Hypothermia is not therapeutic in a neonatal piglet model of inflammation-sensitized hypoxia–ischemia

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BACKGROUND: Perinatal inflammation combined with hypoxia–ischemia (HI) exacerbates injury in the developing brain. Therapeutic hypothermia (HT) is standard care for neonatal encephalopathy; however, its benefit in inflammation-sensitized HI (IS-HI) is unknown.

METHODS: Twelve newborn piglets received a 2 µg/kg bolus and 1 µg/kg/h infusion over 52 h of *Escherichia coli* lipopolysaccharide (LPS). HI was induced 4 h after LPS bolus. After HI, piglets were randomized to HT (33.5 °C 1–25 h after HI, n = 6) or normothermia (NT, n = 6). Amplitude-integrated electroencephalogram (aEEG) was recorded and magnetic resonance spectroscopy (MRS) was acquired at 24 and 48 h. At 48 h, terminal deoxynucleotidyl transferase dUTP nick-end labeling (TUNEL)-positive brain cell death, microglial activation/proliferation, astrogliosis, and cleaved caspase-3 (CC3) were quantified. Hematology and plasma cytokines were serially measured.

RESULTS: Two HT piglets died. aEEG recovery, thalamic and white matter MRS lactate/N-acetylaspartate, and TUNEL-positive cell death were similar between groups. HT increased microglial activation in the caudate, but had no other effect on glial activation/proliferation. HT reduced CC3 overall. HT suppressed platelet count and attenuated leukocytosis. Cytokine profile was unchanged by HT.

CONCLUSIONS: We did not observe protection with HT in this piglet IS-HI model based on aEEG, MRS, and immunohistochemistry. Immunosuppressive effects of HT and countering neuroinflammation by LPS may contribute to the observed lack of HT efficacy. Other immunomodulatory strategies may be more effective in IS-HI.

Pediatric Research (2022) 91:1416–1427; https://doi.org/10.1038/s41390-021-01584-6

IMPACT:
- Acute infection/inflammation is known to exacerbate perinatal brain injury and can worsen the outcomes in neonatal encephalopathy.
- Therapeutic HT is the current standard of care for all infants with NE, but the benefit in infants with coinfection/inflammation is unknown.
- In a piglet model of inflammation (LPS)-sensitized HI, we observed no evidence of neuroprotection with cooling for 24 h, based on our primary outcome measures: aEEG, MRS Lac/NAA, and histological brain cell death.
- Additional neuroprotective agents, with beneficial immunomodulatory effects, require exploration in IS-HI models.

INTRODUCTION
Therapeutic hypothermia (HT) is standard care for neonates with intrapartum-related neonatal encephalopathy (NE) under intensive care settings. Cooling to 33.5 °C for 72 h improves survival free from disability in the short and long term. However, cooling is ineffective in preventing disabilities in a proportion of infants (number needed to treat, 7 (95% confidence interval (CI) 5–10) for reduction in death or neurodisability at 18 months). It is clear that, while disability from cerebral palsy (CP) in NE survivors reduces from 35% to as low as 16% with HT, some level of...
intellectual impairment may remain even in the absence of CP.4
Attenuated efficacy of HT may be due to severity of illness, timing
of insult in relation to initiation of HT, or sensitization with coexisting infection/inflammation.
In preclinical laboratory studies demonstrating HT efficacy, NE
is typically modeled with hypoxic–ischemic (HI) brain injury. However,
in the clinical setting, the etiology of NE is multifactorial, with contributions from antenatal and placental pathology,5
genetic susceptibility,6 and perinatal infection/inflammation7 in addition to sentinel HI events.8 Preclinical9,10 and clinical11,
12 studies suggest that acute infection/inflammation sensitizes the
brain to subsequent HI, lowering the injury threshold and worsening outcome.7,13 Mortality amongst infants with
combined NE and infection is increased compared with NE alone.13 We have recently demonstrated an increase in mortality
and brain cell death in Escherichia coli (E. coli) LPS-sensitized hypoxic brain injury in the piglet model, compared with either E.
coli LPS or hypoxia alone.10 These data concur with studies of small animal models of inflammation-sensitized HI (IS-HI).9,14,15
Given the increase in the severity of brain injury and mortality, effective therapeutic interventions are urgently needed for
infants with IS-HI.

