Cortical neuronal hypertrophy and mTOR pathway activation in CAN regions in SUDEP

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Abstract

Objectives: Dysfunctional connectivity and preexisting structural abnormalities of central autonomic network (CAN) regions have been shown on magnetic resonance imaging (MRI) in sudden unexpected death in epilepsy (SUDEP) and may be mechanistically relevant. In a previous postmortem study we reported increased microglia in CAN regions, including the superior temporal gyrus (STG) in SUDEP. In this current study we investigated mammalian target of rapamycin (mTOR) pathway activation and neuronal c-Fos activation in CAN regions in SUDEP compared to control groups.

Methods: In a series of 59 postmortem cases (SUDEP, n = 26; epilepsy controls [EPCs], n = 14; and nonepilepsy controls [NECs], n = 19), we quantified pS6-240/4, pS6-235/6 (markers of mTOR activation) and c-Fos neuronal densities and labeling index in the STG, anterior cingulate, insula, frontobasal, and pulvinar regions using immunohistochemistry with whole-slide automated image analysis.

Results: Significantly more pS6-positive neurons were present in the STG in cases with a history of recent seizures prior to death and also in SUDEP compared to other cause of death groups. No differences were noted for c-Fos neuronal labeling in any region between cause of death groups. Cortical neuronal hypertrophy in the STG was observed in some SUDEP cases and associated with pS6-240/4 expression. pS6-235/6 highlighted neuronal intranuclear inclusions, mainly in SUDEP cases and in the STG region.

Significance: Neuronal labeling for pS6 in the STG correlated with both seizure activity in the period prior to death and SUDEP. Further investigations are required to explore the significance of this region in terms of autonomic network dysfunction that may increase the vulnerability for SUDEP.

KEYWORDS
C-Fos, mTOR, SUDEP, superior temporal gyrus
INTRODUCTION

Current theories propose that sudden unexpected death in epilepsy (SUDEP) may be precipitated by a suppression or dysfunction of autonomic systems that regulate respiratory and cardiac functions occurring at around the time of a seizure. Recent attention has focused on the central autonomic networks (CANs), which modulate vital brainstem centers. CANs are interconnected cortical and subcortical networks that enable adaptation to physiological challenges, and knowledge of specific functions associated with brain regions continues to evolve through functional imaging, tractography, as well as stimulation studies.\(^1\)\(^-\)\(^5\)

There is growing evidence from magnetic resonance imaging (MRI) studies in SUDEP for specific patterns of alteration in CAN regions, both representing potential risk biomarkers and of mechanistic relevance to understanding causation. Altered gray matter volumes have been reported in SUDEP, including in temporal, cingulate cortex, and thalamus,\(^6\)\(^-\)\(^9\) and altered connectivity between CAN regions on functional MRI (fMRI), suggesting disorganization of interconnections.\(^6\)\(^,\)\(^10\)\(^-\)\(^12\) Furthermore, electrode stimulation studies in CAN regions in patients with epilepsy have provided important insights on regional cortical autonomic functions of relevance in SUDEP.\(^13\)\(^-\)\(^15\)

In a previous postmortem study of CAN in SUDEP, we noted higher microglial densities in the superior temporal gyrus (STG) and thalamic regions compared to control groups.\(^16\) In addition, we noted increased microglia in patients with reported recent seizures occurring in the 10 days prior to death. Recent seizures, including seizure clusters,\(^17\) have been considered as a risk-factor for SUDEP. In the landmark mortality in epilepsy monitoring unit study reviewing SUDEP on monitoring units, one or more generalized seizure (GS) occurred in the 12h prior to death in 12 of 16 cases.\(^18\) The mechanisms and relationships between recent seizures and SUDEP are unclear but transient neuronal dysfunction, including reversible alterations of modulatory neuropeptides,\(^19\) may represent a period of vulnerability, particularly if involving the critical CAN regions.

Our aim was to investigate c-Fos, an early/immediate proto-oncogene and inducible transcription factor used to monitor temporospatial cellular injury including following epileptiform activity\(^20\) in SUDEP postmortem cases. In parallel we investigated mammalian target of rapamycin (mTOR) pathway activation, which regulates diverse and critical cellular functions and is upregulated following seizures,\(^21\) using phosphorylation of the downstream ribosomal protein S6 as a surrogate marker of pathway activity.\(^22\) This would be investigated in selected CAN regions to explore any relationship to recent seizure histories and other clinicopathological factors.

