Specific pattern of maturation and differentiation in the formation of cortical tubers in tuberous sclerosis complex (TSC): evidence from layer-specific marker expression

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Abstract

**Background:** Tuberous sclerosis complex (TSC) is a multisystem disorder that results from mutations in the TSC1 or TSC2 genes, leading to constitutive activation of the mammalian target of rapamycin (mTOR) signaling pathway. Cortical tubers represent typical lesions of the central nervous system (CNS) in TSC. The pattern of cortical layering disruption observed in brain tissue of TSC patients is not yet fully understood, and little is known about the origin and phenotype of individual abnormal cell types recognized in tubers.

**Methods:** In the present study, we aimed to characterize dysmorphic neurons (DNs) and giant cells (GCs) of cortical tubers using neocortical layer-specific markers (NeuN, SMI32, Tbr1, Satb2, Cux2, ER81, and RORβ) and to compare the features with the histo-morphologically similar focal cortical dysplasia (FCD) type IIb. We studied a cohort of nine surgically resected cortical tubers, five FCD type IIb, and four control samples using immunohistochemistry and in situ hybridization.

**Results:** Cortical tuber displayed a prominent cell loss in all cortical layers. Moreover, we observed altered proportions of layer-specific markers within the dysplastic region. DNs, in both tubers and FCD type IIb, were found positive for different cortical layer markers, regardless of their laminar location, and their immunophenotype resembles that of cortical projection neurons.

**Conclusions:** These findings demonstrate that, similar to FCD type IIb, cortical layering is markedly disturbed in cortical tubers of TSC patients. Distribution of these disturbances is comparable in all tubers and suggests a dysmaturation affecting early and late migratory patterns, with a more severe impairment of the late stage of maturation.

**Keywords:** Tuberous sclerosis complex, Cortical layer markers, Epilepsy, Neurosurgery, Neuropathology

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Background
Tuberous sclerosis complex (TSC) is a genetic disease caused by mutations in the TSC1 and TSC2 genes. The two proteins encoded by these genes—tuberin and hamartin—are key regulators of the mammalian target of rapamycin complex 1 (mTORC1) [1]. Therefore, the disease affects multiple organ systems already during development. Brain lesions can be found in about 90 % of patients with TSC and are often associated with intractable epilepsy [2–4]. Only about 40–60 % of all patients with TSC and epilepsy become seizure-free on medication. In a selected subset of TSC patients, epilepsy surgery is considered as a therapeutic option after careful presurgical evaluation [5].

Histology of resected cortical tubers reveals a severely distorted cortex with apparent loss of cortical layers and presence of abnormal cell types (reviewed in [3, 6]). The most common aberrant cell types are dysmorphic neurons (DNs) and bright eosinophilic giant cells (GCs); these are very large cells lying scattered within the cortex and condensed in the white matter. The increase in cellular mass can be explained by the over-activation of anabolic metabolism [7]. Accordingly, the underlying genetic defect results in a constitutive activation of mTORC1 with consequent increase in protein synthesis caused by the production of additional ribosomes and increased rates of messenger RNA translation [8]. The increased knowledge concerning the molecular basis of GC formation has led to novel therapeutic strategies in patients with TSC [9].

However, the origin and phenotype of abnormal cells observed in cortical tubers remain still unclear. It was speculated that in focal cortical dysplasia type IIb (FCD type IIb; [10])—a pathology similar to cortical tubers—these cells arise from radial glial progenitor cells of the ventricular zone representing different stages of maturation [11].

Recently, the recognition of layer-specific neuronal cell markers of the neocortex allowed detailed studies about cortical layer formation. These studies provided insight into the origin and phenotype of cortical neurons in normal cortex but may also contribute to define the pattern of cortical layer disruption in different malformations of cortical development (MCD) in the human brain. These markers specifically labeled subsets of cortical neurons with limited laminar distribution [12]. Furthermore, there is evidence that the identity of each neuron within the neocortex (type of neuron or laminar fate) is predetermined even before the onset of migration [13]. Several studies have been published studying cortical layer markers in FCD. Previous results suggested that FCD samples contain more immature cells than age-matched controls, indicating an activation of progenitor cells that might contribute to the pathophysiology of these lesions [14]. In FCD type IIb, DNs and balloon cells (BC; similar to GCs in cortical tuber) showed distinct expression of layer markers supporting an origin from radial glia and intermediate progenitor cells [14]. Differences in neuronal maturation between FCD subtypes, as well as the existent laminar structure in normal-appearing neurons, led to the conclusion that only a subpopulation of precursor cells is affected [15]. However, another study suggested that these differences might reflect a differential expression of cortical layer markers in various brain regions [16].