The benefit and safety of cooling infants with IS-HI are unclear. HT induces alterations in leukocyte number, microglial
16 cell17 activation,18 cytokine/chemokine profiles,19,20 and hemodynamics,20 it is plausible that immunomodulatory effects of HT
may not be beneficial in the context of inflammation sensitization. In a small prospective study of placental histology
relative to magnetic resonance imaging (MRI) outcomes, HT was less protective for infants with NE where placental histology
showed chorioamnionitis.20 In low-resource settings, where rates of IS-HI are high, HT may be less effective, and potentially
harmful.21,22 In addition, HT has been shown to be deleterious in the presence of meningitis23 and sepsis24 in adult clinical studies. These observations emphasize the critical need for preclinical animal models that assess the safety and efficacy of HT and other neuroprotective agents in IS-HI. Rodent IS-HI models suggest variable, pathogen-dependent, efficacy of HT.14,15,25 This has not previously been examined in a large animal model.
The aim of this study was to assess the safety and efficacy of HT in an E. coli LPS-sensitized HI piglet model. The piglet model
has strong similarities to newborn infants with NE in terms of the evolution and pattern of injury after HI and neuropathology.
26,27 We hypothesized that HT would not provide neuroprotection in IS-HI based on our primary outcome measures:
(i) amplitude-integrated electroencephalogram (aEEG) background activity recovery over 48 h; (ii) proton (1H)
magnetic resonance spectroscopy (MRS) lactate/N-acetylaspar
tate ratio (Lac/NAA) at 48 h in the subcortical white matter and thalamus; and (iii) terminal deoxynucleotidyl transferase dUTP
nick-end labeling (TUNEL) immunolabelling of histological cell
death over the whole-brain and eight brain regions at 48 h after HI.

METHODS
This study was conducted in accordance with the UK Home Office
Guidelines [Animals (Scientific procedures) Act, 1986] and complies with ARRIVE guidelines. The Ethics Committee of
University College London approved the study.

Sample size calculation
Using previous data from our piglet study of LPS plus hypoxia (geometric least-squares mean standard deviation of 17), we
estimated that six piglets per intervention group were required to detect a difference in TUNEL-positive cells of >30 cells/mm², using
a significance threshold of 5 and 80% power.

Animal experiments and surgical preparation
Male large white piglets aged ≤36 h were sedated with intramuscular midazolam (150 μg) and anesthetized with 2–3%
(v/v) isoflurane. Piglets underwent tracheostomy and mechanical ventilation (SLE 2000 Infant Ventilator, Surrey, UK). The bilateral
central carotid arteries were surgically isolated and encircled by inflatable occluders (OC2A, In Vivo Metric). A 2-French venous
catheter (Vygon, Swindon, UK) was inserted into the axillary or brachial vein for later infusion of LPS. Umbilical catheters were
sutured; a 4-French double-lumen umbilical venous catheter (Vygon) was used for maintenance fluids (10% dextrose,
60 ml/kg/day, reduced to 40 ml/kg/day post HI), fentanyl (3 μg/kg/h), and antibiotics (benzylpenicillin 50 mg/kg every 12 h and
gentamicin 2.5 mg/kg every 24 h); and a 2.5-French umbilical arterial catheter (Vygon) for intermittent blood sampling (Abbot
Laboratories, UK) and continuous monitoring of heart rate and
mean arterial blood pressure (MABP). Arterial catheters were
maintained with 0.3 ml/h heparinized saline (1 IU/ml in 0.9% saline solution). Core (rectal) temperature was maintained in
the normothermic range during surgery using a radiant warmer. The duration of surgery was 1–2 h.

Following surgery, piglets were positioned prone within a
bespoke MRI-compatible transport incubator. Intensive care
support and physiological monitoring (SA Instruments, London,
UK) was continued throughout the study. Ventilation was adjusted
to maintain partial pressure of oxygen (PaO₂) and carbon dioxide
(PaCO₂) at 8–13 and 4.5–6.5 kPa, respectively. MABP was
supported with crystalloid (0.9% saline, Baxter, 10 ml/kg bolus)
and inotropes (dopamine 5–25 μg/kg/min, dobutamine 5–20 μg/
k/min, noradrenaline 0.1–1 μg/kg/min, adrenaline 0.1–1 μg/kg/
min) to target MABP >35 mm Hg. Anesthesia was maintained
throughout with fentanyl (3 μg/kg/h infusion) and isoflurane
(2–3% v/v). Hyperkalemia (potassium > 7.0 mmol/l) was treated
with 4 μg/kg intravenous salbutamol over 10 min.

EEG acquisition
Multichannel six-lead EEG (Nicolet, CareFusion, Madison, WI) was
commenced immediately post surgery and maintained through-
out the study. aEEG recordings were retrospectively scored hourly using the pattern classification.26 Flat trace was assigned a score of
0; continuous low voltage 1; burst suppression 2; discontinuous
normal voltage 3; and continuous normal voltage 4. Scores were
averaged every 6 h. Electrographic seizures were treated with
phenobarbitone, initially 20 mg/kg, followed by 10 mg/kg doses
for subsequent seizures, up to a maximum of 40 mg/kg.

