, 3, 6, 24, 26, 27, 36, 42, 48 h after time 0 (end of HI).

Magnetic resonance spectroscopy. ^{31}P and ^1H MRS was performed at 24 and 48 h after HI in a Philips clinical 3 T MRI scanner. ^{31}P metabolites were measured over whole brain (see supplementary file). ^1H MRS metabolites were measured in white matter in the right subcortical region ($8 \times 8 \times 15$ mm) and deep grey matter ($15 \times 15 \times 10$ mm) in the thalamus (Fig. 2). Data was analyzed using jMRUI and Lac/NAA peak area ratio calculated.

Amplitude integrated electroencephalogram (aEEG). A multichannel EEG and aEEG (Nicolet) was acquired at baseline and continued for 48 h post-insult. The aEEG score was based on pattern classification³¹; isoelectric (0), continuous low voltage (1), burst suppression (2), discontinuous normal voltage (3) and continuous normal voltage (4).

Immunohistochemistry. At 48 h after HI, piglets were euthanized with pentobarbital and the brain fixed by cardiac perfusion with cold 4% paraformaldehyde, dissected out and post-fixed at 4 °C in 2% paraformaldehyde for 7 days. Coronal slices (5 mm thick) of the right hemisphere, starting from anterior to the optic chiasma, were embedded in paraffin, sectioned to 8 μm thickness and stained with hematoxylin and eosin to validate the bregma for analysis. For each animal, 2 sections (bregma 00 and -2.0) were stained and 8 brain regions were examined: cingulate cortex, sensorimotor cortex, hippocampus, internal capsule, periventricular white matter, caudate, putamen and thalamus. Histological analysis was performed by an investigator blinded to treatment allocation³².

Please see Supplementary file for details on immunohistochemistry methodology and statistical analysis.

Results

One piglet (HT+M) died at 21 h and one piglet (HT) was excluded due to aEEG recovery within 1 h of HI, suggesting mild HI. Twenty piglets were included; HT (n = 6), HT+V (n = 7); HT+M (n = 7).

Physiological data and insult severity. There were no intergroup differences for bodyweight, baseline heart rate, MABP and core temperature (Table 1). The arterial gases at baseline were similar. Considering HI severity, the duration of (i) hypoxia and carotid occlusion, (ii) isoelectric EEG and (iii) MABP <25 and <30 mmHg were similar (Table 1). At the end of HI, blood pH, lactate, base excess and AUC FiO_2 reduction below 21% were similar (Table 1).

Parameter	HT IS Mean (SEM)	HT+Vehicle IS Mean (SEM)	HT+Melatonin IS Mean (SEM)	P value
Weight (kg)	2.1 (0.05)	2.1 (0.04)	2.0 (0.05)	0.776
Hypoxic ischemic insult				
Duration of HI insult (min)	21.3 (1.1)	20.0 (1.0)	23.1 (1.0)	0.103
Duration of isoelectric EEG during HI (min)	19 (0.08)	18.3 (0.8)	20.9 (0.8)	0.081
Duration of blood pressure <30 mmHg (min)	8.7 (1.4)	10.7 (1.3)	6.9 (1.3)	0.150
Duration of blood pressure <25 mmHg (min)	2.3 (1.2)	3.2 (1.1)	0.7 (1.1)	0.321
Nadir pH	7.2 (0.03)	7.2 (0.03)	7.2 (0.03)	0.433
Nadir Lactate (mmol/l)	14.3 (0.7)	11.9 (0.7)	13.3 (0.7)	0.070
Area under the curve FiO ₂	287.2 (19.6)	271.6 (18.2)	322.4 (18.2)	0.160
Nadir base excess (mmol/l)	-13.2 (2.94)	-9.7 (1.2)	-9.6 (1.2)	0.103
Heart rate (min⁻¹)				
Baseline	179.9 (8.2)	166.2 (7.6)	175.4 (7.6)	0.471
0-1 h after insult	200 (10.0)	197.6 (9.2)	192.6 (9.2)	0.857
1-13 h	181.8 (6.82)	171.3 (6.3)	171.0 (6.3)	0.445
13-25 h	202 (11.5)	195.6 (10.6)	181.4 (10.6)	0.416
25-48 h	212.4 (9.4)	182.1 (7.9)	187.1 (7.9)	0.061
Mean arterial blood pressure (mmHg)				
Baseline	52 (2.6)	51.2 (2.4)	50.2 (2.4)	0.875
0-1 h after insult	53.3 (5.0)	44.3 (4.6)	45.4 (4.6)	0.384
1-13 h	46 (1.6)	42.4 (1.5)	45.8 (1.5)	0.191
13-25 h	45.8 (2.5)	42.6 (2.3)	50.3 (2.3)	0.091
25-48 h	55.0 (2.1)	49.4 (1.8)	52.6 (1.8)	0.152
Rectal temperature (°C)				
Baseline	38.1 (0.3)	38.3 (0.3)	38.0 (0.3)	0.777
0-1 h after insult	38.9 (0.2)	38.5 (0.1)	38.2 (0.1)	0.020
1-13 h	34.0 (0.05)	34 (0.05)	33.9 (0.05)	0.615
13-25 h	36.2 (0.1)	36.5 (0.1)	36.4 (0.1)	0.295
25-48 h	38.4 (0.09)	38.5 (0.08)	38.4 (0.08)	0.794
PaO₂ (kPa)				
Baseline	11.1 (1.3)	9.9 (1.2)	14.2 (1.2)	0.067
End of insult (time 0)	9.3 (1.2)	6.4 (1.1)	7.6 (1.1)	0.241
12h after time 0	11.5 (1.0)	11.3 (0.9)	10.8 (0.9)	0.879
24h after time 0	11.5 (1.0)	11.4 (0.9)	12.8 (0.9)	0.469
48h after time 0	14.1 (0.7)	12.0 (0.6)	12.7 (0.6)	0.119
PaCO₂ (kPa)				
Baseline	5.6 (0.4)	6 (0.4)	5.4 (0.4)	0.506
End of insult (time 0)	5.6 (0.4)	6.1 (0.4)	6 (0.4)	0.628
12h after time 0	4.8 (0.5)	5.4 (0.5)	5.6 (0.5)	0.521
24h after time 0	7 (0.3)	5.6 (0.3)	5.5 (0.3)	0.003
48h after time 0	5 (0.3)	4.5 (0.3)	5.2 (0.3)	0.311
Blood pH				
pH_Baseline	7.4 (0.03)	7.4 (0.03)	7.5 (0.03)	0.301
End of insult (time 0)	7.2 (0.04)	7.2 (0.03)	7.2 (0.03)	0.960
12h after time 0	7.5 (0.04)	7.5 (0.03)	7.5 (0.03)	0.641
24h after time 0	7.3 (0.03)	7.4 (0.02)	7.5 (0.02)	0.009
48h after time 0	7.5 (0.02)	7.5 (0.02)	7.5 (0.02)	0.743
Base excess (mmol/l)				
Baseline	1.8 (1.7)	3.3 (1.6)	5.4 (1.6)	0.319
End of insult (time 0)	-10.0 (1.7)	-9.7 (1.6)	-9.6 (1.6)	0.982
12h after time 0	5.7 (1.5)	4.7 (1.4)	5.7 (1.4)	0.845
24h after time 0	2.2 (2.6)	-0.4 (2.2)	5.1 (2.2)	0.222
48h after time 0	2.8 (1.3)	1.9 (1.1)	3.1 (1.1)	0.689
Lactate (mmol/l)				
Baseline	5.2 (0.7)	3.8 (0.7)	4.1 (0.7)	0.347
End of insult (time 0)	12.9 (1.0)	12.3 (0.9)	13.3 (0.9)	0.756
12h after time 0	3.1 (0.6)	3.9 (0.6)	3.1 (0.6)	0.543
Continued				