Key Points
- The superior temporal gyrus showed significantly more mammalian target of rapamycin (mTOR) pathway activation in sudden unexpected death in epilepsy (SUDEP) postmortems compared to other groups.
- c-Fos and mTOR expression in other autonomic regions, including the anterior cingulate, pulvinar, and insular cortex, did not distinguish SUDEP from other causes of death.
- The superior temporal gyrus showed increased microglial activation in a previous study, highlighting the potential relevance of this region to autonomic dysregulation in SUDEP.

METHOD

2.1 | Case groups

SUDEP cases (n = 26), primarily from adults, were obtained through the Epilepsy Society Brain and Tissue Bank (ESBTB) at University College London (UCL). The study has been ethically approved and cases consented for use in research. All of the SUDEP cases were initially coroner’s autopsies conducted for investigations into the cause of death and subsequently donated for research to the tissue bank. These included definite SUDEP cases, with four cases of possible/probable SUDEP according to criteria of Nashef.\(^23\) We included seven SUDEP cases, in which a degree of cortical neuronal hypertrophy had been noted during diagnostic evaluation on routine examination using hematoxylin and eosin (H&E), Cresyl violet/ Luxol fast blue and neurofilament stains. In none of these seven cases was there a clinical, neuroimaging or genetic diagnosis of focal cortical dysplasia during life. Neuronal hypertrophy was evident mainly in the STG region, with scattered enlarged neurons but without the hallmark pathology criteria of cortical dyslamination, balloon cells, and white matter changes for a definitive diagnosis of focal cortical dysplasia.\(^24\) (Further detail in Table S1).

Non-SUDEP epilepsy cases (n = 14) were used as a control groups, from the ESBTB and the ERUK Corsellis epilepsy brain collection, including a subset with documented recent status epilepticus or seizures prior to death (epilepsy controls, EPCs). Nonepilepsy control (NEC) cases were obtained from the ESBTB and Medical Research Council (MRC) Edinburgh brain bank (n = 19). All postmortem cases were collected between 1971 and 2019 (summary group details are shown in Table 1 and further case details.
2.3 **Tissue preparations**

Sections were cut at a 5 μm thickness from the ACC, STG, insular, and pulvinar regions from both the left and right sides, where available. Sections were stained with Cresyl violet for confirmation of the cytoarchitectural regions, which is particularly relevant for ACC to correlate with grey and white matter. We reported microglia in each region, which was assessed by histochemical staining. The microglial density was measured using morphometric analysis.

**TABLE 1**

| Groups                      | Case detail shown in Table S1 | Significance between cause of death groups p value | Age at death, mean (years); range | Gender M:F | Age at onset of epilepsy, mean (years); range | Cases with recent focal brain pathology (relevant to epilepsy) | Brain weight, mean (g); range | Fixation time, mean (days); range | Postmortem interval, mean (days); range |
|-----------------------------|-------------------------------|-----------------------------------------------|----------------------------------|------------|-----------------------------------------------|------------------------------------------------|-----------------------------|----------------------------------|----------------------------------------|
| SUDEP                       | N = 26                        |                                               | 36.38 (15–68)                   | 15:11      | 20.13 (5–49)                                 | N/S (SUDEP and EPC groups)                               | 1354 ± 812 (1060–1580) | 45 (7–120)                  | 3.4 (1–7)                             |
| Epilepsy controls           | N = 14                        |                                               | 48.9 (1–88)                     | 8:6        | 31.2 (0–82)                                 | N/S (all three groups)                                   | 1209 ± 590 (1912–1421) | 11 (2–140)                  | 2.7 (1–6)                             |
| Nonepilepsy controls        | N = 19                        |                                               | 48.9 (2–88)                     | NA         | NA                                           | N/S (all three groups)                                   | 1354 ± 812 (1060–1580) | 29 (12–140)                 | 4.4 (2–12)                            |

Note: Recent = Seizure reported at any time of death (including status epilepticus).

a Fixation times shown for macroscopic brain examination and diagnostic sampling; detailed fixation times for each region shown in Table S1.