Broadband near-infrared spectroscopy
A broadband near-infrared spectroscopy (bNIRS) system (Mini-
CYRIL, Cytochrome Research Instrument and Application System,
UCL, UK) was used to measure the concentration of oxidized
cytochrome c oxidase (oxCCO).

LPS administration
Baseline physiological, bNIRS, and EEG data were captured prior to
administration of LPS. All piglets received a bolus of 2 μg/kg LPS
(E. coli-derived, Sigma LPS O55:B5, St. Louis, MO), followed by a
continuous infusion of 1 μg/kg/h LPS for the duration of the
experiment (total 52 h LPS).

Transient HI
Four hours after the LPS bolus, the carotid occluders were inflated
and the fraction of inspired oxygen (FiO₂) was reduced to induce
transient HI. FiO₂ was decreased sequentially to 6% over the first 3
min and maintained. The insult was titrated using continuous
MABP, EEG, and bNIR monitoring, and arterial blood gas was
measured at 5-min intervals. Target insult parameters included
MABP between 26 and 30 mm Hg, a 3-fold reduction in oxCCO
from baseline, a sustained isoelectric EEG, and metabolic acidosis

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Magnetic resonance spectroscopy
MRS was acquired on a Philips Achiever 3 T (Philips Healthcare, Best, The Netherlands) scanner at 24 and 48 h after HI. 1H magnetic resonance spectra were acquired using a Philips 32-channel head coil. Chemical shift imaging (CSI) based on a two-dimensional point-resolved spectroscopy excitation used an 8 × 8 matrix with a voxel size of 7.5 × 7.5 mm² and slab thickness of 10 mm. The repetition time was 2 s and the echo time was 288 ms. 1H CSI data was processed using spectroscopy analysis software, TARQUIN, yielding signal amplitudes for NAA and lactate. Two dimensional point-resolved spectroscopy excitation used an 8 × 8 matrix with a voxel size of 7.5 × 7.5 mm² and slab thickness of 10 mm. The repetition time was 2 s and the echo time was 288 ms.

Hematology and cytokine analysis
Arterial blood was collected at baseline, 4 h after LPS bolus, at the end of HI, and at 3, 12, 24, and 48 h post HI. Cerebrospinal fluid (CSF) was obtained by lumbar puncture at baseline and at 48 h. Hematology samples were stored at 4 °C for a maximum of 72 h prior to complete blood film examination (Royal Veterinary College Diagnostic Laboratory, Hawkshead Campus, UK). Hemoglobin (Hb), total white cell (WBC), neutrophil, lymphocyte, monocyte, and platelet count were analyzed. Serum and CSF samples were frozen (−20 °C) for later protein analysis. Multiplex porcine enzyme-linked immunosorbent assay (ELISA) was undertaken to quantify interleukin-1β (IL-1β), IL-6, IL-8, and tumor necrosis factor-α (TNFα) (Q-Plex Porcine Cytokine—High sensitivity (4-Plex), ref.: 119149PC, Tebu-Bio, France), while a porcine ELISA was utilized for IL-10 (Porcine IL-10 ELISA Kit (Cloud-Clone Corp., ref. SEA0566P), Tebu-Bio, France).

Brain histology
Forty-eight hours after HI, piglets were euthanized and histological brain specimens were obtained and prepared as previously described. Cell death was quantified by nuclear DNA fragmentation using histochemistry with TUNEL. Glial activation was quantified using the presence of astrocyte glial fibrillary acidic protein (GFAP) and microglial ionized calcium-binding adaptor molecule 1 (IBA1) immunoreactivity. Cleaved caspase-3 (CC3) immunoreactivity was quantified. Histological analysis was performed by investigators blind to the treatment group. Eight brain regions were assessed (cingulate cortex, sensorimotor cortex, hippocampus, periventricular white matter, internal capsule, caudate nucleus, putamen, and thalamus). For each region, TUNEL-positive nuclei were counted in six fields (three per section, at ×40 magnification, with an area of 0.066 mm²) and the average converted into counts per mm². IBA1-positive cell body count was similarly performed. In addition, IBA1 ramification index was calculated in six fields per region, at ×40 magnification, using a 0.049 mm × 0.049 mm square grid. The microglial ramification index was calculated as (B²/C), where C is the number of cell bodies within the grid, and B is the number of branches crossing the three horizontal and three vertical gridlines. CC3 immunoreactive cells were counted in four fields per region (at ×20 magnification with an area of 0.164 mm²) and the average converted into counts per mm². To quantify the GFAP immunoreactivity optical luminosity values were calculated by deducting mean brightness values of the tissue (four fields per region at ×20 magnification) from the mean brightness of the blank slide.