Parameter	HT LS Mean (SEM)	HT+Vehicle LS Mean (SEM)	HT+Melatonin LS Mean (SEM)	P value
24 h after time 0	3.3 (1.5)	5.3 (1.3)	2.1 (1.3)	0.227
48 h after time 0	1.2 (0.3)	1.7 (0.2)	1.5 (0.2)	0.425
Glucose (mmol/l)				
Baseline	6.4 (0.5)	6 (0.5)	5.6 (0.5)	0.493
End of insult (time 0)	9.3 (1.1)	9.2 (1.0)	9.7 (1.2)	0.942
12 h after time 0	15.2 (1.6)	13.3 (1.5)	11.3 (1.5)	0.254
24 h after time 0	17 (3.9)	13.2 (3.3)	8.9 (3.3)	0.307
48 h after time 0	7.2 (1.0)	6.4 (0.9)	6 (0.9)	0.657

Table 1. Physiological parameters for the piglets in each group. Time zero = time of resuscitation after HI. Least square mean values (SEM) are presented for the three groups: (i) HT (n = 6), (ii) HT+Vehicle(Ethanol) (n = 7), and (iii) HT+Melatonin (n = 7). An analysis of variance (ANOVA) model was fitted to each group at each time point. No differences were observed between any groups at any time point or ranges following Bonferonni multiplicity correction ($p < 0.001$). Bold figures represent those measurements that are significantly different between groups.

Infusions	HT		HT+V		HT+M		P value
	Mean (SD)	Mean (SD)	Mean (SD)	Mean (SD)	Mean (SD)	Mean (SD)	
Dopamine ($\mu\text{g/kg/min}$)	15.3	4.5	16.7	3.5	8.9	4.1	0.005
Dobutamine ($\mu\text{g/kg/min}$)	2.7	5.4	7.1	6.4	0.7	1.9	0.07
Noradrenaline (ng/kg/min)	13.0	23.2	43.5	70.3	17.9	47.4	0.53
Adrenaline (ng/kg/min)	157.0	348.5	192.4	477.9	14.2	37.6	0.60
10 ml/kg Saline Bolus (n)	0.2	0.4	0.4	1.1	0.1	0.4	0.73

Table 2. Average total volume replacement and inotrope infusion for the piglets in the HT, HT+V, HT+M groups during the 48 h after HI.

In the first hour after HI, the mean core temperature was higher in HI versus HI+M (38.9 °C versus 38.2 °C, $p = 0.006$), although temperatures were within 0.4 °C of normal piglet temperature (38.5 °C). For the remainder of the study, there was no temperature difference. The blood pH was more alkalotic and PaCO_2 lower at 24 h after HI in HT+M ($p = 0.001$) and HT+V ($p = 0.003$) versus HT.

There was no difference in saline boluses. Inotropic use with Dopamine and Dobutamine was higher in HT and HT+V versus HT+M. Noradrenaline and adrenaline use did not differ (Table 2).

aEEG recovery and seizures. aEEG background activity scores were 4 (normal) in all piglets before HI and 0 during HI. Following HI, the mean hourly aEEG scores were higher in HT+M versus HT+V and HT from 19–24 h ($p = 0.037$ and $p = 0.025$ respectively) and 25–30 h ($p = 0.022$ and $p = 0.010$) after HI, indicating faster recovery of electrical activity with melatonin-augmented cooling. From 31 h, the vehicle treated group's brain electrical activity recovered and there was no difference between HT+M and HT+V. aEEG background voltage did not recover in HT and was lower than both HT+V and HT+M, with scores of < 1 from 31 h until the study end (Fig. 3).

One piglet in HT and 3 piglets each in HT+V and HT+M had seizures. All electrographic seizures were treated with 20 mg/kg phenobarbitone; short clinical seizures with no electrographic evidence of seizures were not treated. One animal in HT and HT+M and 3 animals in HT+V received phenobarbitone.

3T MRS. 48 h after HI, comparing HT+M to HT, there was higher PCr/Pi ($p = 0.036$) and NTP/epp ($p = 0.049$). Comparing HT+M to HT including 24 and 48 h data, Lac/NAA was lower in the white matter ($p = 0.30$) and grey matter ($p = 0.038$). There were no differences in MRS comparing HT and HT+V or HT+V and HT+M (Fig. 4).

Immunohistochemistry. TUNEL. TUNEL-positive cells/ mm^2 for treatment groups are shown in Fig. 5 with group comparison in Table 3. Over all regions, TUNEL-positive cells/ mm^2 were 123.8 cells/ mm^2 in HT, 97.5 cells/ mm^2 in HT+V and 47.1 cells/ mm^2 in HT+M groups. There were lower TUNEL-positive cells comparing HT+M versus HT+V ($p = 0.024$) and HT+M versus HT ($p = 0.001$). For regional differences; there were lower TUNEL-positive cells in the IC in HT+M (12.2 cells/ mm^2) versus HT (259.6 cells/ mm^2 ; $p < 0.001$). In the same region, there was a partial protective effect with ethanol with lower TUNEL-positive cells in HT+V (98.1 cells/ mm^2) versus HT (259.6 cells/ mm^2 ; $p = 0.006$). In the pvWM there were less TUNEL-positive cells in HT+M (9.7 cells/ mm^2) versus HT (133.4 cells/ mm^2 ; $p = 0.036$).

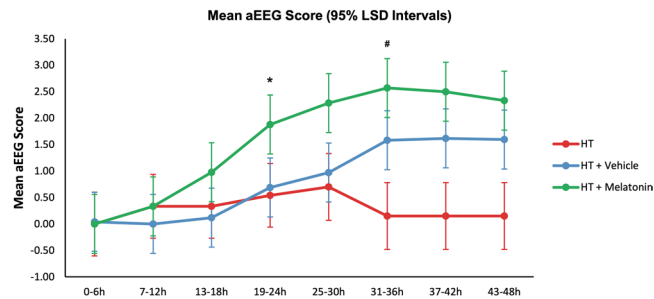


Figure 3. Amplitude-integrated electroencephalogram (aEEG) background activity at baseline, during HI and following HI. The grouped mean hourly aEEG scores per 6 h period with 95% Least Significant Difference (LSD) are shown. Non-overlapping bars show evidence of a significant difference. The mean hourly aEEG scores were significantly higher in the HT+M group versus HT+V and HT from 19–24 h ($p = 0.037$ and $p = 0.025$ respectively) and 25–30 h ($p = 0.022$ and $p = 0.010$ respectively) after HI, indicating faster recovery of brain electrical activity. From 31 h onwards, the vehicle treated group's brain electrical activity recovered and there was no difference between the HT+M and HT+V groups. The aEEG background voltage did not recover in the HT group and was significantly lower than both HT+V and HT+M groups, with scores of <1 from 31 h until the end of the study. *Shows when the HT+M group aEEG recovered compared to HT+V and HT groups. #Shows when the HT+V group aEEG recovered compared to the HT group.