**Rationale for the CAN brain**

The regions selected for this study were based on the existing evidence of their involvement in SUDEP. These regions include the subgenual anterior cingulate cortex (ACC), which is involved in cardiovascular/vagal responses and sympathetic regulation. The subcallosal cortex has known roles in cardiovascular/vagal responses and play a role in the regulation of blood pressure. The superior temporal gyrus (STG) is involved in the integration of CAN networks and is a central hub for the regulation of cardiovascular, sympathetic, and parasympathetic responses. The insular cortex is involved in the regulation of cardiovascular, sympathetic, and parasympathetic responses and is a central hub for the regulation of CAN networks. The pulvinar region of the thalamus is recognized as a respiratory and sympathetic regulatory brain region.
distinguish BA25 from BA33 and BA24. Sections from all regions were immunolabeled for c-Fos (Santa Cruz, Biotechnology SC-166940, 1: 50), from all regions (except ACC) for pS6-240/4 (Cell signaling technology, 5364, 1: 500) and pS6-235/6 (Cell signaling technology, 4857S, 1: 50), and in the pulvinar region for glial fibrillary acidic protein (GFAP) (Dako, Z0334. 1: 2000) using a Leica BondMax (see Appendix S1 for details of immunohistochemistry). For the demonstration of pS6, we utilized two antibodies that recognized different phosphorylation sites of pS6: pS6-240/4 is specific for mammalian target of rapamycin complex 1 (mTORC1) pathway activation, whereas pS6-235/236 may also occur through mTOR independent pathways including Ras-mitogen activated protein kinase pathway activation and both antibodies in conjunction have been widely used in studies.22 Because many cases were archival coronial postmortems, not all regions were available from all cases and some regions were sampled at a later interval. The cases used and fixation times for each region are detailed in Tables 2 and S1. Double immunofluorescence labeling was carried out for pS6-240/4 and pS6-235/6 and with neuronal markers (MAP2 [microtubule associated protein] and synapticophysin), GFAP, and microglial marker Iba1 in the cortical and pulvinar regions in selected SUDEP and NEC cases (see Appendix S1 for details).

2.4 | Quantitative analysis and regions of interest

A digital slide scanner (Leica SCN400F, Leica Microsystems or Hamamatsu 360, Hamamatsu Photonics) was used to scan all slides at ×40 magnification. The scanned whole slide images (WSIs) were stored in a computer server and were uploaded to either Definiens (Definiens AG) for c-Fos or QuPath platform31 for pS6 and GFAP image analysis. The following regions of interest (ROIs) were outlined. ACC: Evaluation of the cytoarchitectural of the subcallosal coronal blocks on CV was designated as either BA25 or BA33/24/32 based on previous studies32; in 10 cases both of these BA regions were available in separate blocks. The entire extent and full thickness (pia to white matter border) of these BA regions were outlined, to include both superficial and deep cortical layers but excluding white matter. A further cortical ROI was drawn on frontobasal cortex (BA11), representing a nonautonomic control brain region. STG: ROIs were placed on either side of the gyrus at the mid-point between

| TABLE 2 | Results of quantitative analysis of c-Fos and pS6 labeling in STG, pulvinar, and insular regions |
|----------|---------------------------------|---------------------------------|---------------------------------|---------------------------------|
| Region   | SUDEP                          | Definite SUDEP                  | Epilepsy controls               | Nonepilepsy controls            |
|----------|--------------------------------|--------------------------------|--------------------------------|--------------------------------|
| c-Fos    |                                |                                |                                |                                |
| STG      | 348.43 (260)                    | 279.9 (172.1)                  | 408.15 (285.5)                  | 298.49 (193.3)                  |
| N = 19   | 52.3% (26)                      | 48.9% (25)                     | 48.9% (26)                      | 48.2% (25)                      |
| Insular  | 277.9 (245)                    | 287.18 (249.4)                  | 325.0 (137.8)                  | 174.57 (164.05)                  |
| N = 19   | 37.7% (24)                      | 38.6% (25)                     | 41.5% (10)                     | 26.8% (19.7)                     |
| Pulvinar | 49.05 (61.8)                   | 40.56 (42.7)                    | 24.2 (2.9)                     | 47.49 (54.87)                    |
| N = 19   | 7.6% (7)                        | 6.9% (6.2)                     | 6.5% (1.5)                     | 8.8% (10.5)                     |
| pS6-240/4 | 28 (40)/mm²                   | 18 (32)/mm²                    | 1 (2)/mm²                      | 6 (10)/mm²                      |
| (N = cases⁴) |                   |                                |                                |                                |
| N = 15   | 6.6% (8)                       | 7% (9.2)                       | 6.6% (2.9)                     | 1.2% (0.53)                      |
| Insular  | 5 (14)/mm²                     | 6 (15)/mm²                     | 0.8 (0.5)/mm²                  | 0.7 (0.7)/mm²                    |
| N = 18   | 3.9% (4.7)                     | 4.1% (5.1)                     | 3.5% (2)                      | 2.1% (1.6)                      |
| Pulvinar | 5 (10)/mm²                     | 6 (11)/mm²                     | 0.1 (0.05)/mm²                 | 0.2 (0.1)/mm²                    |
| N = 17   | 6.9% (8.7)                     | 7.2% (9.1)                     | 3% (1.8)                      | 2.6% (3.5)                      |
| pS6-235/6 | 0.26 (0.35)                   | 0.31 (0.37)                    | 0.075 (0.08)                   | 0.071 (0.06)                    |
| (N = cases⁴) |                   |                                |                                |                                |
| N = 19   |                               |                                |                                |                                |
| Insular  | 0.13 (0.15)                    | 0.13 (0.16)                    | 0.086 (0.1)                    | 0.084 (0.1)                     |
| N = 11   |                               |                                |                                |                                |
| Pulvinar | 0.76 (0.88)                    | 0.65 (0.8)                     | 0.67 (0.3)                     | 0.98 (0.59)                     |
| N = 16   |                               |                                |                                |                                |