(pH < 7.3, lactate > 10). FiO₂ was liberalized in the event of MABP and oxCCO values beyond the target range. The duration of insult was anticipated to be 20 min. At the end of the insult, occluders were deflated, FiO₂ normalized to 21%, and the animal resuscitated as necessary.

Experimental groups: temperature management
Following insult, piglets were randomized, using a computer-generated randomization sequence and opaque sequentially numbered envelopes, to two groups: (i) LPS, HI, and normothermia (NT) and (ii) LPS, HI, and HT (Fig. 1). NT piglets had core temperature maintained in the normothermic range (38.0–39.0 °C) throughout the study. HT piglets were actively cooled to 33.5 °C from 1 to 25 h post insult, and then rewarmed over 10 h at an incremental rate of 0.5 °C/h. HT piglets were subsequently maintained in the normothermic range until the end of the study. Temperature was maintained within target ranges using a water mattress (Tecotherm TSmed200, Inspirational Healthcare, Crawley, UK) manually adjusted according to continuous rectal temperature. The mattress was wrapped around the trunk and limbs.

Magnetic resonance spectroscopy
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Statistical methods
Non-normally distributed data were log-transformed prior to analysis. Using SAS JMP® Pro v14.0.0, analysis of variance models were fitted to give estimates of expected treatment group mean values and difference between means. For TUNEL and histology measurements, the effects of treatment, region, and the interaction between them were fitted to the mean result for each subject averaged across replicate measurements within each region. For EEG, MRS, cytokines, and hematology, a model with terms for treatment, time interval (as a factor) and the interaction between them, and a random subject effect to take into account the repeated-measures structure was fitted to the results for each subject averaged across each time interval. Using the model, a comparison to baseline was calculated for hematology values. GraphPad Prism® v8 was used to calculate Pearson’s correlation between TUNEL and CC3 cell count. Cytokine change from baseline was calculated using Wilcoxon’s matched-pairs signed-rank test. Using IBM SPSS® v22, physiological data and clinical outcomes were analyzed using t test or Fisher’s exact test as appropriate.

RESULTS
Baseline characteristics and physiological parameters
Piglet weight ranged from 1800 to 2250 g. Physiological parameters are shown in Table 1. At baseline groups were similar. During the time epochs 1–25 and 25–35 h, which encompass cooling and rewarming, the HT subjects had lower core temperatures as expected (p < 0.001 and 0.002, respectively).

HI insult
Insult severity, as determined by the duration of insult, MABP < 30, isoelectric EEG, area under the curve (AUC) FiO2, AUC oxCCO, and end HI acid–base status, was similar between groups (Table 2).

Survival and clinical illness severity
During active cooling, HT animals had a lower mean heart rate than NT animals (171 (SD 25) b.p.m. compared with 235 (SD 21) b.p.m., p = 0.001). Mean heart rate was also lower during rewarming for HT animals (p = 0.02) (Table 1). Mean MABP, inotropic support, and saline bolus requirement were not different between groups (Table 3). Mean infusion rates of all inotropes were higher in the HT group; however, confidence intervals are wide. HT had higher mean blood glucose at 24 h (10.3 (SD 3.6) mmol/L compared with 6.5 (SD 1.2) mmol/L, p = 0.033). Three of six NT and four of six HT subjects had hyperkalemia necessitating treatment (p = 0.5).

Two of six piglets died prior to experiment completion in the HT group, compared with none in the NT group (p = 0.23). The two animals had refractory hypotension and worsening metabolic acidosis at the time of cardiac arrest (30 and 33 h). Both were rewarmed early due to the severity of hypotension (at 16 and 12.5 h). One had severe hyperkalemia (K+ > 9). Both had macroscopic necrosis to abdominal solid organs including the liver, kidneys, and bowel.

TUNEL
Mean whole-brain TUNEL-positive cell counts were similar between the NT and HT groups (p = 0.97) (Fig. 2a). The regional assessment was also similar between groups. There was a trend towards greater histological cell death in the internal capsule for the TH group (p = 0.064).

Cleaved caspase-3
HT reduced overall CC3 (Fig. 2b) compared with NT (p = 0.04). The regional analysis demonstrates lower CC3 in the internal capsule,
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Table 1. continued