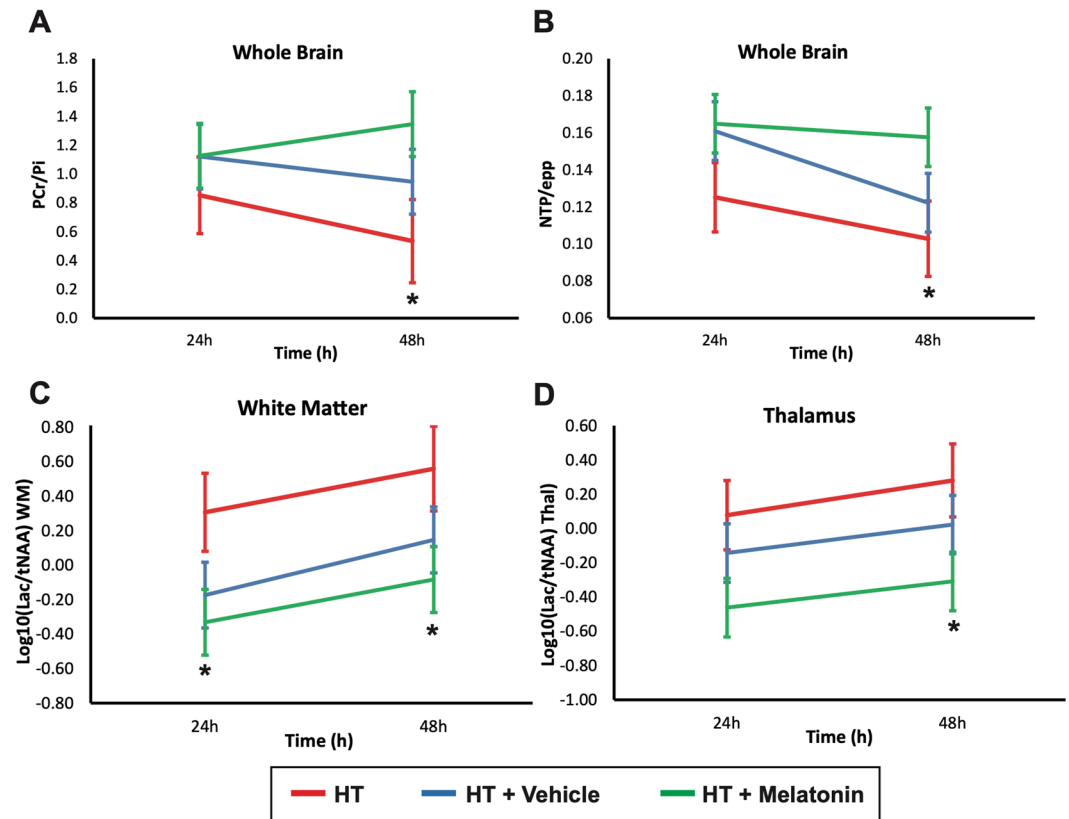


Figure 4. 1H Magnetic resonance spectroscopy of the brain at 24 and 48 h after HI. Least square mean plots with 95% Least Significant Difference (LSD) bars for the NTP/epp and PCr/Pi in whole-forebrain, and Lac/NAA in thalamus and white matter; non-overlapping bars show evidence of a significant difference. NTP/epp (A) and PCr/Pi (B) means were significantly higher in the HT+M group compared to HT at 48 h ($p = 0.049$ and 0.036 respectively). Comparing the HT+M group to the HT group including both 24 and 48 h, Lac/NAA was lower in the white matter ($p = 0.30$) and grey matter ($p = 0.038$). There were no significant differences in MRS measures comparing HT and HT+V groups and HT+V and HT+M groups. epp = exchangeable phosphate pool; Lac = lactate; NAA = N-acetyl aspartate; Thal = thalamus; WM = white matter; HI = hypoxia ischemia; M = melatonin * $p < 0.05$.

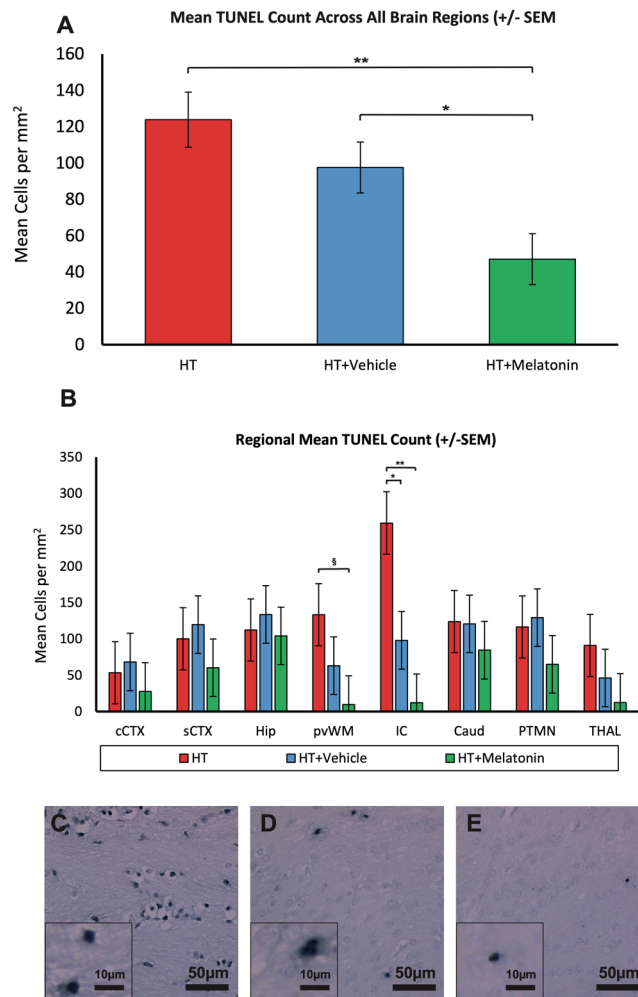


Figure 5. (A) TUNEL immunohistochemistry, all brain regions. Co-treatment of cooling with melatonin decreased overall TUNEL positive cell death at 48 h after HI when compared to cooling and vehicle ($p = 0.012$) and cooling alone ($p = 0.003$). $*p = 0.012$ HT+M versus HT+V. $**p = 0.003$ HT+M versus HT. (B) TUNEL immunohistochemistry from 8 brain regions. There were less TUNEL-positive cells in the IC in HT+M compared to both HT+V and HT groups. There were fewer TUNEL-positive cells in the pvWM in the HT+M group versus HT alone. $**p < 0.001$ HT+M versus HT. $*p < 0.01$ HT+V versus HT. $^{\$}p > 0.05$ HT+M versus HT. Internal capsule = IC; pvWM = periventricular white matter. Error bars represent standard error.

Iba1. There was no difference in overall ramification index (higher index suggests less activation) in HT+M versus HT+V or HT (Fig. 6A). On regional analysis the caudate showed increased IBA1 ramification index in HT+V versus HT (Fig. 6E).

GFAP. On GFAP luminosity, there was no difference in overall astrogliosis (Fig. 6B), but localized increases in cCTX, IC and caudate (Fig. 6F).