⁴Refers to the number of cases evaluated in each group for each region (not all regions were available in all cases).
the crown and sulcus, representing the full thickness of cortex but not including white matter. Insular cortex: The entire extent and full cortical thickness representing BA 13/4 were outlined on the sections, overlying the claustrum, with care not to include the white matter or adjacent gyri. Pulvinar: A rectangular ROI of ~45 mm² was placed within the nucleus deep to the subependymal surface in all cases. (Further details of ROIs are detailed in Figure S1A–D.)

For c-Fos sections using Definiens Tissue Studio Software (Definiens AG) thresholds were set following analysis of a pilot series of a range of labeling intensities and densities to delineate single neuronal cells, and the neuronal density (ND) (positive cells/area) was calculated (Figure S1C). Different thresholds were set for the ACC regions from the STG, insular, and pulvinar regions, but the thresholds were kept constant for each region across all cases. Scanned images of pS6-240/4 and pS6-235/6 were uploaded onto the Qupath platform, thresholds similarly optimized and neuronal density/mm² (or ND) and overall labeling index (LI) measured. The LI represents the field fraction of immunolabeling for the whole region and, therefore, provides a measure of the overall labeling of cells and processes. In pulvinar GFAP sections, using similar ROIs, the GFAP LI was evaluated. For c-Fos and pS6-240/4 in the insular, pulvinar, and STG regions, positively labeled neurons as a percentage of all neuronal cells in each region were also calculated (see Appendix S1 and Figure S1E).

Statistical analysis between the cause of death groups was carried out using SPSS (version 25, IBM incorporation) using nonparametric tests (Mann-Whitney, Kruskal-Wallis tests), Spearman’s correlation, and linear regression analysis. Graph pad (Prism, version 9) was used for the graphical representation of the data.

3 | RESULTS

3.1 | Qualitative patterns

3.1.1 | c-Fos

Neuronal labeling (nuclear and cytoplasmic) was present in the cortical regions in epilepsy and control groups but variable between cases. In the ACC (BA33, 24, 32, and 25) common patterns were labeling of layer II neurons alone, a “tram-track: pattern of layer II and layer V/VI neurons, or pan-cortical labeling, the latter pattern particularly noted in BA25 (Figure 1A). In the adjacent control BA11 region, variable laminar neuronal-staining patterns (II, V/VI or pan-cortical) was also observed. In insular cortex and STG, scattered positive neuron, laminar patterns of labeling (layers II, IV, V/VI), or pan-cortical labeling was variably observed but with no pattern dominating in any cause of death group (Figure 1B–D). There was no evidence of glial labeling with c-Fos. In the pulvinar, the impression was of overall fewer c-Fos-positive neurons compared to the cortical regions (Figure 1E).

3.1.2 | pS6-240/4

In the STG region, neuronal cytoplasmic labeling primarily of pyramidal cortical cells was observed both in epilepsy cases and controls, including the hypertrophic neurons described in SUDEP cases (Figure 2A). The extent of labeling in cases varied from scattered neurons to laminar patterns involving layers II, III, V, and VI, with relative sparing of layer IV (Figure 2B) and rare astroglial labeling. pS6-240/4 neuronal labeling was present in the insular region but appeared less extensive than STG (Figure 2C), whereas the pulvinar region showed predominantly infrequent scattered positive neurons with rare astroglial labeling (Figure 2D).

3.1.3 | pS6-235/6

Neuronal cytoplasmic labeling with pS6-235/6 was observed in the STG and to a lesser extent in the insular cortex. Scattered small and large neurons were variably present, randomly scattered through laminae and often in perivascular locations rather than a laminar distribution (Figure 2G). Peri-neuronal granular, synaptic-like labeling was also evident (Figure 2H). In the pulvinar, there was less frequent neuronal labeling and a more prominent synaptic-like labeling (Figure 2I). Cortical astroglial labeling was noted, particularly in layer I astrocytes (Figure 2I). pS6-235/6-positive, linear to round intranuclear neuronal inclusions were noted mainly in the STG region (Figure 2K,L) and present in 57% of SUDEP cases compared to 16% of NECs but not in EPCs.