|                      | Normothermia Mean (SD) | Hypothermia Mean (SD) |
|----------------------|------------------------|-----------------------|
| Lactate (mmol/L)     |                        |                       |
| Baseline             | 3.6 (1.7)              | 2.9 (1.2)             |
| 4 h LPS              | 4.1 (2.0)              | 3.4 (1.3)             |
| End of insult        | 13.1 (1.7)             | 11.0 (2.2)            |
| 12 h                 | 3.0 (0.6)              | 6.6 (5.0)             |
| 24 h                 | 2.3 (0.8)              | 6.7 (7.1)             |
| 48 h                 | 1.1 (0.2)              | 1.3 (0.3)             |
| Glucose (mmol/L)     |                        |                       |
| Baseline             | 4.9 (1.3)              | 5.2 (1.2)             |
| 4 h LPS              | 6.0 (2.2)              | 5.6 (0.5)             |
| End of insult        | 8.9 (3.0)              | 7.9 (1.0)             |
| 12 h                 | 6.8 (1.2)              | 11.5 (5.6)            |
| 24 h                 | 6.5 (1.2)              | 10.3 (3.6)*           |
| 48 h                 | 4.5 (1.2)              | 4.9 (0.7)             |
| Potassium (mmol/L)   |                        |                       |
| Baseline             | 4.1 (0.7)              | 4.5 (0.4)             |
| 4 h LPS              | 4.5 (0.4)              | 5.0 (0.4)             |
| End of insult        | 4.5 (0.5)              | 5.2 (0.8)             |
| 12 h                 | 6.3 (0.7)              | 6.3 (1.0)             |
| 24 h                 | 6.7 (0.7)              | 6.7 (1.4)             |
| 48 h                 | 5.7 (1.2)              | 6.2 (1.3)             |

MABP mean arterial blood pressure, BE base excess.

*P < 0.05, **P < 0.005.

Table 2. HI insult severity measures.

|                      | Normothermia Mean (SD) | Hypothermia Mean (SD) |
|----------------------|------------------------|-----------------------|
| Duration HI (min)    | 20.7 (1.4)             | 21.2 (2.1)            |
| AUC FiO₂ (%)         | 295.8 (20.1)           | 290.5 (43.2)          |
| Duration of EEG < 5 μV (min) | 15.5 (2.9)          | 17.7 (1.8)            |
| Duration of MABP < 30 (min) | 7.3 (2.9)            | 10.0 (3.8)            |
| AUC oxCCO            | 146 (25)               | 124 (50)              |
| End of HI arterial gas |                       |                       |
| pH                   | 7.22 (0.11)            | 7.25 (0.13)           |
| pCO₂ (kPa)           | 6.3 (1.8)              | 5.6 (0.9)             |
| pO₂ (kPa)            | 3.6 (0.4)              | 3.7 (0.9)             |
| Base excess          | −9.0 (2.9)             | −8.7 (4.9)            |
| Lactate (mmol/L)     | 13.1 (1.7)             | 11.0 (2.2)            |

No comparisons were p < 0.05

HI hypoxia-ischemia, EEG electroencephalogram, MABP mean arterial blood pressure, oxCCO oxidized cytochrome c oxidase.

caudate nucleus, and hippocampus (p = 0.006, 0.01, and 0.02, respectively). There was no correlation between TUNEL-positive cell count and CC3-positive cell count (r = −0.42, 95% CI −0.54 to −0.30).

Glial fibrillary acidic protein
Mean whole-brain and regional GFAP luminosity was similar between the NT and HT groups (p = 0.89) (Fig. 2b).

Table 3. Inotrope and saline bolus use throughout the experiment.

|                      | Normothermia Mean (SD) | Hypothermia Mean (SD) |
|----------------------|------------------------|-----------------------|
| Dopamine (μg/kg/min) | 10.1 (4.8)             | 15.6 (6.4)            |
| Dobutamine (μg/kg/min) | 2.8 (4.3)              | 4.5 (6.9)             |
| Noradrenaline (ng/kg/min) | 16 (37)               | 96 (149)              |
| Adrenaline (ng/kg/min) | 0 (0)                  | 484 (811)             |
| 10 ml/kg Saline bolus (n) | 0.7 (0.8)             | 0.8 (1.3)             |

No comparisons were p < 0.05.

Ionized calcium-binding adaptor molecule 1
IBA1-positive cell count was no different between the NT and HT group throughout the whole brain (p = 0.41) and in each brain region (p ≥ 0.08) (Fig. 2b). Overall microglial activation state (ramification index) was also similar between groups (p = 0.81). There was an increase in microglial activation (lower ramification index) in the caudate nucleus with HT (p = 0.029). Microglia for both groups appeared partially activated, with enlarged cell bodies, a reduced number of processes, and thickening of the remaining processes.

Representative photomicrographs from the internal capsule area at ×40 magnification for all histological stains are shown in Fig. 2c.

Amplitude-integrated electroencephalogram
All subjects had a normal (score 4) aEEG at baseline (Fig. 3). The mean aEEG score for both groups was suppressed following HI and remained suppressed throughout the experiment. There was no effect of HT on aEEG recovery. Two of the six NT and one of the six HT piglets had electrographic seizures (p = 0.5). Seizure onset was at 12 and 15 h post HI for the two NT piglets, and at 36 h for the one HT piglet with seizures. Seizure burden was no different between groups (range 0–47 min) (p = 0.75). Three of the six NT and one of the six HT piglets were treated with phenobarbitone. Phenobarbitone use and dose per kg were not different between groups (p ≥ 0.34).

