



REGULAR ARTICLE

A simple strategy for culturing morphologically-conserved rat hypothalamic tanycytes

Pablo Nicolás De Francesco¹ · Daniel Castrogiovanni² · Maia Uriarte¹ · Victoria Frassa¹ · Francina Agosti³ · Jesica Raingo³ · Mario Perello¹

Received: 25 August 2016 / Revised: 2 March 2017 / Accepted: 6 March 2017 / Published online: 17 April 2017
© Springer-Verlag Berlin Heidelberg 2017

Abstract Hypothalamic tanycytes are specialized bipolar ependymal cells that line the floor of the third ventricle. Given their strategic location, tanycytes are believed to play several key functions including being a selective barrier and controlling the amount of hypothalamic-derived factors reaching the anterior pituitary. The *in vitro* culture of these cells has proved to be difficult. Here, we report an improved method for the generation of primary cultures of rat hypothalamic tanycytes. Ependymal cultures were derived from tissue dissected out of the median eminence region of 10-day-old rats and cultured in a chemically defined medium containing DMEM:F12, serum albumin, insulin, transferrin and the antibiotic gentamycin. After 7 days *in vitro*, ~30% of the cultured cells exhibited morphological features of tanycytes as observed by phase contrast or scanning electron microscopy. Tanycyte-like cells were strongly immuno-reactive for vimentin and dopamine-cAMP-regulated phospho-protein (DARPP-32) and weakly immune-reactive for glial fibrillary acidic protein. Tanycyte-like cells displayed a stable negative

resting plasma membrane potential and failed to show spiking properties in response to current injections. When exposed to fluorescent beads in the culture medium, tanycyte-like cells exhibited a robust endocytosis. Thus, the present method effectively yields cultures containing tanycyte-like cells that resemble *in vivo* tanycytes in terms of morphologic features and molecular markers as well as electrical and endocytic activity. To our knowledge, this is the first protocol that allows the culturing of tanycyte-like cells that can be individually identified and that conserve the morphology of tanycytes in their natural physiological environment.

Keywords Cell culture · Ependymal cells · Hypothalamus · Median eminence · Glia

Introduction

The hypothalamic tanycytes are a specialized type of bipolar ependymal cells that form a unicellular epithelium lining the ventral part of the third ventricle, at the level of the median eminence (ME; Goodman and Hajihosseini 2015; Rodríguez et al. 2005). Based on their localization, the directionality of their processes and the expression of differential markers, four tanycyte subtypes can be recognized (Rodríguez et al. 2005). Tanycytes play important physiological functions. On the one hand, tanycytes form a selective barrier that controls the flow of molecules between the cerebrospinal fluid (CSF) and some hypothalamic nuclei and the ME (Rodríguez et al. 2005, 2010). It has been reported that tanycytes are able to take up circulating factors (i.e., leptin) that have diffused to the external layer of the ME from the fenestrated capillaries, mediating their transport into the CSF (Balland et al. 2014). Tanycytes also seem to facilitate the transport of molecules present in the CSF into specific hypothalamic nuclei by mechanisms

✉ Mario Perello
mperello@imbice.gov.ar

¹ Laboratory of Neurophysiology of the Multidisciplinary Institute of Cell Biology [IMBICE, dependent of the Argentine Research Council (CONICET) and Scientific Research Commission, Province of Buenos Aires (CIC-PBA)], Calle 526 entre 10 y 11, PO Box 403, La Plata 1900, Buenos Aires, Argentina

² Cell Culture Facility of the Multidisciplinary Institute of Cell Biology [IMBICE, dependent of the Argentine Research Council (CONICET) and Scientific Research Commission, Province of Buenos Aires (CIC-PBA)], La Plata, Buenos Aires, Argentina