Double labeling confirmed neuronal co-localization of pS6-240/4 and pS6-235/6 in a proportion of neurons (Figure 2O,P), and co-expression with mainly neuronal markers MAP2 and synaptophysin (Figure 2E,F) over glial markers (Figure 2M,N).

4 | QUANTITATIVE ANALYSIS

4.1 | Cause of death groups

4.1.1 | c-Fos

There was no significant difference between the mean c-Fos neuronal densities for any region (anterior cingulate...
regions [BA25, BA33/24/32], BA11, STG, insular, and pulvinar) between SUDEP, EPCs, and NECs (Tables 2 and S2, Figure 1F). This was also the case when evaluating definite SUDEP cases compared to control groups. Significantly lower c-Fos neuronal densities and also the percentage of c-Fos-positive neurons were confirmed in the pulvinar compared to the STG and insular regions in both SUDEP and NECs, reaching the greatest significance in the SUDEP group \( (p < .001 \) and \( p < .0001 \) (Figure 1F). Considering left or right sides separately between the cause of death groups, the mean c-Fos ND (and percentage positive neurons) was found to be significantly lower in the right STG in SUDEP (and for definite SUDEP) compared to NECs \( (p \leq .05 \) (Figure 1G). Comparing left and right sides within cause of death groups, significantly higher c-Fos ND was noted on the left compared to the right insular cortex in the SUDEP group only \( (p = .02 \).

### 4.1.2 | pS6-240/4 and pS6-235/6

Higher mean neuronal densities were noted with pS6-240/4 in SUDEP compared to other cause of death groups in all regions, with the highest values in the STG region (Table 2, Figure 3A). Significantly higher mean LI for pS6-235/6 was present in the STG in SUDEP cases (and definite SUDEP) compared to NECs \( (p = .004 \) and \( p = .003 \) [definite SUDEP]) (Table 2, Figure 3B). In contrast, in the pulvinar region the mean pS6-235/6 LI was lower in the SUDEP group (and definite SUDEP cases) compared to NECs \( (p = .046 \), Figure 3B). Considering hemispheres

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**Figure 1** c-Fos immunostaining patterns and quantitative analysis in sudden unexpected death in epilepsy (SUDEP). (A) Low-power image of Brodmann area 25 (BA25) and c-Fos labeling showing a pan-cortical to laminar labeling pattern of neurons. (B) Insular cortex (INS) with neuronal labeling in layers II, IV, and V/VI (arrowheads) showing a laminar pattern. (C) Superior temporal gyrus (STG) with neuronal labeling predominantly in layers II and VI (arrowheads). (D) Distinct neuronal labeling, including pyramidal cells with a lack of glial labeling was observed with c-Fos immunohistochemistry. (E) In comparison, the pulvinar region (PULV) showed only scattered positive neuronal cells. (F) Radar plots of c-Fos mean neuronal densities were generated to illustrated relative differences between regions in cause of death groups (SUDEP, D-SUDEP (definite), EP-Cont (Epilepsy controls) and NECs (nonepilepsy controls). Because different thresholds for detection were used in the anterior cingulate regions to other areas, the mean neuronal density is expressed as a ratio of the maximum value for each region. Lower levels of c-Fos were noted in the pulvinar region in all cause of death groups. (G) Scatter plots of c-Fos neuronal densities in the STG region were significantly lower in SUDEP cases on the right side compared to NECs. Bar in (A) = 2 mm, Bar shown in (B) approximately equivalent in (B and C) to 300 μm and in (D and E) to 100 μm.
separately between cause of death groups, higher LI for pS6-235/6 in SUDEP than NECs was noted for both the right and left STG (p < .0001, p < .005). Comparing hemispheres within cause of death groups, a lower percentage of positive pS6-240/4 neurons in the left than the right insula was noted in SUDEP cases (p = .03). There was a significant correlation between pS6-235 and pS6-240 labeling in the STG and insular regions (p < .05 and p = .003, Spearman’s correlation) but not in the pulvinar region. There was a correlation between the percentage of positive neuron labeling with c-Fos and pS6-240/4 in epilepsy groups in the pulvinar region (p < .05) but not in controls or other regions.