In the present study, we aimed to investigate the expression of a panel of layer-specific markers covering all cortical layers (Satb2, Cux2, RORβ, ER81, Tbr1, SMI32, and NeuN) in cortical tubers of patients with TSC compared to perilesional cortex of the same patients, as well as to FCD type IIb or aged- and location-matched controls.

Methods
Subjects
In total, we collected nine cortical tubers and five FCD type IIb samples from various epilepsy surgery centers (Department of Pediatric Neurology, Brain Center Rudolf Magnus, University Medical Center Utrecht; Department of Pediatric Neurology, Charles University, 2nd Faculty of Medicine, Motol University Hospital, Prague; Evangelisches Krankenhaus Mara Bethel-Bielefeld, Bielefeld, Germany; Department of Neuropathology, University Hospital Erlangen, Erlangen, Germany) for this study. Extensive presurgical evaluation including 24 h to 5 days video-EEG monitoring, high-resolution MRI, and neuropsychological testing was performed in each patient in order to identify the epileptogenic zone (EZ) before tailored surgical resection. Perilesional cortex (if available in both pathologies) was also included as defined by the lack of dysmorphic neurons and giant/balloon cells in the cortex.

We included an age-matched autopsy control group (n = 4). None of these patients had a history of seizures or other neurological diseases.

Tissue was obtained and used in accordance with the Declaration of Helsinki and the AMC Research Code provided by the Medical Ethics Committee and approved by the science committee of the UMC Utrecht Biobank. The local ethical committees of all participating centers gave permission to undertake the study.

Tissue preparation and immunohistochemistry
The tissue was carefully oriented, cut perpendicular to the pial surface, fixed overnight in 4 % formaldehyde, and routinely processed into liquid paraffin. Sections were cut at 4–6 μm with a microtome (Microm, Heidelberg, Germany) and mounted on positively charged slides (Superfrost + Menzel, Germany). Each specimen
was histopathologically examined using hematoxylin and eosin (H & E). In autopsy controls, an additional Nissl staining was performed to calculate overall neuronal densities.

An immunohistochemical examination of all surgical specimens was performed using the following panel of antibodies: NeuN (neuronal nuclei, 1:100, clone A60, Chemicon, Temecula, CA, USA), non-phosphorylated neurofilament H (1:1000, clone SMI32, Sternberger, Lutherville), Satb2 (1:400, clone SATBA4B10, Abcam, Cambridge, UK), and Tbr1 (1:50, polyclonal rabbit, Abcam, Cambridge, UK).

The slides were air-dried overnight at 37 °C. All immunohistochemical stainings were performed with a Ventana semi-automated staining machine (Benchmark ULTRA; Ventana, Illkirch, France) and the Ventana DAB staining system according to the manufacturer’s protocol.

**In situ hybridization**

In situ hybridization (ISH) for Cux2, RORβ, and ER81 were performed using a 5′–3′ double digoxygenin (DIG)-labeled superior probes (containing locked nucleic acid [LNA]) and 2 O-Methyl [2′OMe] RNA moieties; “l” indicates LNA and “m” indicates 2OMe RNA; Ribotask ApS, Odense, Denmark):

- Cux2: 5′DIG-lAmAmUItMUmCITmCmU1GmCmAl GmCmAlAmGmGmUmUIUT-DIG
- RORβ: 5′DIG-lAmAmUItMUmCICmUmUIGlGmGlU1 TmCmU1AmU1AmGlGIC-DIG
- ER81: 5′DIG-lAmAmUItMUmCITmCmA1TmAmGl TmAmAI1tAmGmCmGlGIG-DIG

The hybridizations were done on 5-μm sections of paraffin-embedded materials as described previously [17, 18]. The probes were hybridized at 62 °C for 1 h, and the hybridization was detected with alkaline phosphatase (AP)-labeled anti-DIG (Roche Applied Science, Basel, Switzerland), NBT (nitro-blue tetrozolium chloride)/BCIP (5-bromo-4-chloro-3′-indolylphosphate p-toluidine salt) was used as chromogenic substrate for AP. Negative control assays were performed without probes and without primary antibody (sections were blank).

**Quantitative analysis of the immunopositive cells and the ISH signals**

To study the alteration of cortical layering, the cortex was divided into three regions: supragranular layers (L1-3, SG), layer 4 plus infragranular layers (L4-6, IG), and deep white matter (>500 μm WM). The number of immunopositive cells was counted at ×10 magnification in five representative fields of L1-3 and L4-6 as well as 10 visual fields of WM. If applicable, the tuber samples were divided into perilesional cortex (PLx) and full-blown tuber (Tb). Perilesional was considered if the cortex showed a normal cortical architecture and no signs of DNs or GCs. A variable degree of giant cells in the white matter was always present.