³ Laboratory of Electrophysiology of the Multidisciplinary Institute of Cell Biology [IMBICE, dependent of the Argentine Research Council (CONICET) and Scientific Research Commission, Province of Buenos Aires (CIC-PBA)], La Plata, Buenos Aires, Argentina

dependent on transcytosis (Peruzzo et al. 2004; Rodríguez et al. 2005). In addition, tanycytes can modulate the activity of several neuroendocrine axes. For instance, tanycytes affect the thyroid axis activity since they express the ectoenzyme pyroglutamyl peptidase II, which degrades the thyrotropin-releasing hormone and regulates the amount of neuropeptide reaching the anterior pituitary, as well as the enzyme thyroxine deiodinase type II, which generates bioactive triiodothyronine from thyroxine (Fekete et al. 2000; Lazcano et al. 2015). Tanycytes also seem to regulate the release of gonadotropin-releasing hormone (GnRH) into the ME as their ablation prevents the pulsed release of this neuropeptide into the portal blood, the peak of luteinizing hormone and ovulation (Rodríguez et al. 2005). Despite the very important physiological roles of the tanycytes, the biology of these cells remains widely unexplored.

Cell cultures provide a simpler model to study complex biological systems that are hard to approach *in vivo*. The culture of hypothalamic tanycytes, however, has proved to be difficult to achieve due to several factors, including their limited amount *in vivo* and the cellular complexity of their normal environment, among others. Some research groups have developed methods to culture hypothalamic tanycytes (Chauvet et al. 1996; Gabrion et al. 1988). Gabrion and colleagues set up the conditions to culture dissociated cells derived from fetal mouse or rat hypothalami in serum-supplemented or chemically defined media and found that a small fraction of these cells were tanycytes. However, the major cell type present in the cultures corresponds to non-tanycytes ependymal cells, which steadily grow and form monolayers (Gabrion et al. 1988). Chauvet and colleagues also developed a culture strategy that has been extensively used by other researchers (García et al. 2003; Prevot et al. 2003). These cultures are derived from the ME of newborn 10-day-old rats and immunostaining patterns suggest that they mainly contain tanycytes (Chauvet et al. 1996). Importantly, this procedure uses medium supplemented with fetal bovine serum (FBS) and involves a secondary sub-culture. Thus, a high number of cells can be generated using this strategy but they seem to be less differentiated than *in vivo* tanycytes as they do not display their typical morphology. Here, we used a combination of the above referred procedures in order to obtain cell cultures enriched in tanycyte-like cells that could provide a useful tool to study this particular cell type.

Materials and methods

Animals

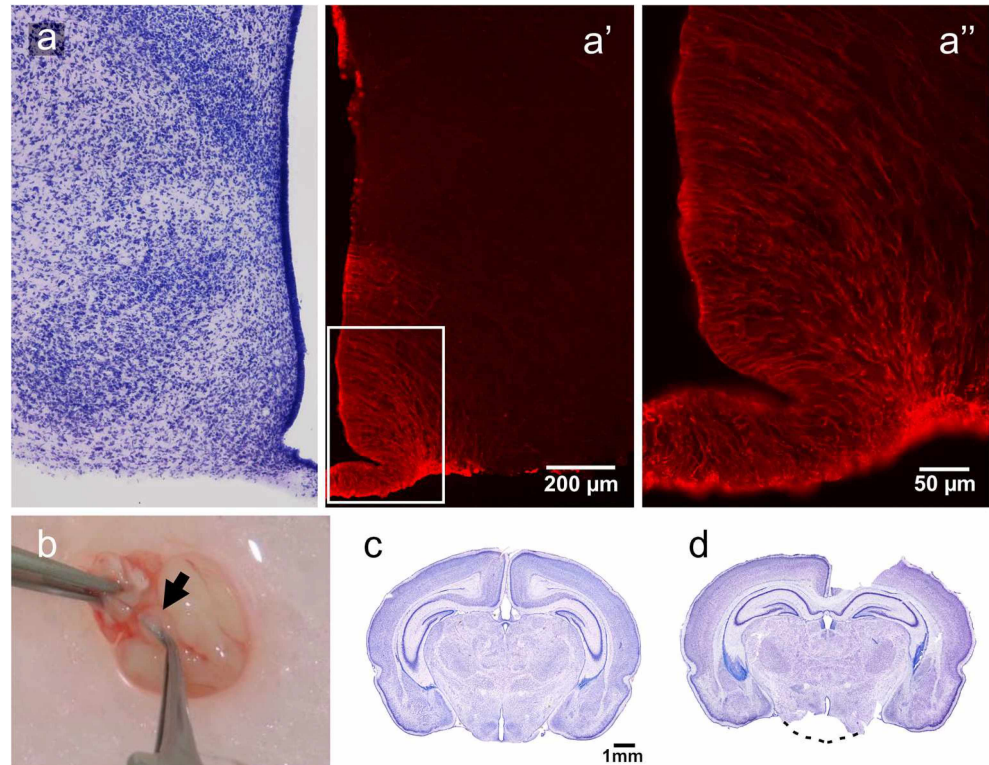
Sprague–Dawley rats were bred at the animal facility of the IMBICE. Animals were housed in a 12-h light/dark cycle with regular chow and water available *ad libitum*. This study was