Magnetic resonance spectroscopy
Lac/NAA was no different between the cooled and normothermic piglets at either time point, in either the white matter or thalamus (Fig. 4, p ≥ 0.35 for all comparisons). Note that due to early death, two of six HT subjects were not scanned at 48 h. Compared to all others, these two deceased piglets had the highest thalamic Lac/NAA at 24 h.

Hematology
In both groups, platelet count fell below baseline by time 0 (p ≤ 0.0024) (Fig. 5). NT platelet count returned to baseline levels, before a late thrombocytosis at 48 h post HI (p < 0.001). HT platelet count continued to fall below baseline for the duration of cooling therapy (p ≤ 0.007), before a later recovery to baseline at 48 h. At 24 and 48 h after HI, platelet count was suppressed in the HT group compared with NT (p = 0.005 and 0.001, respectively).

The WBC increased from baseline for NT subjects at 3, 12, and 24 h (p ≤ 0.016) (Fig. 5). This increase was attenuated in HT subjects. There was no significant difference between WBC for NT and HT animals at any single time point. Neutrophils were the predominant leukocyte present and the analysis closely mirrored that for WBC. There was no effect of cooling or time on lymphocyte number. The monocyte count was raised at 24 h in the HT group compared to baseline (p = 0.002). Monocytes were otherwise unaltered by HT or time.

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Fig. 2  Brain histology.  

a  TUNEL histology. There was no difference between whole-brain or regional estimated least-squares mean (LSM) TUNEL-positive cells per mm².  

b  LSM whole-brain and regional cleaved caspase-3 (CC3), GFAP, IBA1-positive cell count, and IBA1 ramification index. Hypothermia reduced CC3 throughout the brain and in the internal capsule, caudate nucleus, and hippocampus. Hypothermia did not alter astrogliosis. Cooling had no effect on the overall microglial number or activation state. There was an increase in microglial activation (lower ramification index) in the caudate with HT.  

c  Representative sections for each stain (TUNEL, CC3, GFAP, and IBA1) from the internal capsule are shown at ×40 magnification. cCTX cingulate cortex, sCTX sensorimotor cortex, HIP hippocampus, PvWM periventricular white matter, IC internal capsule, CAUD caudate, PTMN putamen, THAL thalamus. Data are displayed as analyzed, on a log 10 scale. Error bars represent 95% CI. *p < 0.05, **p < 0.01.
There was no intergroup difference for Hb at any time point. NT Hb was stable and then fell below baseline from 24 h (p = 0.002). Hb increased from baseline in the HT group at 4 h after LPS and at 3 and 12 h after HI (p ≤ 0.012) (Fig. 5). At 48 h, both HT and NT Hb were below baseline (p ≤ 0.022).

Cytokines
The pattern of IL-6, IL-8, and TNFα in the plasma and CSF is shown in Fig. 6. Plasma IL-6, IL-8, and TNFα increased from baseline pre-HI (p ≤ 0.0034) in response to LPS. HT plasma IL-6 was higher than NT at 3 h post HI (p = 0.018). There were no other differences between groups and no difference in CSF cytokines. IL-1β and IL-10 were below the limit of detection for the majority of samples and therefore were not analyzed.

**DISCUSSION**

In this piglet model of IS-HI, we did not observe protection with 24 h HT (from 1 to 25 h after HI) based on pre-defined primary outcome measures; aEEG, MRS, and histological cell death (TUNEL-positive cells). Microglial activation and glial cell proliferation were unaltered, although microglial activation was increased in the caudate nucleus with HT. Compared to NT, HT suppressed the platelet and white cell count response. Plasma and CSF cytokine response to LPS and HI was largely unchanged with HT. HT reduced CC3 throughout the brain; in our model, CC3 is a poor marker of cell death,36 and in this and previous studies, a regional reduction in CC3 is associated with increased TUNEL-positive cells.