**Statistical analysis**

Statistical analysis was performed on SPSS 21 (IBM, PASW Statistics, USA). The total number of positive cells as well as the ratio of positive cells divided by the total amount of neurons was used for computing the analysis. Descriptive statistics (median, range) and non-parametric testing were performed to evaluate between-group differences. \( p < 0.05 \) was considered significant.

**Results**

**Demographic patient data**

Our cohort of epilepsy surgery patients consisted of six female and eight male patients. The control group consisted of two females and two males. Cortical tubers were resected temporally in five cases and frontally in four cases. The somatic mutation in the TSC gene was known in three of the patients (TSC1 = 1; TSC2 = 2). The site of FCD type IIb was frontal in four cases and occipital in one. The control group comprised two frontal, one temporal, and one occipital sample. The median age at surgery was 5.00 years (range = 3.00–36.00) in patients with TSC, 12.00 years (range = 1.00–36.00) in patients with FCD type IIb, and 8.50 years (range = 07.00–12.00) in controls.

The first step of the analysis included a general assessment of cortical layering in our different cohorts.

**Cortical layer disruption in TSC cortical tubers and FCD type IIb**

In order to determine the extent of dyslamination in TSC cortical tubers, we first addressed the total number of neurons in cortical layers as well as in the white matter from cortical tubers and FCD type IIb (Fig. 1a–i). In our cohort, we detected a significant neuronal cell loss in cortical tubers and FCD type IIb when compared to perilesional tissue and non-epileptic autopsy specimens (Kruskal-Wallis, \( p = 0.000 \), Table 1, Fig. 1j). In addition, there was a significant increase in heterotopically placed neurons in deep white matter in both pathologies compared to our control samples (Kruskal-Wallis, \( p = 0.033 \), Table 1, Fig. 1k, Additional file 1: Figure S1). As previously reported [19], DNs are characterized by a pathologic accumulation of non-phosphorylated neurofilament (SMI32) in both FCD type IIb and cortical tubers, and we observed significantly more DNs present in the neocortex of FCD type IIb than cortical tubers (Kruskal-Wallis, \( p = 0.017 \), Table 1, Fig. 1l, Additional file 1: Figure S1).
Next, we looked at neuronal fate determination in the remaining neuronal populations utilizing layer-specific markers.

**Satb2 and Cux2 expression**

Neurons in the upper cortical layers were defined by their immunoreactivity for Satb2+ (known to be expressed in L2, L3, L4, and L5; Fig. 2a–h) and Cux2 (normally expressed in L2 und L3, Fig. 2i–p) [12, 14–16, 20, 21]. Overall, the amount of Satb2+ neurons was decreased in perilesional cortex, cortical tubers, and FCD type IIb compared to non-epileptic autopsy controls (Kruskal-Wallis, \( p = 0.003 \), Table 1, Fig. 2q, Additional file 1: Figure S1). The proportions of neurons that were Satb2+ and subsequently corrected for the total amount of neurons showed a trend towards loss of Satb2+ neurons in lower cortical layers in TSC cortical tubers and FCD type IIb. A larger amount of dysmorphic neurons can be detected in FCD type IIb compared to TSC cortical tubers. SG supragranular layers, IG infragranular layers, WM white matter, Co control, PLx perilesional cortex, Tb cortical tuber, IIb FCD type IIb.