carried out in strict accordance to the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Research Council and all efforts were made to minimize suffering. All experimentation received approval from the Institutional Animal Care and Use Committee of the IMBICE.

Cell cultures

The hypothalamic ME region was dissected from 10-day-old (P10) rats, in which tanycytes are already present lining the ventral part of the ventricle (Altman and Bayer 1978; De Seranno et al. 2004; Fig. 1a–a”). Specifically, rat brains were removed from the skull, cleaned of meninges and placed on their dorsal face to remove the median eminence–ARC region with fine-tip forceps (Fig. 1b). The optic chiasm and rostral edge of mammillary bodies were used as rostral and caudal limits, respectively, while ~1 mm from the lateral ends of the median eminence and 1 mm deep were used as lateral and ventral limits, respectively (Fig. 1c, d). Then, the resulting blocks of tissue containing the ME/arcuate nucleus (ARC) region were washed three times with Hank’s solution and afterwards cells were dissociated with a solution containing trypsin 0.25 mg/ml (Sigma-Aldrich, cat# T4799) and deoxyribonuclease I (DNAse I) 0.28 mg/ml (Sigma-Aldrich, cat# D5025,) at 37 °C for 10 min, including a light dissociation with a glass pipette after the first 5 min. Afterwards, the same volume of FBS (Internegocios S.A.) supplemented with DNAse I (5:1) was added to stop the digestion and the cells were spun for 5 min at 150 g and resuspended in culture medium (see composition below) plus 10% FBS and mechanically dissociated by forcing the suspension through glass pipettes and hypodermic needles with progressively smaller diameters. After these steps, the cell number and viability was assessed in a hemocytometer by mixing the cell suspension in a 1:1 ratio with a solution of trypan blue 0.4% in saline. Cells incorporating the dye were counted as dead. The cells were then centrifuged for 5 min at 150 g and resuspended in a suitable volume of serum-free culture medium for plating. Cells were plated on either 12-mm-diameter round cover glasses (50,000–60,000 cells) or in 22 × 22 mm cover glasses (75,000–100,000 cells). These cover glasses had been treated with poly-L-lysine 0.03 mg/ml (Sigma-Aldrich, cat# P8920) for 1 h at room temperature, washed three times with sterile water and air-dried prior to use. Cells were incubated at 37 °C in 5% CO₂ atmosphere with DMEM/F12 1:1 medium without phenol red (Microvet, cat# P3042-001) supplemented with 0.18 UI/ml insulin (Nordisk Pharm Ind), bovine serum albumin 50 µg/ml (Sigma-Aldrich, cat# A9418), transferrin 12.5 µg/ml (Calbiochem, cat# 616420) and 40 µg/ml gentamicin sulfate salt (Richtel). On day 3, half of the medium was replaced with a fresh one, plus the addition of thrombin to a