4.1.3 GFAP

In view of a lower level of neuronal labeling in the pulvinar compared to cortical regions, quantitative evaluation of GFAP was also carried out. There was no correlation between GFAP labeling index, pS6-235/6, or pS6-240/4 in any cause of death group. No difference in the mean

**FIGURE 2** Phospho-S6 immunohistochemistry patterns in the central autonomic regions. (A–D), pS6 240–44: (A) Scattered hypertrophic neurons were noted, particularly in the superior temporal gyrus region (STG) in a proportion of sudden unexpected death in epilepsy (SUDEP) cases, but lacking the balloon cells of other specific pathological features of a focal cortical dysplasia. These neurons were often pS6 positive (inset). (B) Low power of STG in a SUDEP case showing a laminar pattern of neuronal labeling with pS6-240. (C) In the insular region, frequent neuronal pS6-240 expression was also noted. (D) Scattered but variable numbers of pS6-positive neurons were observed in the pulvinar region (PULV), and in some cases astrogial labeling was noted (inset, arrowed); there was focal colocalization of labeling of pS6-240 with neuronal markers MAP2 (E) and synaptophysin (F). (G–L), pS6 235–6: (G) A prominent perivascular distribution of pS6 labeling was noted in the insular cortex, rather than a laminar pattern. (H) In addition to neuronal perikaryal labeling, a striking perineuronal synaptic granular pattern of labeling was noted, shown here in the STG. (I) Labeling of glial cells in layer I was noted (arrowed) in some cases. (J) In the pulvinar region, a more striking granular labeling of the parenchyma was appreciated. (K and L) Intraneuronal linear inclusions in neurons in the STG were more frequently seen in SUDEP cases. (M and N) Double labeling of pS6 markers with GFAP showed an overall prominent neuronal pattern with minimal localization with glial processes. (O and P) Focal co-labeling of pS6-240 and pS6-235 was observed in neuronal cells. The bar shown in (A) for (A, C, E, F, M, O, P, and H) is equivalent to ~50 microns; in (D, G, I and H) to ~150 microns, and in (B) to ~300 microns.
pulvinar GFAP labeling index between cause of death groups was noted, with lowest mean values in the SUDEP group (Figure S3).

4.2 | Clinical and pathological correlations

Regarding seizure history, there was no significant difference between in regional pS6-235/6 and pS6-240/4 measurements in epilepsy cases with or without a history of status epilepticus prior to death. However, in cases with a history of generalized seizures reported in the 10 days prior to death, significantly higher pS6-235/6 was observed in the STG than cases with no seizures ($p = .004$, Mann-Whitney test, Figure 3C). Higher pS6-240/4 neuronal densities were also noted in all regions in cases with recent seizures, but without reaching significance. There was no significant association of c-Fos neuronal densities in any region in relation to status epilepticus or seizures reported 10 days prior to death.

A significant positive correlation between pulvinar GFAP labeling with age at death was observed in the SUDEP group only ($p = .006$) (Figure S3) but was not noted with other markers. In patients with epilepsy with any focal brain pathologies relevant to seizures in any...
region (including old surgical scars, hippocampal sclerosis [see Table S1 for detail]), a significantly higher LI with pS6-235/6 was noted in the STG (p = .02, Figure 3D) and pS6-240/4 neuronal density in the pulvinar (p = .016, Figure 3E) compared to neuropathology-negative epilepsy cases. In SUDEP cases with neuronal hypertrophy, although higher pS6- pS6-240/4 and 235/6 labeling was noted in the STG and insular cortex, there were no statistically significant difference from epilepsy cases without this feature (Figure S2A,B).

There was no statistical association between c-Fos ND and the presence of an underlying brain pathology or neuronal hypertrophy.

### 4.3 Effects of fixation times and postmortem intervals

There were no significant differences in mean postmortem intervals between the cause of death groups (Table 1) and there were no significant associations between postmortem interval and c-Fos, pS6 240/4, or 235/6 measurements. There was a significant variation in mean brain fixation times between the cause of death groups (Table 1, p < .05), with the shortest fixation times in NECs. There was no correlation between regional tissue fixations times and pS6 labeling in the pulvinar or insular sections. There was an inverse correlation between pS6-240/4 neuronal densities in the STG and fixation time; using multivariate analysis to factor the variable of fixation time, linear regression analysis confirmed significantly higher pS6-240/4 labeling in the STG in SUDEP compared to other cause of death groups (p = .02, Figure 3A). There were no significant associations in c-Fos measures and fixation time for any brain region (Figure S1F).