**Fig. 1** Cortical disorganization in TSC cortical tubers. a Normal control cortex with classic 6-layered lamination. b, c Normally shaped neuronal cells of supra- and infragranular layers. d Deep white matter in a control patient. e Normal appearing cortex in a perilesional sample. f Cortical dyslamination of FCD type IIb in a patient with difficult-to-treat frontal lobe epilepsy. g Dysorganization of the cortex in a cortical tuber of a patient with genetically proven TSC. h i Dysmorphic neurons (arrows) are present throughout the whole cortex. Scale bar in i equals 500 \( \mu \text{m} \) and applies also to a, e, and f. Scale bar in l equals 200 \( \mu \text{m} \) and applies also to b–d. j Loss of neurons throughout all cortical layers in TSC cortical tubers and FCD type IIb. k Significant increase in heterotopic neurons in the deep white matter in patients who underwent surgery for FCD Type IIb or cortical tuber resection. l A larger amount of dysmorphic neurons can be detected in FCD type IIb compared to TSC cortical tubers. SG supragranular layers, IG infragranular layers, WM white matter.
| Pathology | Location | Total neurons [median] | Dysmorphic neurons [median] | Total Tbr1+ cells [median (range)] | Total Satb2+ cells [median (range)] | Total Cux2+ cells [median (range)] | Total RORβ+ cells [median (range)] | Total ER81+ cells [median (range)] | Tbr1+ cells per total neurons [%] | Satb2+ cells per total neurons [%] | Cux2+ cells per total neurons [%] | RORβ+ cells per total neurons [%] | ER81+ cells per total neurons [%] |
|-----------|----------|-----------------------|----------------------------|-----------------------------------|-----------------------------------|-----------------------------------|-----------------------------------|-----------------------------------|---------------------------------|-------------------------------|-----------------------------|-----------------------------|-----------------------------|
| Tuber SG  | 399.50   | 110.00–238.00         | 41.00 (0.00–126.00)       | 63.00 (49.00–161.00)             | 43.00 (0.00–145.00)             | 88.00 (5.00–198)                  | 161.00 (99.00–706.00)            | 8.00                             | 19.00                          | 13.00                         | 23.00                       | 48.00                       |
| IG        | 348.00   | 111.50 (4.00–204.00)  | 94.00 (0.00–178.00)       | 2.00 (0.00–136.00)               | 13.00 (0.00–112.00)             | 134.00 (3.00–165.00)             | 206 (96.00–565.00)              | 30.00                            | 1.00                           | 4.00                          | 27.00                       | 49.00                       |
| FCD IIb SG| 476.00   | 217.00 (175.00–275.00) | 44.00 (13.00–103.00)     | 181.00                           | 88.00 (0.00–173.00)             | 156.00 (15.00–200.00)            | 373.00 (175.00–799.00)          | 9.00                             | 23.00                          | 20.00                         | 25.00                       | 85.00                       |
| IG        | 523.00   | 192.00 (113.00–206.00) | 61.00 (58.00–287.00)     | 204.00                           | 76.00 (44.00–123.00)            | 166.00 (98.00–187.00)            | 231.00 (182.00–314.00)          | 12.00                            | 32.00                          | 12.00                         | 26.00                       | 49.00                       |
| PLx SG    | 916.00   | 176.00 (0.00–460.00)   | 193.00 (136.00–315.00)   | 345.00 (78.00–660.00)            | 121.00 (0.00–264.00)            | 227.00 (12.00–652.00)            | 16.00                             | 19.00                            | 37.00                          | 10.00                         | 27.00                       |
| IG        | 1000.00  | 420.00 (246.00–662.00) | 14.00 (9.00–215.00)      | 99.00 (12.00–276.00)             | 452.00 (120.00–688.00)          | 519.5 (169.00–987.00)            | 44.00                             | 3.00                             | 16.00                          | 55.00                         | 64.00                       |
| Control SG| 952.00   | 14.50 (5.00–24.00)     | 548.00 (420.00–676.00)   | 367.00                           | 49.00 (35.00–63.00)             | 65.00 (58.00–73.00)              | 0.00                              | 36.00                            | 39.00                          | 12.00                         | 19.00                       |
| IG        | 971.00   | 321.50 (300.00–343.00) | 407.50 (362.00–453.00)   | 55.00                            | 465.00 (412.00–501.00)          | 450.00 (407.00–493.00)           | 23.00                             | 30.00                            | 6.00                           | 50.00                         | 59.00                       |

Overview over the quantitative analysis. Our findings indicate a tendency towards a loss of Satb2+ neurons in layer 5 and an increase of Tbr1+ cells throughout all cortical layers.

SG supragranular, IG infragranular, FCD focal cortical dysplasia, PLx perilesional cortex.
Fig. 2 (See legend on next page.)

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cortex when showing strong expression in L2 and L3 (Fig. 2i–l) [21–23]. In contrast to normal Cux2 distribution, only in upper layers L2 and L3 in controls, cortical tubers showed less Cux2+ cells in the upper layers (Kruskal–Wallis, p = 0.023). Corrected for neuronal number, there was however no difference. DNs were strongly Cux2+ throughout all layers of cortical tubers. Nonetheless, in four out of nine cases, all normally appearing neurons also expressed Cux2 (Fig. 2m–p). Expression was never detected in giant cells. In FCD type IIb, we observed a similar pattern, although the expression pattern was limited to the upper cortical layers in one case.