Fig. 1 **a, a'** Coronal slice from a P10 rat brain that was subjected to either Nissl staining or immunofluorescence staining for vimentin-Alexa 594, respectively. A higher magnification of the region outlined in (**a'**) is shown in (**a''**). Tancytic processes are already visible and fully structured at this stage. **b** The blunt dissection of the most ventral part of the hypothalamic region (*arrow*) using fine-tipped forceps. Coronal slice from a P10 rat brain before (**c**) and after (**d**) the ME/ARC collection procedure. *Dashed line* the missing portion of tissue



final concentration of 1.25 UI/ml (FloSeal, Baxter, cat# 934057). Cells were cultured for a total period of 7 days.

Neuroanatomical analysis of brain sections

Intact or ME/ARC-microdissected brains were used for neuroanatomical analysis. In this case, brains were fixed overnight with 4% formaldehyde in phosphate buffer saline (PBS; pH 7.4) and then cryoprotected by overnight immersion in 20% sucrose in PBS. Coronal sections of 40 μm were cut using a cryostat and used for either Nissl staining or immunohistochemistry. In the former case, the sections were mounted, stained with thionin (Sigma-Aldrich, cat# T7029), dehydrated in an ascending alcohol series, cleared in xylene and cover-slipped. For immunostaining, free floating sections were blocked with a solution containing 3% normal donkey serum and 0.25% Triton X-100 in PBS and then incubated with a chicken anti-vimentin antibody (Millipore, cat# AB5733, 1:2000) overnight at 4 $^{\circ}\text{C}$. Then, the sections were treated with an Alexa 594 goat anti-chicken antibody (Molecular Probes, cat# A11042, 1:2500) for 2 h at room temperature. Finally, the brain sections were sequentially mounted and cover-slipped with mounting medium. Fluorescence images were acquired with a Nikon Eclipse 50i microscope and a DS-Ri1 Nikon camera.

Acridine orange staining

Seven-day in vitro cultured cells were intravitally stained with acridine orange 1 μM (Sigma-Aldrich, cat# A6014) in culture medium for 15 min at 37 $^{\circ}\text{C}$ and washed afterwards with medium. The live-stained cells were directly observed with a Nikon TiS microscope equipped with a DS-Ri2 Nikon camera and epifluorescence and phase contrast images were taken. Acridine orange fluorescence (green to orange) was observed with a blue excitation low-pass filter cube (Nikon B-2A).

Oil Red O staining

Seven-day in vitro cultured cells were fixed with 4% paraformaldehyde in PBS at 4 $^{\circ}\text{C}$ and then washed once with PBS and once with 60% isopropyl alcohol in water. The cells were then stained in Oil Red O (Sigma-Aldrich, cat# O0625) 3 mg/ml in 60% isopropyl alcohol for 15 min. They were subsequently washed with 60% isopropyl alcohol, with PBS and finally cover-slipped in aqueous mountant. Cells were observed and photographed in brightfield mode with a Nikon Eclipse 50i microscope and a DS-Ri1 Nikon camera.

Scanning electron microscopy analysis

Seven-day in vitro cultured cells were fixed with 4% paraformaldehyde in PBS for 30 min at room temperature and then

gradually dehydrated stepwise through ethanol from 10 to 100%. The samples were dried in a critical-point CO₂ drying apparatus (Baltec CP-30) and coated with gold (Jeol Fine Ion Sputter Coater JFC-1100; Jeol, Akishima). Finally, the cells were examined with a JEOL scanning electron microscope (JSM 6360 LV) and images were captured with a Nikon D90 camera.