These results differ from non-sensitized HI piglet models, where HT for 24 h started at 2 h after HI reduced brain injury based on TUNEL-positive cell counts17 and HT for 12–24 h started immediately after HI ameliorated secondary energy failure, reduced seizure activity, and improved neurological function and histopathology scores.31,32 In the present study, a trend towards increased TUNEL-positive cell death in the internal capsule with HT was observed. Concurring with the histological data, aEEG and 1H MRS were not improved by HT. Comparing NT and HT groups, there was no difference in aEEG score or seizure burden. The mean aEEG score for HT trended upwards from 31 h; this may be attributed to the loss of two subjects who had flat traces (score 0). In NE babies, a recovery of aEEG within 48 h is associated with a good outcome.63 1H MRS Lac/NAA peak area ratios in the white matter and thalamus were not improved by HT. In non-sensitized piglet models, HT for 24 h improves MRS biomarkers, including Lac/NAA31,34,35 which is a robust marker of 2-year neurodevelopmental outcome in NE babies.36

Suppression of microglial activity is a favorable marker7 and a key mechanism by which HT is neuroprotective.37,38 Reduced microglial activation following HT is observed in non-sensitized HI models, including the pig,57 sheep,38 and rodent.39 In this IS-HI model, microglial number and morphology were not suppressed by HT; indeed, in the caudate nucleus, microglial morphology was indicative of a pro-inflammatory activation state with HT, possibly reflecting increased injury. Our findings concur with those from a rodent LPS-sensitized HI cooling model.40 The balance between activated microglia with a pro-inflammatory phenotype and those with an anti-inflammatory restorative phenotype is complex and needs further study.7,12,41

Astrogliosis, measured by GFAP, was not impacted by HT. In non-sensitized HI models, HT reduces astroglial activation.39 LPS enhances astrogliosis by stimulating the release of TNFα, IL-1α, and IL-6, mediators of astrocyte proliferation and activation.42,43 A failure of HT to suppress astrogliosis has also been shown in an LPS-sensitized rodent model.46

CC3 was reduced by HT, both overall and specifically in the internal capsule, caudate nucleus, and hippocampus. As in our previous LPS + hypoxia study10 and other recent piglet neuroprotection studies,44,45 CC3 was not associated with TUNEL-positive cell death. The use of male piglets may partly explain these data; cell death is sexually dimorphic, and in males, cell...
resulting in apoptotic cell death. This CC3 response to LPS may then conclude that CC3 is, therefore, a poor marker of cell death in this model. LPS is known to increase CC3 without demonstrating in neonates undergoing HT and was associated with poorer outcome. Thus, alterations in immune cell number may be an important factor in the efficacy and safety of TH in IS-HI.

HI, HT, and LPS each influence the hematological response. Infection/inflammation triggers thrombocytopenia and leukocytosis in neonates and piglets. HI also stimulates leucocytosis. HT is known to reduce circulating platelets and WBC. A meta-analysis of HT clinical trials demonstrated an increase in thrombocytopenia (relative risk (RR) 1.21 (95% CI 1.05–1.40)) with HT, and an increase in leukopenia with whole-body HT (RR 5.7 (95% CI 1.02–31.82)). Similarly, in this study, HT suppressed both platelet and leukocyte (neutrophil) count. The impact of this suppression is uncertain in the setting of IS-HI. In a mouse model of LPS-sensitized HI, neutrophils were central to injury pathways, leading to leukocyte infiltration, pro-inflammatory cytokine production, and brain atrophy. By contrast, an increase in circulating neutrophils, and later lymphocytes, are necessary components of the innate and adaptive immune response to infectious stimuli. Leukopenic “immune paralysis” has been demonstrated in neonates undergoing HT and was associated with poorer outcome. Thus, alterations in immune cell number may be an important factor in the efficacy and safety of TH in IS-HI.

Many immune cells are functionally heat-sensitive and a key mechanism of HT neuroprotection is inhibition of the pro-inflammatory cascade. In a cohort of babies with NE, HT infants had reduced serum IL-6 and IL-4, and increased vascular endothelial growth factor (VEGF) compared with NT infants. In the same cohort, there was a negative correlation between the duration of cooling prior to sampling, and levels of IL-6, IL-10, TNFα, and interferon-gamma (IFNγ). In another cohort, HT lowered IL-6, and increased IL-10 at 48 h. Conversely, in a phase II cooling trial, cooled infants had a biphasic pattern of elevated IL-6, IL-8, IL-10, and chemokine (C-C motif) ligand 2 in comparison with NT infants although another prospective cohort showed a trend towards an increase in TUNEL-positive cell death. HI cooling trial, cooled infants had a biphasic pattern of elevated IL-6, IL-8, IL-10, and chemokine (C-C motif) ligand 2 in comparison with NT infants although another prospective cohort showed a trend towards an increase in TUNEL-positive cell death.

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Fig. 4 MRS. There is no effect of hypothermia on the ratio of lactate to N-acetylaspartate (Lac/NAA) in the thalamus or white matter at 24 or 48 h after HI. In all subjects, log 10 Lac/NAA data points are shown in the graphs on the left-center, and are summarized using log 10 least-squares (LS) means plot with 95% CI error bars in the boxes on the right. The crosses indicate the two HT piglets who died prior to the 48 h scan.
hypoxia alone. In this study, the two cardiac arrests were preceded by refractory hypotension, which may have been contributed to by HT-induced bradycardia. Indeed, in a meta-analysis of cooling trials, HT increased sinus bradycardia, although there was no increase in hypotension or major arrhythmias. The HT group was hyperglycemic during the cooling phase. HT reduces insulin secretion and increases insulin resistance in preclinical models and adult HT studies. Deranged postnatal glycemic control is an important risk factor for the adverse outcome as seen in the Coolcap study and is associated with specific patterns of brain injury in NE. Two piglets in the HT group had macroscopic abdominal organ necrosis, including the bowel, at post mortem.