**RORβ expression**

Neurons of the internal granular layer (L4) can be assessed by ascertaining the expression pattern of RORβ. In our control samples and perilesional cortex, expression of RORβ was restricted to L4 with some scattered neurons in L6 (Fig. 3a–d). We observed reduced expression of RORβ in the lower layers of cortical tubers and FCD type IIb cases (Kruskal–Wallis, p = 0.024). However, strong expression was found in DNs of all cortical layers whereas only in two cases of TSC and in one case of FCD type IIb GCs/BCs were slightly stained on ISH (Table 2). Some of the remaining normal-appearing neurons showed also expression in L4–L6 (Fig. 3e–h).

**Tbr1 and ER81 expression**

The lower cortical layers (L5/6) can be determined via the expression of Tbr1 and ER81. In general, we observed a decrease in the total numbers of Tbr1+ neurons in cortical tubers and FCD type IIb samples (Fig. 4a–d).

| Table 2 Expression pattern of cortical layer markers in aberrant cell forms |
|-----------------|-----------------|-----------------|
|                  | TSC cortical tubers | FCD type IIb |
|                  | DN    | GC    | DN    | BC    |
| Satb2            | 20 %  | 80 %  | 15 %  | 80 %  |
| Cux2             | 80 %  | 0 %   | 80 %  | 0 %   |
| RORβ             | 80 %  | 10 %  | 80 %  | 5 %   |
| Tbr1             | 0 %   | 0 %   | 0 %   | 0 %   |
| ER81             | 80 %  | 15 %  | 80 %  | 0 %   |

Only minor differences between TSC cortical tubers and FCD type IIb cells

Discussing cortical layer formation in malformations of cortical development has recently become possible by the use of neocortical layer-specific markers [12]. The present study provides insights in the pattern of cortical layering in TSC cortical tubers. In summary, we observed
an overall loss of neurons in all cortical layers. We observed further a proportional loss of Satb2+ neurons in lower cortical layers. In contrast, Cux2 was expressed in all cortical layers of tubers and FCD type IIb. In addition, we detected an increase of Tbr1+ and ER81 expressing cells in upper cortical layers. The distinct patterns of disruption of upper and lower cortical layers indicate that dysmaturation of the cortex is a key feature also in TSC brain. However, our data suggests that the upper cortical layers are more severely hampered due to the remnants of neurons that were supposed to relocate in deeper cortical layers implying a more serious impairment of their late stage of maturation.

Satb2, Cux2, RORβ Tbr1, and ER81 expression patterns in TSC: loss of projection neurons in cortical tubers

Satb2 is an AT-rich DNA-binding protein that has been shown to represent a postmitotic determinant for upper-layer neuron specification in the neocortex [28, 29]. Previous studies reported that Satb2+ cells are present in the layers 2 and 3. Additionally, it was suggested that they disappear after 3 months of age in the human brain [20, 29]. On the contrary, it was also reported that Satb2+ cells remain as a subpopulation in the upper cortical layers of the human brain but also extending towards layer 5 (having a predisposition for the frontal lobe) [30, 31]. In our study, expression of Satb2 was detected in L2, 3, and 5 in both perilesional samples and postmortem controls. In TSC cases (tuber and peritubetal cortex), we observed a lower number of Satb2+ neurons compared to controls and there was a tendency towards a loss of Satb2+ neurons in L5 compared to both controls and FCD type IIb. These findings are therefore suggestive for a specific loss of cells in a subpopulation of cortical neurons in TSC indicating an abnormal formation of pyramidal cells. However, abnormal Satb2 expression in the superficial region of the neocortex has been previously reported in FCD type II, showing a different pattern compared to hemimegalencephaly, which is characterized by a diffused Satb2 cortical expression pattern [20]. Altered Satb2 expression patterns by random migration were also reported in various lissencephalies [29]. Another study focusing on FCD showed Satb2+ neurons diffusely distributed in the cortex and/or the white matter of FCD type II cases, without differences in Satb2+ cell densities between FCD type IIa and FCD type IIb [15]. In addition, we observed Satb2 expression in both DNs and GCs/BCs. However, previous studies evaluating Satb2 expression in FCD case do not specify the cell subtypes (DNs, BCs, normal-appearing neurons); thus, comparison cannot be performed.

Cux proteins, putative markers of upper cortical layer neurons [22], have been shown to be critically involved in the regulation of normal dendritic development of L2-3 neurons [23]. In agreement with previous studies
in human neocortex [21], in both control and TSC-perituberal cortex, Cux2 mRNA was expressed in L2-3. However, in cortical tubers, this pattern was disrupted and the expression of Cux2 was extensively heterogeneous with variable expression in normal-looking neurons and DNs (also in FCD IIb), but not in GCs in our cohort. It seems therefore that the fate determination of these neurons is altered and they stay diffusely behind throughout the whole cortex without finding its proper way towards the upper layers. In addition, expression of Cux2 mRNA in both abnormal and DNs has been also reported in a larger cohort of FCD cases, without...
revealing any differences between type Ila and IIb [21]. In contrast, Cux1 protein expression has not been detected in DNs [14].