Immunocytochemical analysis

Seven-day in vitro cultured cells were fixed with 4% paraformaldehyde in PBS for 30 min at 4 °C. They were then treated with blocking solution (3% normal donkey serum and 0.25% Triton X-100 in PBS) and incubated with blocking solution containing antibodies against vimentin (mouse clone V9, Biogenex AM074-10 M ready-to-use, 1/10) and either dopamine-cAMP-regulated phospho-protein (DARPP-32, goat, Santa Cruz, sc-8483, 1:50) or glial fibrillary acidic protein (GFAP, rabbit, Sigma G9269, 1:200) overnight at 4 °C. Then, the cells were washed twice with PBS and incubated for 1 h at room temperature with the corresponding secondary antibodies conjugated to Alexa 488 or Alexa 594 (Molecular Probes, cat# A21202 1:1000, cat# A11058 1:500, or cat# A21207, 1:500 in blocking solution) and rinsed with PBS. For the assessment of the presence of neuronal cells, an antibody against NeuN (Millipore, mouse Mab377, 1:400) followed by an anti-mouse Alexa 594-conjugated antibody (Molecular Probes cat# A21203, 1:1000) were used. Also, for the assessment of the presence of microglial cells, an antibody against Iba1 (Wako, rabbit polyclonal cat# 019-19741, 1:2000) followed by an Alexa 594-conjugated antibody (Molecular Probes, anti-rabbit cat# A21207, 1:1000) were used. Cover glasses were mounted using an anti-fading solution containing bis-benzimide 10 mg/ml (Hoeschst 33258, Sigma-Aldrich, cat# B1155) to stain the cell nuclei. The slides were visualized within a week and stored at 4 °C. Fluorescence images were acquired with a Nikon Eclipse 50i microscope and a DS-Ri1 Nikon camera. FIJI was used to adjust the contrast and brightness of the microphotographs and to prepare the composite panels.

Electrophysiological analysis

Tanycyte-like cells were patched in a whole-cell configuration between 5 and 12 days in culture. The external solution composition was (in mM): 135 NaCl, 4.7 KCl, 1.2 MgCl₂, 2.5 CaCl₂, 10 HEPES and 10 glucose, pH = 7.4. The internal solution composition was (in mM): 140 potassium gluconate, 5 KCl₂, 5 NaCl, 10 HEPES, 4 MgCl₂, 3 Na-ATP, 0.3 Na-GTP and 0.6 EGTA, pH = 7.2. Borosilicate glass capillaries were used to generate the micropipettes. Leak currents less than 50 pA and resistances between 3 and 6 MΩ when electrodes were filled with internal solution were accepted. An Axopatch 200

amplifier (Molecular Devices) was used and data were sampled at 20 kHz and filtered at 10 kHz (−3 dB) using PCLAMP8.2 software. Measurement of resting membrane potential was done in track mode and membrane potential values that remained constant for at least 5 min during the recordings were included. These measurements were not corrected for the liquid junction potential. Voltage change recordings were done in current clamp mode using current injections from −100 to +100 pA in square pulses of 200 ms.

Endocytic activity analysis

Seven-day in vitro cell cultures were exposed to a suspension of 2 μg/ml carboxylate-modified 40-nm red fluorescent microspheres (Fluospheres, Molecular Probes, cat# F8793) in culture medium at 37 °C or 4 °C. After 15 min, the cells were washed twice with PBS, fixed with 4% paraformaldehyde in PBS for 30 min at 4 °C and mounted with anti-fading solution containing bis-benzimide 10 mg/ml. Widefield fluorescence image Z-stacks were acquired in 200-nm steps with a motorized automated rig on a Nikon Eclipse 50i microscope using a DS-Ri1 Nikon camera. The widefield datasets were deconvolved with Huygens (SVI). Immunostaining for Rab5 was also performed for the 37 °C condition following a procedure similar to that described above. A rabbit antibody against Rab5 (Cell Signaling, cat #3547, 1:1000) and an anti-rabbit Alexa 488-conjugated antibody (Invitrogen, cat# A-11008, 1:1000) were used. Images were acquired using a Zeiss AxioObserver D1 with an Apotome.2 module. FIJI was used to generate maximum intensity and orthogonal projections.