Cleaved caspase 3 (CC3). There was an overall increase in CC3 in HT+M versus HT and HT+M versus HT+V (Fig. 6C). Localized differences with increased CC3 mainly in HT+M versus HT+V and HT were seen in cCTX, sCTX, pvWM, caudate, putamen, thalamus (Fig. 6G).

OLIG2. The estimated mean OLIG2 cells/mm² for are shown in Fig. 6D. Over all brain regions, OLIG2 cells/mm² were 324.8 cells/mm² in HT, 532.0 cells/mm² in HT+V and 496.5 cell/mm² in HT+M. There were higher numbers of OLIG2 cells in HT+V and HT+M versus HT (both $p < 0.0001$). For regional differences, there were higher OLIG2 counts in internal capsule in HT+V (1204.5 cells/mm²) versus HT (831.2 cells/mm²; $p < 0.0001$) and HT+M (1204.9 cells/mm²) versus HT (831.2 cells/mm²; $p < 0.0001$) (Fig. 6H). There were higher OLIG2 counts in the pvWM in HT+V (1451.6 cells/mm²) versus HT (716.3 cells/mm²; $p < 0.0001$) and HT+M (1316.4 cells/mm²) versus HT (831.2 cells/mm²; $p < 0.0001$). Excluding IC and pvWM regions (variability was larger than other regions), we observed higher OLIG2 counts in: hippocampus, HT+V (260.9 cells/mm²) versus HT (142.3 cells/mm²; $p = 0.041$); putamen, HT+V (259.7 cells/mm²) versus HT (137.6 cells/mm²; $p = 0.036$) and

TUNEL positive cells per mm ² Least square mean (SEM)					
	HT	HT+V	HT+M	P values for difference in means	
cCTX	53.4 (42.8)	68.3 (39.6)	27.7 (39.6)	HT v HT+M	p = 0.47
				HT v HT+V	p = 0.80
				HT+V v HT+M	p = 0.66
sCTX	100.2 (42.8)	119.7 (39.6)	60.4 (39.6)	HT v HT+M	p = 0.50
				HT v HT+V	p = 0.74
				HT+V v HT+M	p = 0.29
Hip	112.4 (42.8)	133.7 (39.6)	104.3 (39.6)	HT v HT+M	p = 0.89
				HT v HT+V	p = 0.72
				HT+V v HT+M	p = 0.60
pvWM	133.4 (42.8)	63.3 (39.6)	9.7 (39.6)	HT v HT+M	p = 0.036
				HT v HT+V	p = 0.23
				HT+V v HT+M	p = 0.34
IC	259.6 (42.8)	98.1 (39.6)	12.2 (39.6)	HT v HT+M	p = 0.001
				HT v HT+V	p = 0.006
				HT+V v HT+M	p = 0.13
Caudate	124.1 (42.8)	120.8 (39.6)	84.6 (39.6)	HT v HT+M	p = 0.50
				HT v HT+V	p = 0.96
				HT+V v HT+M	p = 0.52
Putamen	116.6 (42.8)	129.4 (39.6)	65.1 (39.6)	HT v HT+M	p = 0.38
				HT v HT+V	p = 0.83
				HT+V v HT+M	p = 0.25
Thalamus	91.0 (42.8)	46.4 (30.6)	12.6 (39.6)	HT v HT+M	p = 0.18
				HT v HT+V	p = 0.45
				HT+V v HT+M	p = 0.55
Overall	123.8 (15)	97.5 (14)	47.1 (14)	HT v HT+M	p = 0.0003
				HT v HT+V	p = 0.200
				HT+V v HT+M	p = 0.012

Table 3. Least Square mean (SEM) TUNEL positive cells (per mm²) in each brain region and overall. P values for the difference in means are shown in the right hand column; significant values are shown in bold. SEM = standard error of the mean; cCTX = cingulate cortex; sCTX = sensorimotor cortex; pvWM = periventricular white matter; IC = internal capsule.

thalamus, HT+V (441.1 cells/mm²) versus HT (242.0 cells/mm²; p = 0.001) and HT+M (372.3 cells/mm²) versus HT (242.0 cells/mm²; p = 0.02).

Pharmacokinetics. 18 mg/kg melatonin infusion over 2 h starting at 1 h after HI led to plasma target therapeutic levels of melatonin at 2 h after HI. The peak melatonin level in the first 24 h was at 3 h after HI (18.84 µg/ml) and in the 2nd 24 h at 27 h after HI (21.84 µg/ml). Individual subject profiles are in Fig. 7A and mean group plasma levels with 95% CI are in Fig. 7B.

Discussion

Compared to therapeutic hypothermia alone, we observed improved cerebral protection with the addition of 18 mg/kg melatonin given intravenously at 1 h and 25 h after HI, based on aEEG recovery from 19 h, improved cerebral energy metabolism at 48 h on ³¹P and ¹H MRS and reduced TUNEL-positive cells (estimated mean 47.1 cells/mm² in HT+M versus 123.8 cells/mm² in HT). An important finding was that in HT+V, ethanol, used to improve melatonin solubility, was associated with partial protection, based on aEEG recovery from 31 h and reduced TUNEL-positive cells (97.5 cell/mm², midway between HT and HT+M). Increased oligodendrocytes were seen in both HT+V and HT+M versus HT, suggesting ethanol drove oligodendrocyte protection. Melatonin levels reached the putative therapeutic range (15–30 mg/l) at 2 h and 26 h after HI and remained within range 12–18 h after each infusion.

Melatonin was safe and did not lead to any physiological change at this high dose with a rapid infusion rate. Indeed, more dopamine and dobutamine were required to maintain the mean blood pressure in HT and HT+V versus HT+M, but there was no difference in fluid bolus requirements. Our previous piglet studies did not observe any effect of melatonin on blood pressure at doses up to 30 mg/kg^{13,14}. Melatonin therapeutic levels (15–30 mg/l) were reached at 2 h using this rapid infusion started 1 h after HI. It is feasible that in babies with NE, resuscitation would be completed and venous access in place by 1 h, enabling the start of melatonin infusion. Oral melatonin has been used in NE babies undergoing HT and is absorbed^{33,34}, however for rapid achievement of therapeutic levels and optimal protection, intravenous administration is more reliable. In the piglet model we have observed that brain protection is dependent on the time after HI that therapeutic levels are achieved¹⁴. This fits with melatonin's diverse anti-oxidative effects which act upstream in the neurotoxic cascade to prevent