The RORβ gene is known as a selective marker for the internal granular layer (L4) of mammalian neocortex [12, 32, 33]. According to previous studies [16, 21, 33], also in our cohort of controls and perilesional TSC cortex, expression of RORβ mRNA was intensely expressed in L4. In cortical tubers remaining normal-appearing neurons maintained the expression RORβ in L4. These results are partly in line with the reported pattern in FCD II cases [16, 21]. In our cohort, similar pattern was observed in tubers with different cortical localization, whereas differences between frontal and temporal resections have been reported in FCD cohorts [16]. RORβ expression was also detected in DNs, distributed throughout all layers within the tuber. Expression of RORβ mRNA in DNs has been also reported in FCD type II [21].

Tbr1 and ER81 are well-known neocortical layerspecific markers used to identify the lower cortical layers (L5/6) [16, 34, 35]. Tbr1 gene encodes a transcription factor necessary for the differentiation of neural stem cells in developing brain and highly expressed in glutamatergic early-born cortical neurons [12, 34, 36, 37]. According to these previous studies, Tbr1 was prominently expressed in the lower cortical layers. In both TSC and FCD cases, we observed a lower number of Tbr1+ neurons compared to controls and perilesional cortex; however, if corrected for the total number of neurons, there was a tendency towards an increased number of neurons in the upper cortical layers and in the white matter of epilepsy surgery specimens. Abnormal distribution of Tbr1+ cells throughout the cortex has been reported in both FCD and hemimegalencephaly [14, 20], and localization of Tbr1+ cells in the superficial cortical region has been observed in FCD type II cases [20]. In our cohort, we also observed an increase in Tbr1+ neurons in the perilesional cortex. Interestingly, increasing evidence indicates the extension of structural and molecular abnormalities in the perilesional cortex which is an integral part of the epileptic network in TSC brain [38–41].

The transcription factor ER81 represents another marker of the projection neurons of L5-6 [12, 42, 43]. ER81 mRNA expression pattern was preserved in the perilesional cortex. In cortical tubers, the constitutive expression pattern in normal-appearing neurons was relatively preserved; however, strong ER81 expression was detected in DNs throughout the dysplastic cortex. Expression of Er81 in both DNs and normal-appearing neurons has been also observed in FCD type II cases [21, 35]. In agreement with previous reports [16, 35], ER81 expression was also observed in normal-appearing neurons in the white matter of all surgical epilepsy specimens and in the upper cortical layer of FCD cases [35]. Nonetheless, these findings should be carefully interpreted in the light of an overall reduction of neurons in cortical tubers and FCD type IIb [44].

How can the altered layer-specific marker expression be interpreted in the context of the pathogenesis of the lesions? The data in TSC and FCD type II suggest an
aberrant migration with a greater impairment in late stage cortical development as a common etiopathogenic mechanism underlying the pattern of cortical layer disruption observed in both lesions. The relative layer preservation reported in other types of FCD (such as FCD IIIa; [16]) indicates more a lesion-induced cause rather than simply the effect of seizure activity. The constitutive activation of mTOR (observed in both TSC and FCD type II) as a consequence of the underlying genetic defects may represent the common molecular link resulting in abnormal cell morphology, migration, and differentiation during corticogenesis and contributing to the aberrant protein expression observed in these lesions.

Cortical layer disruption in TSC cortical tubers
Evaluation of the neuronal densities (with a pan-neuronal marker) in our cohort of TSC patients revealed significant neuronal cell loss in cortical tubers compared to both perilesional tissue and non-epileptic autopsy controls. A similar reduction of neuronal densities is also detected in FCD type IIb cases. Our findings are in line with previous stereological studies performed in FCD cases (including both pediatric and adult cases). These already showed lower cortical neuronal densities in the region of dysplasia (particularly in FCD type IIb) compared to the adjacent cortex or non-epilepsy controls, regardless of age and cortical localization [35, 44]. Whether reduced neuronal densities may simply reflect

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**Fig. 6** Schematic overview of layer marker expression. 