Statistical analysis

Twelve independent sets of cultures, which included a total of 135 pups, were performed using the above-described conditions. The analysis of each type of imaging study was performed on a minimum of 20–40 cells per cover glass, using 4–6 cover glasses that were derived from at least two different experiments. Data are expressed as the mean ± SEM.

Results

Overall performance of primary cultures

Typical average yield of the tissue dissociation was estimated in 60,000–90,000 cells per ME/ARC region collected. Cell viability monitored by trypan blue indicated that the vast majority of the cells were alive right before seeding. Cell cultures were plated using a density of ~150 cells/mm² (i.e., 75,000 cells in a 22 × 22 mm cover glass). A few minutes after plating, most cells were attached and showed a spherical

morphology with some of them already displaying a single and short cellular process. After 1 day in vitro, tancyte-like and non-tancyte-like cells were observed surrounded by aggregates of cell debris and dead cells. After 3 days in vitro, most cell debris and dead cells were removed when replacing part of the medium by fresh culture medium containing thrombin. After 4 days in vitro, a density of ~ 25 cells/mm² remained alive and healthy in the culture as indicated by phase contrast microscopy. Neither the cell density nor the overall appearance of the cell cultures changed significantly over the next 5 days in vitro and, therefore, the characterization of the cultured cells was performed by day 7 in vitro.

Morphological characterization of tancyte-like cells

At day 7 in vitro, a cell population with tancyte-like morphology could be easily visualized by phase contrast microscopy analysis (Fig. 2a–a'). The tancyte-like cells represented around 20–30% of the total number of cells. The non-tancyte-like cells included a heterogeneous group of flat and polygonal epithelial-like cells, round epithelial-like cells, fibroblast-like cells and macrophage-like cells. Importantly, no ciliated cells were found in the cell cultures, as assessed by high-magnification phase contrast observation. In addition, neither NeuN- nor Iba1-immunoreactive cells were found in the cultures, suggesting that neuronal and microglial cells were not present in the cultures. Tancyte-like cells represented a highly homogenous set of cells that displayed a nonciliated cell body and a long cytoplasmic shaft. The cell body of the tancyte-like cells was slightly elongated (average length 41.5 ± 1.8 μm , average width 19.5 ± 0.9 μm) and displayed an eccentric nucleus. The process of the tancyte-like cells varied both in length and diameter and typically ended as a swollen varicosity or footplate, which was never found in association with other cells of the culture. The cell length displays an asymmetric unimodal distribution with a median value of 124 μm (25% percentile: 93 μm ; 75% percentile: 190 μm) and a small fraction of cells extending past the 500- μm range. Phase contrast analysis showed that tancyte-like cells contained abundant highly refractive intracellular vesicle-like structures, which were bigger in the cell body and smaller in the processes (see Fig. 2a'). Within the cell body, these structures were preferentially located at the pole where the shaft forms as well as at the apical end. In order to test if these vesicles were acidic, such as lysosomes or late endosomes, cells were processed for acridine orange staining, which showed orange fluorescent compartments within the cell body, in most cases surrounding the nucleus, as well as in some small sections along the cell shaft and in the terminal foot plate (Fig. 2b–b'''). Notably, none of the highly refractive vesicles were of acidic nature upon comparison of fluorescence with phase contrast images. Since tancytes are known to contain lipids droplets (Flament-Durand and Brion

1985; Peruzzo et al. 2004; Rodríguez et al. 2005), the cells were also stained with the fat-soluble dye Oil Red O (Fig. 2c–c''). Oil red O staining revealed a strong red staining pattern coincident with the distribution of these highly refractive structures, which indicates the lipid-containing nature of these vesicles.

Scanning electron microscopy analysis showed that the cell body of tancyte-like cells displayed small spherical protrusions (or blebs) and extensive adhesion processes to the glass surface, while their cytoplasmic shafts displayed thin adhesion processes with small branching extensions, especially at the most distal part (Fig. 2d–e''). Thus, the experimental conditions described in the current study allow obtaining a cell culture containing tancyte-like cells that resemble some morphological features of tancytes in vivo.