### 5 DISCUSSION

There is evidence implicating CAN in the peri-ictal autonomic dysfunction that precipitates SUDEP. Using a postmortem series to evaluate four autonomic brain regions, we have shown increased mTOR pathway activation in the STG region in SUDEP that also correlated with recent seizures prior to death. In contrast, there was less evidence that c-Fos neuronal expression patterns in CAN distinguished SUDEP from other causes of death. In a previous study on a different postmortem cohort, increased microglial densities were identified in the STG in SUDEP, which also associated with recent seizures.16 These current observations reinforce that the STG is of potential relevance to the autonomic pathways linked to SUDEP.

Identification of specific tissue biomarkers for acute neuronal dysfunction and their application in the investigation of epilepsy-related and sudden deaths has been an area of recent research, as both a much-needed diagnostic tool as well as furthering our understanding of brain regions that may be critically involved in SUDEP.33,34 mTOR is a major kinase that forms two complexes—mTORC1 and mTORC2—with different activation pathways, regulating many fundamental cellular functions35,36 and influencing neuronal size and synaptic neurotransmission. Activation of this pathway is observed in broad spectrum of focal epilepsy pathologies and nonlesional tissue in both neuronal and glial cell types.37-39 Experimental epilepsy models confirm mTOR activation following seizures,40 occurring within 2 h following status epilepticus in the kainic acid model,41 and mTOR activation may play a critical role in cellular responses and epileptic networks following status epilepticus, although little explored in human tissues to date.42

We utilized quantitative immunohistochemistry for phosphorylation of the downstream ribosomal protein S6 at Ser235/6 and Ser240/4, as mTOR-dependent and mTOR-independent phosphorylation sites, as used widely in previous studies to assess pathway activation.22 Differences in pS6 patterns were noted across the SUDEP cases, which may reflect the clinical and pathological heterogeneity of this group as well as dysfunctional cellular mechanism prior to death. For example, we noted greater pS6 labeling in relation to any underling focal neuropathology identified postmortem. However, we also observed a relationship with reported seizures occurring in the last 10 days before death, which may suggest a more sustained mTOR activation in these cases. In experimental models, mTOR activation evaluated by pS6 immunohistochemistry occurs from 6 h to 5 weeks following seizure onset.21 Furthermore, mTOR activation following experimental seizures induces inflammatory pathways and activates microglia42; of interest, our previous observations of increased regional microglia in SUDEP, particularly with recent seizures prior to death,16 could be linked mechanistically to the observed mTOR activation. In seven SUDEP cases we also noted hypertrophic neurons in the cortical regions, particularly the STG, which did not amount to focal cortical dysplasia based on pathology criteria, although genetic testing had not been conducted. We also observed pS6 expression in these hypertrophic neurons similar to the ubiquitous expression in hypertrophic and dysmorphic neurons reported in focal cortical dysplasias associated with mTOR pathway mutations.22 Of interest, increased rates of SUDEP are reported in some focal epilepsies with DEPDC5 mutations, an mTOR pathway inhibitor,43 and associated with focal dysplasia,44 although the precise cellular mechanisms leading to the sudden
death and any autonomic dysfunction is as yet unclear.\textsuperscript{45} Hypertrophic neurons have long been recognized in acquired epilepsy pathologies,\textsuperscript{46,47} considered to reflect altered neuronal function, metabolism, cellular stress, or connectivity as a result of seizures. The identification of isolated hypertrophic neurons and more widespread mTOR activation in our SUDEP series is unlikely to represent mTOR pathway mutations but raises potential mechanistic links with DEPDC5-related epilepsies as a future line of investigation.

This study highlights the STG as a brain region of relevance in SUDEP (Figure S4). Cortical thinning, including progressive atrophy,\textsuperscript{48} has been demonstrated in the STG in large epilepsy cohorts on MRI.\textsuperscript{26} There are few data on specific autonomic regulatory functions associated with the STG, but parasympathetic autonomic regulation is recognized\textsuperscript{1} to be of relevance to SUDEP. In addition, tractography studies of the CAN place the temporal lobe, particularly the STG as an integral part of autonomic networks including the parieto-anterior-temporal pathway, which connects STG to amygdala and insular cortex.\textsuperscript{4} In this current study only a minority (eight of the SUDEP cases) were also included in our previous study reporting increased microglia in the STG in SUDEP; this reinforces that pathological alteration in this region may be relevant to SUDEP across cohorts. Concerning lateralization, there was some evidence in this series for greater mTOR activation on the right than left STG and lower c-Fos activation in the left STG in SUDEP compared to controls. In a recent study of patients with severe ictal-hypoxia during generalized seizures, considered a risk-factor for SUDEP, lower gray matter volumes were observed on the right but not on the left STG.\textsuperscript{7} Although not all SUDEP cases showed increased pS6 labeling, likely reflecting the heterogeneity of underlying causes, these collective studies suggest that further study of specific autonomic functional networks associated with the right STG would be relevant for some cases.