- **A** Cortical layer markers are expressed in normal cortex as follows: Cux2: layers II and III (blue); Satb2: layers II, III, IV, and V (green); RORβ in layer IV and to a lesser extent layer VI (orange); ER81 in layers V and VI (purple), and Tbr1 in layers V and VI and few white matter neurons (red). 
- **B** Expression pattern of cortical layer markers in tubers and FCD type IIb: dysmorphic neurons strongly expressed Cux2, RORβ, and ER81 and to a lesser extent Satb2 throughout all layers of the cortex. They did not express Tbr1. Giant cells of cortical tubers often expressed Satb2, ER81, or none of the markers. Balloon cells of FCD type IIb showed a similar pattern although to a lesser extent. 
- **C** Expression of Tbr1 and ER81 was increased in the upper cortical layers and was also present in normal-appearing neurons of the white matter. Triangles neurons, circles giant/balloon cells, SG supragranular layers, IG infragranular layers.
cortical volume, expansion is still a matter of discussion since archival paraffin blocks are not suitable for accurate evaluation of the total volume of the dysplastic region [35, 45]. However, in TSC patients (but not in FCD patients), lower gray matter NeuN densities have been shown to be correlated with reduced MRI gray and white matter volumes, suggesting a reduced neuronal production during corticogenesis (failure of progenitor cell proliferation) [46]. However, we cannot exclude selective vulnerability of neurons in which the mTOR signaling is activated. Accordingly, hyperactivation of this pathway has been shown to contribute to cell damage, promoting aging-related changes (reviewed in [47]. Our recent observations suggest that the abnormal activation of mTOR may contribute to apoptosis signaling pathways and premature activation of mechanisms of neurodegeneration in both FCD type II and TSC [48]. Interestingly, although DNs were detected in all cortical layers in both FCD type IIb and TSC cortical tubers, their number was lower in cortical tubers compared to FCD.

We also observed a significant increase in heterotopic neurons in the deep WM in both FCD and TSC specimens compared to controls. This, however, represents a common finding in epilepsy cases, including FCD [46, 49], as well as mild malformation of cortical development (mMCD) [50]. Moreover, the mechanisms underlying the presence of heterotopic WM neurons (i.e., local migratory failure or to incomplete regression of subplate neurons), their relationship with seizure activity, and their role in epileptogenesis remain still unclear [50, 51].

Conclusions

In summary, our findings provide insight in the abnormal pattern of lamination observed in cortical tubers (summarized in Fig. 6). We provide evidence for an excess of early stage migrational neurons (ER81 and Tbr1) shifting towards the upper layers and therefore hampering late stage migration albeit only the subpopulation of CUX2 expressing cells being affected and leaving the Satb2 population intact. The existence of hidden cortical lamination involving normal-looking neurons has also been recently reported in FCD type II [21]. Moreover, similar to FCD type II [16, 21, 35], DNs (and to a lesser extent GCs) in cortical tubers displayed variable expression of different cortical layer markers (Satb2, Cux2, RORβ, and Er81) regardless of their laminar location, supporting their altered migratory pattern. The immunophenotype of DNs in TSC, similar to FCD type II cases [35], resembles that of cortical projection neurons and suggests an alteration of a selected population of intermediate progenitor cells. Several hypotheses concerning the contribution of DNs and BCs/GCs to local pattern of cortical lamination have been proposed [52, 53], suggesting that both cell-autonomous and non-cell-autonomous mechanisms may be operative during corticogenesis in TSC brain.

Additional file

Additional file 1: Figure S1. A. Loss of neurons throughout all cortical layers in TSC cortical tubers and FCD type IIb. B. A larger amount of dysmorphic neurons can be detected in FCD type IIb compared to TSC cortical tubers. C. Significant increase in heterotopic neurons in the deep white matter in patients who underwent surgery for FCD Type IIb or cortical tuber resection. D. Quantitative analysis of Satb2+ neurons revealed significant overall cell loss in all epilepsy surgery specimens. E. Relative loss of Satb2+ neurons in TSC tissue. F. Quantitative increase in Tbr1+ neurons in perilesional cortex (PLx). G. Quantitative increase of Tbr1+ cells in the upper cortical layers. (TIF 26642 kb)

Competing interests

CMcK gave scientific advice to Eisai (Frankfurt, Germany) and UCB (Monheim, Germany); undertook industry-funded travel with the support of Eisai (Frankfurt, Germany), UCB (Monheim, Germany), Desitin (Hamburg, Germany), and Grifols (Frankfurt, Germany); obtained honoraria for speaking engagements from Eisai (Frankfurt, Germany), UCB (Monheim, Germany), Desitin (Hamburg, Germany), dialed (Köln, Germany), Fresenius Medical Care (Bad Homburg, Germany); and received research support from Astellas Pharma (Münichen, Germany), Octapharma (Langenfeld, Germany), dialed (Köln, Germany), and Fresenius Medical Care (Bad Homburg, Germany). The other authors declare that they have no competing interests.