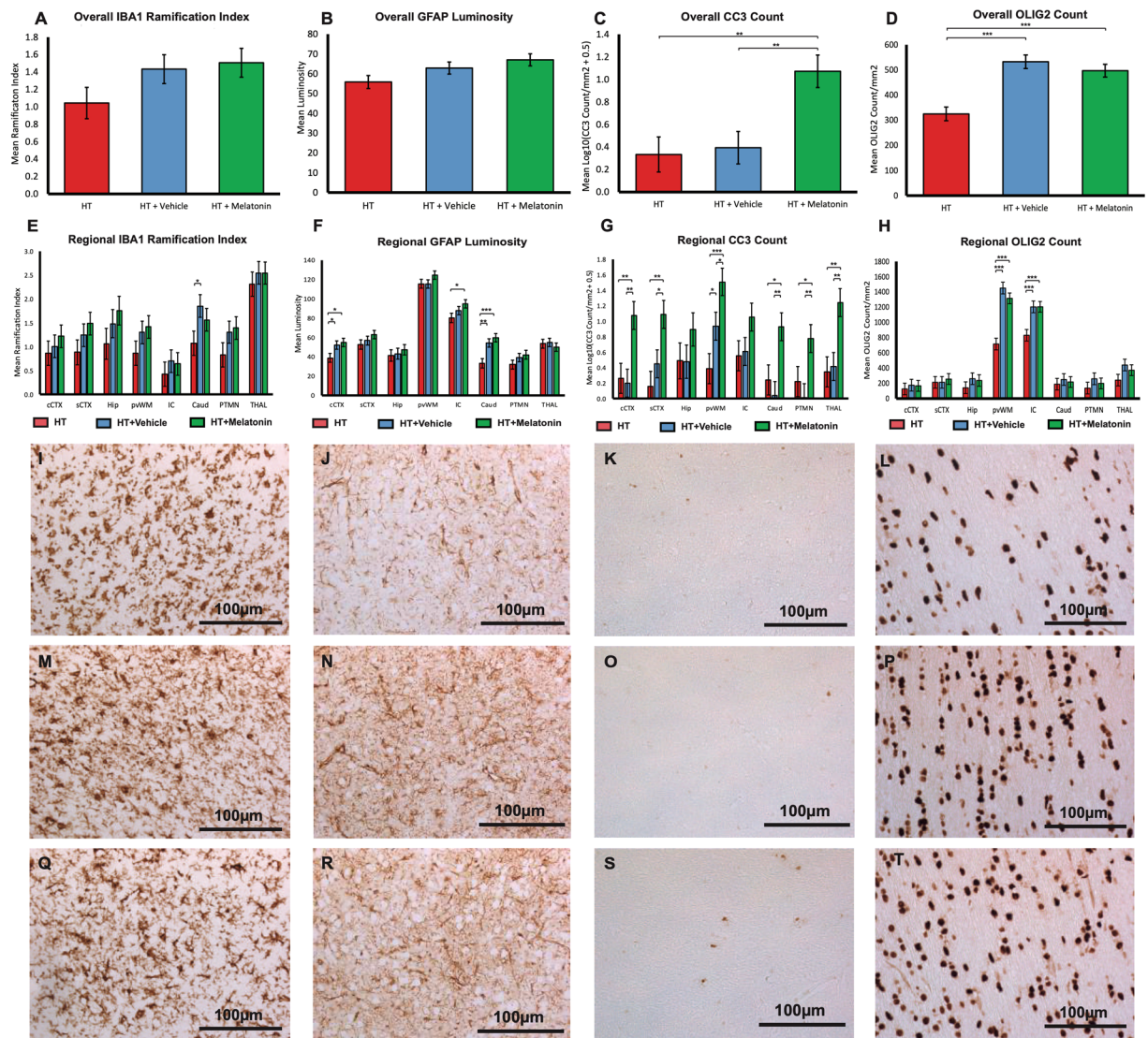


Figure 6. Overall and regional immunohistochemistry for HT, HT+V and HT+M groups for IBA1 ramification index, GFAP luminosity, Cleaved Caspase 3 (CC3) and OLIG2. For IBA1 ramification index, there was no difference in overall ramification index in the HT+M versus HT+V or HT alone (A). On regional analysis the caudate showed an increase in IBA1 ramification index in the HT+V versus HT (E). Representative photomicrographs are shown for HT (I), HT+V (M) and HT+M (Q). There was no difference in overall astrogliosis (GFAP) (B), but localized increases in astrogliosis in the cCTX, IC and caudate as shown (F). Representative GFAP photomicrographs are shown for HT (J), HT+V (M) and HT+M (R). There was an overall increase in CC3 in the HT+M versus HT and HT+M versus HT+V (C). Localized differences with increased CC3 mainly in the HT+M group were compared to both HT+V and HT alone were seen in cCTX, sCTX, pvWM, caudate, putamen, thalamus (G). Representative CC3 photomicrographs are shown for HT (K), HT+V (O) and HT+M (S). There was an overall increase in OLIG2 counts in the HT+M and HT+V compared to HT (D). Localised differences were seen with increased OLIG2 mainly in the IC and pvWM (H). Representative OLIG2 photomicrographs are shown for HT (L), HT+V (P) and HT+M (T). Cingulate cortex = cCTX; Sensorimotor cortex = sCTX; Hippocampus = HIP; Periventricular white matter = PvWM; Internal capsule = IC; Caudate = CAUD; Putamen = PTMN; Thalamus = THAL. Error bars represent standard error *** $p < 0.001$, ** $p < 0.01$, * $p < 0.05$.

free radical-induced oxidative damage to the electron transport chain and mitochondrial DNA³⁵. The newborn infant is at heightened risk for free radical production and injury from oxidative stress. Melatonin and its metabolites work as free radical scavengers, by enhancing anti-oxidant enzyme expression and activity³⁶. Further downstream, melatonin maintains mitochondrial energy production by increasing complex I and IV electron transport chain activity³⁷. Melatonin subsequently prevents apoptosis by preventing nitro-oxidative damage to membrane lipids and inhibiting pro-apoptotic proteins such as BAX; these actions prevent cytochrome c leakage and propagation of intrinsic apoptotic cascades.

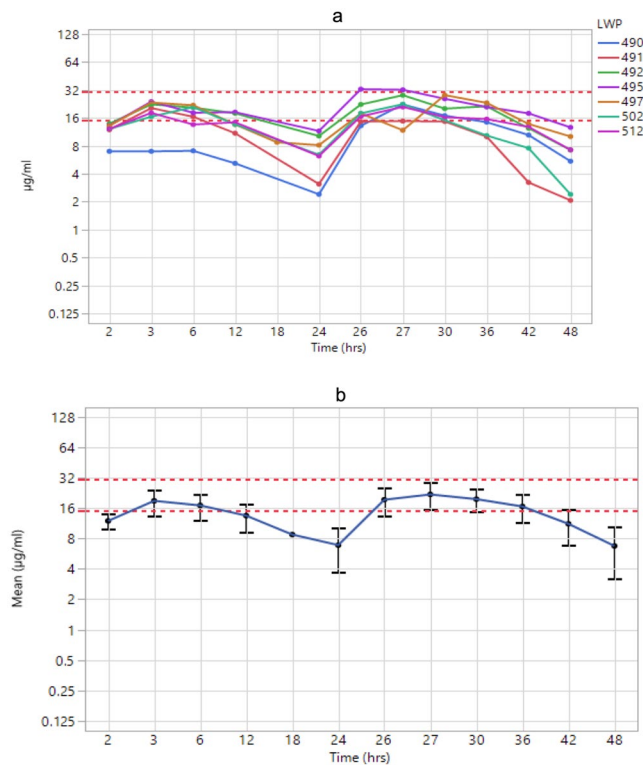


Figure 7. Melatonin plasma levels for individual piglets (A) and Mean (95% CI) Melatonin levels for the HT+M group (B). Melatonin reached the target therapeutic range at 2 h and 26 h after HI (1 h after each infusion started) and remained within this range for up to 18 h.

aEEG is used to monitor babies undergoing HT and the aEEG background voltage and rate of recovery after HI predict outcome^{28,38}. In our study, mean aEEG scores recovered more quickly in HT+M versus HT from 19 h, but surprisingly, we saw improvement in HT+V versus HT from 30 h, corresponding to partial protection also seen with TUNEL-positive cells with HT+V.