Immunocytochemical characterization of tancyte-like cells

Cell cultures were characterized by immunocytochemical methods in order to test the presence of cell markers known to be present in tancytes in vivo. All tancyte-like cells were markedly positive for vimentin, even though this staining was also present in most of the other cells within the culture. This finding should be interpreted with caution, as vimentin can be induced in astroglial primary cultures by in vitro culture environments (Ciesielski-Treska et al. 1988). All tancyte-like cells were also immunoreactive for GFAP; however, the intensity of this signal was weaker as compared to the signal observed in a small fraction of cells present in culture that did not possess a tancyte-like morphology (Fig. 3a–c''). As expected, vimentin and GFAP stainings displayed a fairly similar pattern since both proteins have the same subcellular localization as part of the type III intermediate filaments and are known to form heteropolymers (Abd-el-Basset et al. 1992). Importantly, $\sim 90\%$ of tancyte-like cells positive for vimentin were also positive for DARPP-32 (Fig. 3d–d''), while the rest of the cells were not immunoreactive for this marker.

Electrophysiological characterization of tancyte-like cells

Whole-cell patch-clamp recordings of cultured tancyte-like cells indicated that they display an average resting membrane potential of -49.9 ± 4.7 mV ($n = 13$; Fig. 4a, b). As expected, tancyte-like cells failed to display either spontaneous fluctuations in their membrane potential or spike-like voltage changes in response to current injections ranging from -100 to $+100$ pA (Fig. 4c). Thus, tancyte-like cells display the electrophysiological response expected for this subtype of glial cell.