Authors’ contributions

AM, AI, JS, and JA contributed to the selection and preparation of material for immunohistochemistry and in situ hybridization and assisted in the technical application. In situ hybridization and analysis of the data were performed by AM, FJ, TV, WS, FS, PK, JZ, CB, TP, HV, RC, IB, and EA helped in the collection of brain tissues and clinical data. EA, AI, JS, and MF helped AM in drafting and preparing the manuscript for submission. AM and EA conceived and designed the study. All authors read and approved the final manuscript.

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comparative tissue analysis of cortical tubers and perituberal cortex. Epilepsia. 2014;55(4):539-50. doi:10.1111/epi.12545.

41. Prabowo AS, Anink JJ, Lammens M, Nellist M, van den Ouweland AM, Adle-Biassette H, et al. Fetal brain lesions in tuberous sclerosis complex: TORC1 activation and inflammation. Brain Pathol. 2013;23(1):45–59. doi:10.1111/j.1750-3639.2012.00616.x.

42. Yoneshima H, Yamasaki S, Voelker CC, Molnar Z, Christophe E, Audinat E, et al. ErkB1 is expressed in a subpopulation of layer 5 neurons in rodent and primate neocortices. Neuroscience. 2006;137(2):401–12. doi:10.1016/j.neuroscience.2005.08.075.

43. Watakabe A, Ichinose N, Ohawa S, Hashikawa T, Komatsu Y, Rockland KS, et al. Comparative analysis of layer-specific genes in mammalian neocortex. Cereb Cortex. 2007;17(8):1918–33. doi:10.1093/cercor/bhl102.

44. Mühlebner A, Coras R, Kobow K, Feucht M, Czech T, Stefan H, et al. Neuropathologic measurements in focal cortical dysplasias: validation of the ILAE 2011 classification system and diagnostic implications for MRI. Acta Neuropathol. 2012;123(2):259–72. doi:10.1007/s00401-011-0920-1.

45. Abdijadid S, Mathern GW, Levine MS, Cepeda C. Basic mechanisms of epileptogenesis in pediatric cortical dysplasia. CNS Neurosci Ther. 2015;21(2):92–103. doi:10.1111/cns.12345.

46. Chandra PS, Salamon N, Nguyen ST, Chang JW, Huynh MN, Cepeda C, et al. Infantile spasm-associated microencephaly in tuberous sclerosis complex and cortical dysplasia. Neurology. 2007;68(6):438–45. doi:10.1212/01.wnl.0000252952.62543.20.

47. Iyer A, Prabowo A, Anink J, Spilet WG, van Rijen PC, Aronica E. Cell injury and premature neurodegeneration in focal malformations of cortical development. Brain Pathol. 2013. doi:10.1111/bpa.12060.

48. Andres M, Andre VM, Nguyen S, Salamon N, Cepeda C, Levine MS, et al. Human cortical dysplasia and epilepsy: an ontogenetic hypothesis based on volumetric MRI and NeuN neuronal density and size measurements. Cereb Cortex. 2005;15(2):194–210. doi:10.1093/cercor/bhh122.

49. Liu JY, Ellis M, Brooke-Ball H, de Tisi J, Eriksson SH, Brandner S, et al. High-throughput, automated quantification of white matter neurons in mild malformation of cortical development in epilepsy. Acta Neuropathol Commun. 2014;2:72. doi:10.1186/2051-5966-2-72.

50. Blumcke I, Vinters HV, Armstrong D, Aronica E, Thom M, Speafico R. Malformations of cortical development and epilepsies: neuropathological findings with emphasis on focal cortical dysplasia. Epileptic Disord. 2009;11(3):181–93. doi:10.1684/epid.2009.0261.

51. Yasin SA, Latack K, Becherini F, Ganapathi A, Miller K, Campos O, et al. Balloon cells in human cortical dysplasia and tuberous sclerosis: isolation of a pathological progenitor-like cell. Acta Neuropathol. 2010;120(1):85–96. doi:10.1007/s00401-010-0677-y.

52. Kim SH, Speirs CK, Solnica-Krezel L, Ess KC. Zebrafish model of tuberous sclerosis complex reveals cell-autonomous and non-cell-autonomous functions of mutant tuberin. Dis Model Mech. 2011;4(2):255–67. doi:10.1242/dmm.005587.