MRS biomarkers predict 2-year neurodevelopmental outcome in NE infants^{23,24}. Higher NTP and PCr on ³¹P MRS is associated with better 2-year outcome in clinical studies²⁶; we saw higher ATP and PCr in HT+M versus HT at 48 h. ¹H MRS Lac/NAA is a validated translational biomarker; high levels of thalamic Lac/NAA on MRS in NE babies predict poor 2-year outcomes^{23,24}. We saw lower Lac/NAA on WM MRS with HT+M at 24 and 48 h and in grey matter at 48 h versus HT.

We saw an overall reduced number of TUNEL-positive cells in HT+M versus HT+V and HT. On regional assessment, most protection was observed in IC and pvWM. The high level of protection in these regions may relate to their increased vulnerability to free radical attack, localised severe injury and the early achievement of therapeutic melatonin levels. Although excitotoxic mediated injury may affect all neuronal cells, the myelin-producing oligodendrocytes are particularly vulnerable to injury. We observed that ethanol is likely to have particularly driven the oligodendrocyte protection in IC and pvWM. Other regions with lower levels of injury, such as the hippocampus, putamen and thalamus, also showed preserved oligodendrocytes with ethanol vehicle, but the effect was smaller. Here we did not measure changes in immature or mature oligodendrocytes or myelination. As melatonin is highly lipophilic and easily penetrates brain and organelles, protection is unlikely to be due to vascular factors³⁹. Protection of IC and pvWM with melatonin-augmented cooling as seen here would lead to improvements in cognitive and language outcomes with preservation of brain growth.

Compared to our recent study where therapeutic levels were achieved at 8 h after HI¹⁴, earlier administration of melatonin and therapeutic levels by 2 h in this current study provided enhanced protection. Here, pvWM and IC protection were similar to that achieved with melatonin administered at 10 mins after HI previously where we also saw protection in grey matter regions¹³. Such widespread protection may have been related to the overall more severe injury, expanding the potential for brain protection with melatonin-augmented cooling. Taken together, it is likely melatonin-augmented cooling would be effective in severe injuries and protect both white and grey matter.

There was clear partial protection seen with HT+V (ethanol) whereby mean TUNEL-positive cells/mm² (97.5) were midway between HT (123.8) and HT+M (47.1). There was preservation of oligodendrocytes with HT+V mainly in IC and pvWM and ethanol is likely to have driven protection seen with HT+M in these regions. Ethanol vehicle protection was maximal in IC where injury was most severe. Ethanol has been used as a vehicle for several melatonin neuroprotection pre-clinical and clinical studies, including the piglet¹³, fetal sheep^{15,18,19} and human baby¹⁶. Ethanol modulates GABA and increases GABAergic neurotransmission⁴⁰, this being inhibitory in adults but excitatory in immature brain⁴¹. Neuroprotection with acute low-dose ethanol given up to 4 h after

middle cerebral artery occlusion is described in adult stroke studies with 1.0–1.5 g/kg ethanol producing blood levels of 80–100 g/dl²². In adolescent but not adult rats, high ethanol doses to model binge drinking, upregulated Toll like receptor signalling (TLRS) and increased proinflammatory cytokines⁴², supporting the concept that the immature brain is more vulnerable to ethanol's toxic effects. This increased vulnerability of the developing brain to ethanol, related to inhibition of cortical activity⁴³, mandates ethanol-free medicines for newborn infants⁴⁴.

Therefore, it is likely that ethanol influenced protection in one¹³ but not another¹⁴ previous piglet study and in fetal sheep neuroprotection studies^{15,17–19,45}. Only one fetal sheep study¹⁵ had an ethanol-only control group where ethanol exposure was associated with improved neuronal survival in striatum but reduced survival in hippocampus. In this study, ethanol was associated with suppression of cell proliferation and increased microglial activation, suggesting it is imperative to study melatonin with safe excipients in future studies.

We saw no effect of HT+M versus HT on brain microglial activation^{13,14}. It is possible that, despite no visible change, gene expression was altered¹³. There was no effect of HT+M on GFAP. There were clear increases in CC3 in HT+M versus HT+V and HT. Such CC3 increases were evident throughout the brain; this may reflect caspase's non-apoptotic functions, promoting microglial and lymphocyte function, cell differentiation and autophagy^{46–48}. Increased CC3 expression, not linked the cell death, has been observed in the contralateral hemisphere in a unilateral hypoperfusion P7-rat model⁴⁹. We have previously observed discrepancies between TUNEL-positive cell death and CC3 in our piglet model³². The use of male piglets may partly explain these data; cell death is dimorphic, and in males, apoptosis occurs via caspase-independent pathways^{50,51}. It is likely that the observed TUNEL-positive cell death occurred independently of caspase^{34,7}, such as necrosis, necroptosis, and non-caspase mediated apoptosis⁵². CC3 is therefore a poor cell death/apoptotic marker in this male piglet model.

There are limitations to this study. The inclusion of male piglets only was to minimize variability; inclusion of both sexes is necessary for future development of the model. Termination at 48 h, may not have allowed sufficient time for evolution of apoptotic cell death. Importantly, the rectal temperature was higher in HI versus HI+M (38.9 °C versus 38.2 °C). This is a potential confounder; cooling trials have shown that death/disability is increased 3.6–4.0 fold for each 1 °C increase in temperature in controls⁵³. Duration of temperature rise influences outcome; the difference in our study was only 1 h; over the next 48 h, there was no difference. Nevertheless, temperature rise after HI, exacerbates injury^{54,55}; and this could have worsened brain injury in HT. The blood pH was more alkalotic and PaCO₂ lower at 24 h after HI in HT+M and HT+V versus HT. High variability in CO₂, in particular hypocapnia^{56,57}, is associated with NE adverse outcome and this could exacerbate HT+M brain injury. The contribution of phenobarbitone to brain injury is unclear^{58,59}.

In the search for safe and effective therapies to improve outcomes in NE babies, it will be important to tailor therapies based on sex, inflammatory state and injury severity⁶⁰. Melatonin may be an intervention which targets severe injury if started early enough. With its potent anti-oxidative effects at high dose, low side effect profile and lack of cold-chain storage requirement, melatonin has potential to improve NE outcomes. Phase I safety studies and careful incremental dose studies are needed prior to RCTs. There is a growing interest in the combination of agents with different neuroprotection profiles; melatonin with erythropoietin in preterm brain injury shows promise⁶¹.

In conclusion, 18 mg/kg melatonin 1 h after HI, combined with cooling, reduced brain injury based on faster aEEG recovery 19 h after HI, improved brain energy metabolism on ³¹P and ¹H MRS over 48 h and reduced TUNEL positive cells with particular protection in most severely damaged regions, IC and pvWM. The vehicle, ethanol, used to improve melatonin solubility, led to partial protection based on aEEG recovery 31 h after HI, reduced TUNEL positive cells and increased oligodendrocytes in IC and pvWM versus HT. Melatonin is a promising and safe neuroprotective agent which augments HT if target therapeutic levels are achieved ~2 h after HI. It is imperative to study melatonin with ethanol-free excipients in future pre-clinical and clinical neuroprotection studies.