The subcallosal (or subgenual) cingulate gyrus comprises BA25 as well as parts of BA33, 24, and 32 as cytoarchitectonically distinct regions\textsuperscript{32} with different functional connectivity profiles; BA25 is activated with autonomic processing and BA33/24 is co-activated with the amygdala and anterior cingulate, among other regions.\textsuperscript{49} In tractography studies, the ACC forms part of the rostral limbic pathway, which is connected to the insular region.\textsuperscript{4} BA25 has been regarded as the principal site of autonomic regulation in the frontal lobe,\textsuperscript{50} including cardio-vagal\textsuperscript{1,3} as well as sympathetic regulation.\textsuperscript{5} Stimulation of Brodmann area 25 (subcallosal neocortex) in patients with epilepsy produced striking systolic hypertensive changes.\textsuperscript{14} Furthermore, reduced functional connectivity of the ACC in fMRI has been associated with SUDEP risk\textsuperscript{11} and network alterations in SUDEP.\textsuperscript{10} In structural MRI studies, increased thickness of subgenual cingulate cortex was shown in patients with GS\textsuperscript{8} and increased gray matter volumes reported in the subcallosal cortex in SUDEP vs Control groups.\textsuperscript{25} This is the first pathological study of BA25 in epilepsy but despite striking laminar neuronal labeling with c-Fos in SUDEP cases, we failed to identify distinct difference from controls, supported by quantitative analysis.

The insula is also regarded as a central hub in connectome studies of CAN using tractography.\textsuperscript{4} Cortical thickening in the insular region has been reported in patients with generalized seizures on MRI,\textsuperscript{8} increased functional connectivity recognized in SUDEP,\textsuperscript{11} and surgical resection associated with autonomic dysfunction.\textsuperscript{27} There is robust evidence for a role of the insula in the regulation of heart rate,\textsuperscript{51} with inconsistent evidence for lateralization\textsuperscript{52}; PET hypometabolism in the right posterior insula was observed in patients with epilepsy with ictal asystole\textsuperscript{29} but electrode implantation and stimulation of either left or right insula was shown to induce changes in heart rate and cardiac output but without effects on blood pressure or respiration.\textsuperscript{5} We noted some lateralization of pathology with greater c-Fos labeling and reduced pS6-240 labeling in the left compared to the right insula in the SUDEP group, but no overall significant differences from controls. The pulvinar region showed less mTOR and c-Fos expression than cortical regions but again no specific patterns emerged in SUDEP cases. In addition, despite evidence for reduced posterior thalamic volumes in SUDEP in structural MRI,\textsuperscript{9,25} we also did not identify increased gliosis in the pulvinar region in SUDEP, only a significant association with age, and the pathological basis for these imaging alterations remains unclear.

There are several limitations to this postmortem study, in that not all regions were available in all cases and only limited CAN regions were selected for this study. We included cases in the study with reports of a seizure occurring in the last days prior to death from the clinical records; we cannot exclude that in cases without this history, a recent seizure occurred but was not documented. Furthermore, in many cases, details of seizure types (including for status epilepticus), duration, and localization through recent EEG and MRI investigations were not always available to further stratify the cases. In addition, different fixation times for brain regions were factored into our analysis. This is a coroner’s postmortem SUDEP cohort, and genetic testing is not available to correlate with pathology findings.

In summary, neuronal labeling for pS6 in the STG correlated with seizure activity in the period prior to death and was enhanced in SUDEP, highlighting this as a potential critical region. Further clinical and functional studies,
including the role of this region within the CAN, is therefore warranted in SUDEP.

**AUTHOR CONTRIBUTION**

SP and HEH were involved in tissue sampling protocols and preparation. SP, FC, and IS carried out immunostaining and quantitation. YL developed image analysis protocols. SP and MT were involved in data analysis, and all the authors were involved in manuscript preparation.

**ACKNOWLEDGMENTS**

University College London (UCL) is part of the Center for SUDEP Research (CSR) supported through a National Institute of Neurological Disorders and Stroke of the National Institutes of Health (NIH; Award Numbers neuropathology of SUDEP: 5U01NS090415 and SUDEP admin core grant: U01-NS090405). All tissues are provided by the Epilepsy Society through the Katy Baggott Foundation. Supports the UCL core grant: U01-NS090405.

**CONFLICT OF INTEREST**

None of the authors has any conflict of interest to disclose.

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