There are limitations to this study. By using E. coli LPS to stimulate inflammation, we modeled only gram-negative perinatal infection. Rodent models of IS-HI, comparing LPS with PAM3CSK4 (a toll-like receptor (TLR) 2 agonist, which simulates gram-positive infection), demonstrate the differences in temperature, cytokine expression, CC3, and glial cell response. Importantly, the efficacy of HT was pathogen-dependent; HT was neuroprotective with PAM3CSK4 IS-HI, but not with LPS IS-HI. Two of six piglets in the HT group were rewarmed early due to refractory hypotension,
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consistent with clinical practice. These two piglets died prematurely, reducing the available outcome data at later time points. With reduced numbers, the study may have been underpowered to demonstrate differences in later outcome measures, including aEEG and MRS. Histological outcomes measures were available for all 12 subjects, including the two piglets that died prematurely. Only male piglets were used to reduce intergroup variation. We therefore were not able to examine the impact of sex on outcome. Future studies using this model of IS-HI will be conducted using both male and female piglets. The porcine immune system is an ideal model for immunology research, resembling humans by >80%. However, there are differences, for example, piglets are born without transplacental antibodies to and preventing injury, and the complex mechanisms underlying the evolution of HI injury, therapies targeting immunomodulation, rather than inflammation suppression, require exploration.

Novel neuroprotective therapies require exploration in IS-HI models. In experimental rodent models of IS-HI, promising therapies include melatonin, N-acetylcysteine, and inflammatory cascade inhibitors/antagonists of IL-1 receptor, nuclear factor kappa beta, plasmogen activator protein-1, and histone deacetylase. Other possible therapies include erythropoietin, mesenchymal stem cells (MSCs), TLR modulators, cyclooxygenase-2 inhibitors, and immunomodulatory antibiotics such as azithromycin and minocycline. It will be important to assess therapies with and without HT; of concern is the negated benefit, or even exacerbation of injury, observed with therapies such as IL-1 receptor antagonists and MSCs when combined with HT.

Unlike the protection seen previously with 24 h HT (from 2 to 26 h) in the piglet HI model, we did not observe protection with 24 h HT (from 1 to 25 h) in a model of IS-HI. Our observation was based on EEG, MRS, and TUNEL-positive cell death. The reduced efficacy of HT, in the context of IS-HI, may relate to the inability of HT to alter glial cell function and pro-inflammatory cytokine profile as well as systemic effects of hyperglycemia and hemodynamic instability. Hypothermic suppression of leukocytes, platelets, and CC3 may be detrimental to systemic inflammatory illness. Novel neuroprotective agents with immunomodulatory properties require exploration in IS-HI models.

ACKNOWLEDGEMENTS
We thank Debbie Kraus (PRISMTC) for her statistical analysis and Cornelius Bauer (UCL) for his expertise with bNIRS acquisition. This project was supported by a grant from the Medical Research Council (grant number MR/M006743/1).

Fig. 6 Cytokines. Mean values over time for IL-6, IL-8, and TNFα in the plasma and cerebrospinal fluid (CSF) (±SEM). Pale blue shading represents cooling. **P < 0.01 for difference at timepoint between the NT and HT group.

Our results, suggesting no protection with HT in IS-HI, may be related to (i) failure of HT to inhibit microglial activation, astrogliosis, and inflammatory cytokine production; (ii) attenuation of neutrophil numbers by HT; (iii) secondary brain injury from hemodynamic instability; and (iv) abnormal glycemic control. Following HI, HT inhibits both protective and damaging cellular pathways; in IS-HI, the balance of inhibition may be detrimental. Given the dual role of the immune response in both contributing to and preventing injury, and the complex mechanisms underlying the evolution of HI injury, therapies targeting immunomodulation, rather than inflammation suppression, require exploration.

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ACKNOWLEDGEMENTS
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AUTHOR CONTRIBUTIONS
All authors met the Pediatric Research authorship requirements. Substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data: all authors. Drafting the article or revising it critically for important intellectual content: K.A.M. drafted the manuscript, with contributions to the methods from C.M., D.P., and M.S. N.J.R. revised the manuscript. Final approval of the version to be published: all authors.

ADDITIONAL INFORMATION
Competing interests: The authors declare no competing interests.

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