Received: 15 November 2019; Accepted: 17 February 2020;

Published online: 03 March 2020

References

- Lee, A. *et al.* Intrapartum-related neonatal encephalopathy incidence and impairment at regional and global levels for 2010 with trends from 1990. *Pediatr. Res.* **74**(Suppl 1), 50–72 (2013).
- Kurinczuk, J., White-Koning, M. & Badawi, N. Epidemiology of neonatal encephalopathy and hypoxic-ischaemic encephalopathy. *Early Hum. Dev.* **86**(6), 329–338 (2010).
- Jacobs, S. *et al.* Cooling for newborns with hypoxic ischaemic encephalopathy. *Cochrane Database Sys. Rev.* **1**, CD003311, <https://doi.org/10.1002/14651858.CD003311.pub3> (2013).
- Jary, S. *et al.* Motor performance and cognitive correlates in children cooled for neonatal encephalopathy without cerebral palsy at school age. *Acta Paediatr.* **108**(10), 1773–1780 (2019).
- Lee-Kelland, R. *et al.* School-age outcomes of children without cerebral palsy cooled for neonatal hypoxic-ischaemic encephalopathy in 2008–2010. *Arch. Dis. Child. Fetal Neonatal Ed.* **105**(1), 8–13 (2020).
- Shankaran, S. *et al.* Effect of depth and duration of cooling on deaths in the NICU among neonates with hypoxic ischemic encephalopathy: a randomized clinical trial. *JAMA.* **312**(24), 2629–2639 (2014).
- Shankaran, S. *et al.* Effect of Depth and Duration of Cooling on Death or Disability at Age 18 Months Among Neonates With Hypoxic-Ischemic Encephalopathy: A Randomized Clinical Trial. *JAMA.* **318**(1), 57–67 (2017).
- Macleod, M., O'Collins, T., Horky, L., Howells, D. & Donnan, G. Systematic review and meta-analysis of the efficacy of melatonin in experimental stroke. *J. Pineal Res.* **38**(1), 35–41 (2005).
- Husson, I. *et al.* Melatoninergic neuroprotection of the murine periventricular white matter against neonatal excitotoxic challenge. *Ann. Neurol.* **51**(1), 82–92 (2002).
- Kilic, U., Kilic, E., Reiter, R., Bassetti, C. & Hermann, D. Signal transduction pathways involved in melatonin-induced neuroprotection after focal cerebral ischemia in mice. *J. Pineal Res.* **38**(1), 67–71 (2005).
- Luchetti, F. *et al.* Melatonin signaling and cell protection function. *FASEB J.* **24**(10), 3603–3624 (2010).

12. Carloni, S., Fachinetti, F., Pelizzi, N., Buonocore, G. & Balduini, W. Melatonin acts in synergy with hypothermia to reduce oxygen-glucose deprivation (OGD) in rat hippocampus organotypic slice cultures. *Neonatology*. 2018;in press.
13. Robertson, N. *et al.* Melatonin Augments Hypothermic Neuroprotection in a Perinatal Asphyxia Piglet Model. *Brain*. **136**(Pt 1), 90–105 (2013).
14. Robertson, N. *et al.* Melatonin as an adjunct to therapeutic hypothermia in a piglet model of neonatal encephalopathy: A translational study. *Neurobiol. Dis.* **121**, 240–251 (2019).
15. Drury, P. *et al.* Partial neural protection with prophylactic low-dose melatonin after asphyxia in preterm fetal sheep. *J. Cereb. Blood Flow. Metab.* **34**(1), 126–135 (2014).
16. Fulia, F. *et al.* Increased levels of malondialdehyde and nitrite/nitrate in the blood of asphyxiated newborns: reduction by melatonin. *J. Pineal Res.* **31**(4), 343–349 (2005).
17. Welin, A. *et al.* Melatonin reduces inflammation and cell death in white matter in the mid-gestation fetal sheep following umbilical cord occlusion. *Pediatr. Res.* **61**(2), 153–158 (2007).
18. Yawno, T. *et al.* The Beneficial Effects of Melatonin Administration Following Hypoxia-Ischemia in Preterm Fetal Sheep. *Front. Cell Neurosci.* **11**, 296 (2017).
19. Aridas, J. *et al.* Systemic and transdermal melatonin administration prevents neuropathology in response to perinatal asphyxia in newborn lambs. *J. Pineal Res.* 2018; <https://doi.org/10.1111/jpi.12479>. [Epub ahead of print].
20. Su, F. *et al.* Low-Dose Ethanol Preconditioning Protects Against Oxygen-Glucose Deprivation/Reoxygenation-Induced Neuronal Injury By Activating Large Conductance, Ca²⁺-Activated K⁺ Channels *In Vitro*. *Neurosci. Bulletin*. **33**(1), 28–40 (2017).
21. Aronowski, J., Strong, R., Shirzadi, A. & Grotta, J. Ethanol plus caffeine (caffeinol) for treatment of ischemic stroke: preclinical experience. *Stroke*. **34**(5), 1246–1251 (2003).
22. Wang, F. *et al.* Neuroprotective effect of acute ethanol administration in a rat with transient cerebral ischemia. *Stroke*. **43**(1), 205–210 (2012).
23. Mitra, S. *et al.* Proton magnetic resonance spectroscopy lactate/N-acetylaspartate within 2 weeks of birth accurately predicts 2-year motor, cognitive and language outcomes in neonatal encephalopathy after therapeutic hypothermia. *Arch. Dis. Child. Fetal Neonatal*. <https://doi.org/10.1136/archdischild-2018-315478> (2018).
24. Lally, P. *et al.* Magnetic resonance spectroscopy assessment of brain injury after moderate hypothermia in neonatal encephalopathy: a prospective multicentre cohort study. *Lancet Neurol.* 2018;pii: S1474-4422(18)30325-9.
25. Azzopardi, D. *et al.* Moderate hypothermia within 6 h of birth plus inhaled xenon versus moderate hypothermia alone after birth asphyxia (TOBY-Xe): a proof-of-concept, open-label, randomised controlled trial. *Lancet Neurol.* **S1474-4422**(15), 00347–6 (2015).
26. Azzopardi, D. *et al.* Prognosis of newborn infants with hypoxic-ischemic brain injury assessed by phosphorus magnetic resonance spectroscopy. *Pediatr. Res.* **25**(5), 445–451 (1989).
27. Roth, S. *et al.* Relation between cerebral oxidative metabolism following birth asphyxia, and neurodevelopmental outcome and brain growth at one year. *Dev. Med. Child. Neurol.* **34**(4), 285–295 (1992).
28. Thoresen, M., Hellström-Westas, L., Liu, X. & de Vries, L. Effect of hypothermia on amplitude-integrated electroencephalogram in infants with asphyxia. *Pediatrics*. **126**(1), e131–139 (2010).
29. Lorek, A. *et al.* Delayed (“secondary”) cerebral energy failure after acute hypoxia-ischemia in the newborn piglet: continuous 48-hour studies by phosphorus magnetic resonance spectroscopy. *Pediatr. Res.* **36**, 699–706 (1994).
30. Lingam, I. *et al.* Short-term effects of early initiation of magnesium infusion combined with cooling after hypoxia-ischemia in term piglets. *Pediatr. Res.* **86**(6), 699–708 (2019).
31. de Vries, L. & Hellström-Westas, L. Role of cerebral function monitoring in the newborn. *Arch. Dis. Child. Fetal Neonatal Ed.* **90**(3), F201–207 (2005).
32. Martinello, K. *et al.* Acute LPS sensitization and continuous infusion exacerbates hypoxic brain injury in a piglet model of neonatal encephalopathy. *Sci. Rep.* **15**(91), 10184 (2019).
33. Aly, H. *et al.* Melatonin use for neuroprotection in perinatal asphyxia: a randomized controlled pilot study. *J. Perinatol.* **35**(3), 186–191 (2015).
34. Balduini, W. *et al.* Melatonin pharmacokinetics and dose extrapolation after enteral infusion in neonates subjected to hypothermia. *J. Pineal Res.* **66**(4), e12565 (2019).
35. Reiter, R. *et al.* Melatonin as an antioxidant: under promises but over delivers. *J. Pineal Res.* **61**(3), 253–278 (2016).
36. Galano, A., Tan, D. & Reiter, R. On the free radical scavenging activities of melatonin’s metabolites, AFMK and AMK. *J. Pineal Res.* **54**(3), 245–257 (2013).
37. Leon, J. *et al.* Melatonin and mitochondrial function. *Life Sci.* **75**(7), 765–790 (2004).
38. Csekő, A. *et al.* Accuracy of amplitude-integrated electroencephalography in the prediction of neurodevelopmental outcome in asphyxiated infants receiving hypothermia treatment. *Acta Paediatr.* **102**(7), 707–711 (2013).
39. Paradies, G., Petrosillo, G., Paradies, V. & Reiter, R. Melatonin, cardiolipin and mitochondrial bioenergetics in health and disease. *J. Pineal Res.* **48**(4), 297–310 (2010).
40. Kelm, M., Criswell, H. & Breese, G. Ethanol-enhanced GABA release: a focus on G protein-coupled receptors. *Brain Res. Rev.* **65**(2), 113–123 (2011).
41. Ben-Ari, Y., Khalilov, I., Kahle, K. & Cherubini, E. The GABA excitatory/inhibitory shift in brain maturation and neurological disorders. *Neuroscientist*. **18**(5), 467–486 (2012).
42. Pascual, M., Pla, A., Miñarro, J. & Guerri, C. Neuroimmune activation and myelin changes in adolescent rats exposed to high-dose alcohol and associated cognitive dysfunction: a review with reference to human adolescent drinking. *Alcohol. Alcohol* **49**(2), 187–192 (2014).
43. Lotfullina, N. & Khazipov, R. Ethanol and the Developing Brain: Inhibition of Neuronal Activity and Neuroapoptosis. *Neuroscientist*. **24**(2), 130–141 (2018).
44. Whittaker, A. *et al.* Toxic additives in medication for preterm infants. *Arch. Dis. Child. Fetal Neonatal Ed.* **94**(4), F236–240 (2009).
45. Miller, S., Yan, E., Castillo-Meléndez, M., Jenkin, G. & Walker, D. Melatonin provides neuroprotection in the late-gestation fetal sheep brain in response to umbilical cord occlusion. *Dev. Neurosci.* **27**(2–4), 200–210 (2005).
46. Abraham, M. & Shaham, S. Death without caspases, caspases without death. *Trends Cell Biol.* **14**(4), 184–193 (2004).
47. Northington, F., Chavez-Valdez, R. & Martin, L. Neuronal cell death in neonatal hypoxia-ischemia. *Ann. Neurol.* **69**(5), 743–758 (2011).
48. McComb, S., Mulligan, R. & Sad, S. Caspase-3 is transiently activated without cell death during early antigen driven expansion of CD8(+) T cells *in vivo*. *PLoS One*. **22**(5), e15328 (2010).
49. Villapol, S. *et al.* Unilateral blood flow decrease induces bilateral and symmetric responses in the immature brain. *Am. J. Pathol.* **175**(5), 2111–2120 (2009).
50. Zhu, C. *et al.* Different apoptotic mechanisms are activated in male and female brains after neonatal hypoxia-ischaemia. *J. Neurochem.* **96**(4), 1016–1027 (2006).
51. Charriaud-Marlangue, C. *et al.* Sex differences in the effects of PARP inhibition on microglial phenotypes following neonatal stroke. *Brain Behav. Immun.* **73**, 375–389 (2018).
52. Zhu, C. *et al.* Involvement of apoptosis-inducing factor in neuronal death after hypoxia-ischemia in the neonatal rat brain. *J. Neurochem.* **86**(2), 306–317 (2003).

53. Lupton, A. *et al.* Elevated temperature after hypoxic-ischemic encephalopathy: risk factor for adverse outcomes. *Pediatrics*. **122**(3), 491–499 (2008).
54. Yager, J., Armstrong, E., Jaharus, C., Saucier, D. & Wirrell, E. Preventing hyperthermia decreases brain damage following neonatal hypoxic-ischemic seizures. *Brain Res*. **1011**(1), 48–57 (2004).
55. Mishima, K. *et al.* Effects of hypothermia and hyperthermia on attentional and spatial learning deficits following neonatal hypoxia-ischemic insult in rats. *Behav. Brain Res*. **15**(1-2), 209–217 (2004).
56. Lopez Laporte, M. *et al.* Association between hypocapnia and ventilation during the first days of life and brain injury in asphyxiated newborns treated with hypothermia. *J. Matern. Fetal Neonatal Med.* **32**(8), 1312–1320 (2019).
57. Lingappan, K., Kaiser, J., Srinivasan, C. & Gunn, A. Relationship between PCO₂ and unfavorable outcome in infants with moderate-to-severe hypoxic ischemic encephalopathy. *Pediatr. Res*. **80**(2), 204–208 (2016).
58. Torolira, D., Suchomelova, L., Wasterlain, C. & Niquet, J. Phenobarbital and midazolam increase neonatal seizure-associated neuronal injury. *Ann. Neurol*. **82**(1), 115–120 (2017).
59. Krishna, S., Hutton, A., Aronowitz, E., Moore, H. & Vannucci, S. The effects of adding prophylactic phenobarbital to therapeutic hypothermia in the term-equivalent hypoxic-ischemic rat. *Pediatr. Res*. **83**(2), 506–513 (2018).
60. Hassell, J., Ezzati, M., Alonso-Alconada, D., Hausenloy, D. & Robertson, N. New Horizons for Newborn Brain Protection: enhancing endogenous neuroprotection. *Arch. Dis. Child. Fetal Neonatal Ed.* **100**(6), F541–552 (2015).
61. Robinson, S. *et al.* Extended Combined Neonatal Treatment With Erythropoietin Plus Melatonin Prevents Posthemorrhagic Hydrocephalus of Prematurity in Rats. *Front. Cell Neurosci.* **12**, 322 (2018).

Acknowledgements

This project was supported by the Medical Research Council (MR/M006743/1) and the National Institute for Health Research, University College London Hospitals Biomedical Research Centre.

Author contributions

N.J.R. wrote the grant for funding of the project, N.J.R., X.G., P.G., B.F., B.K. contributed to the concept and design, L.S., C.M., K.M., I.L., A.A.-B., M.H., A.B., M.S., D.P., M.T. acquired the data, N.J.R. and C.M. analyzed and interpreted the data, N.J.R. and L.S. wrote the first draft of the manuscript, all authors have critically reviewed and approved the article.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information is available for this paper at <https://doi.org/10.1038/s41598-020-60858-x>.

Correspondence and requests for materials should be addressed to N.J.R.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2020