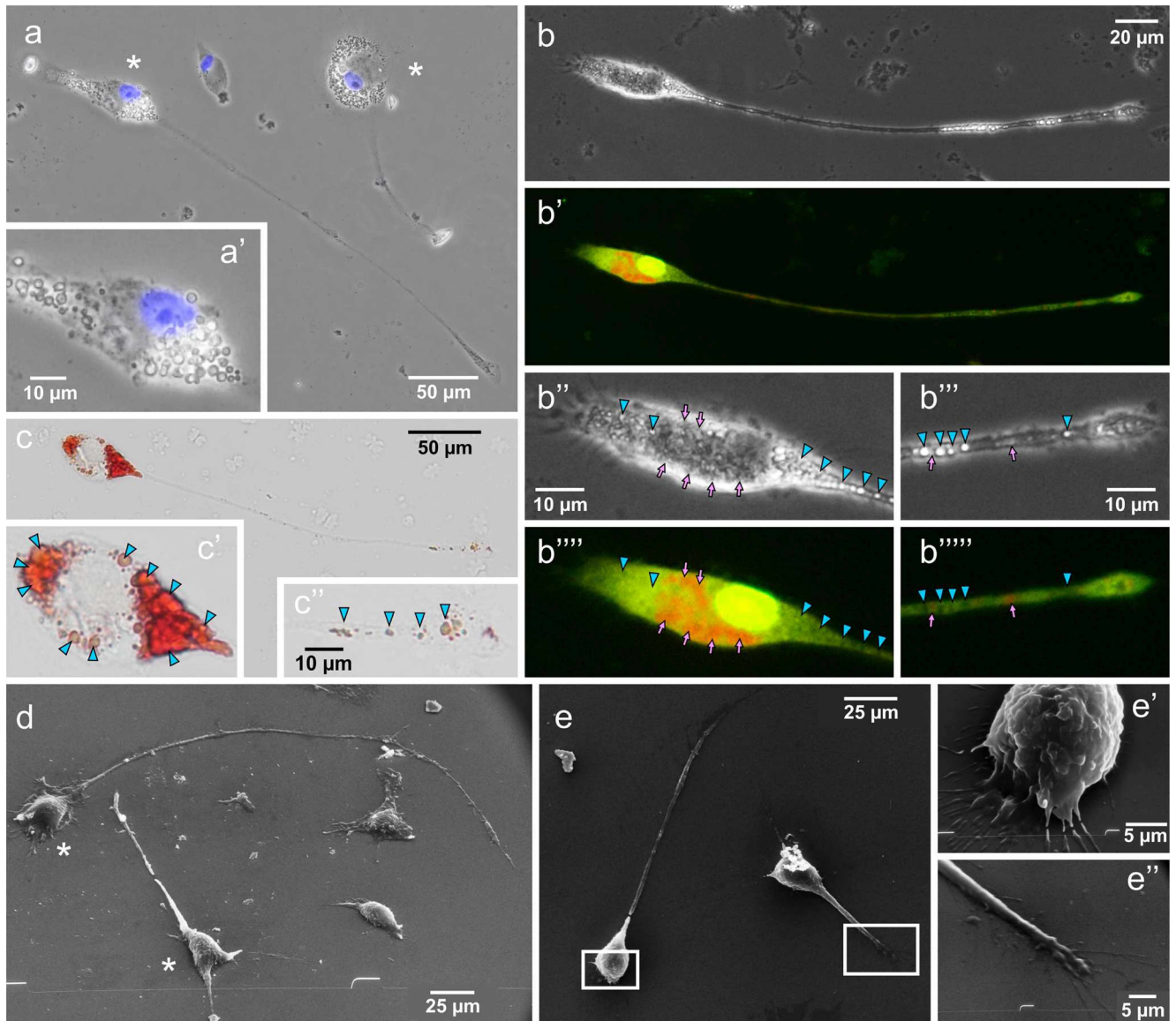


Fig. 2 Morphological and ultrastructural characterization of tancyte-like cells. **a** Representative phase contrast microscopy image from cells present in the cultures from day 7. The *grayscale* image is superimposed with a Hoechst fluorescent nuclear stain (*blue*). *Asterisks* label tancyte-like cells, while the other cell present in the image is a non-tancyte-like cell. A magnified inset is presented in (**a'**), showing the body of one of these cells where the intracellular presence of highly refractive vesicle-like structures is evident. **b** Representative fluorescence image of a tancyte-like cell after acridine orange staining. A corresponding phase contrast (**b'**) image is shown, along with paired *insets* depicting the cell body (**b''**, **b'''**) and the terminal portion of the projection with its foot plate (**b''''**, **b'''''**). *Cyan arrowheads* point at highly refractive vesicle-like structures, while *magenta arrows* point at acidic acridine orange-loaded

structures. **c** Representative image of a tancyte-like cell after staining with Oil Red O. *Insets* show a magnified view of the cell body (**c'**) and the terminal portion of the shaft (**c''**). *Cyan arrowheads* point at Oil Red O-stained vesicle-like structures. **d** Representative low-magnification scanning electron microscopy image from cells present in the cultures from day 7. Tancyte-like cells are labeled with an *asterisk* and present small branching extensions all along their body length. **e** Long and middle-length tancyte-like cells, along with higher magnification details (**e'**–**e''**). For the former, very thin adhesion processes are clearly visible at the distal end. The detail of the cell body shown in (**e'**) presents small cytoplasmic blebs and extensive adhesion processes. Magnified in (**e''**), the most distal part of the shaft has clear varicosities and branching

Endocytic activity of the tancyte-like cells

Since the hypothalamic tancytes are known to display endocytosis *in vivo* (Peruzzo et al. 2004), the active endocytosis was tested in tancyte-like cells. The incubation of primary cultures for 15 min at 37 °C with carboxylate-

modified red fluorescent microspheres led to the appearance of a punctate signal pattern within the cell bodies of several cell types within the culture including the tancyte-like cells (Fig. 5d–f). This intracellular punctate fluorescent pattern was not observed when primary cultures were incubated with fluorescent microspheres at 4 °C (Fig. 5a–c),

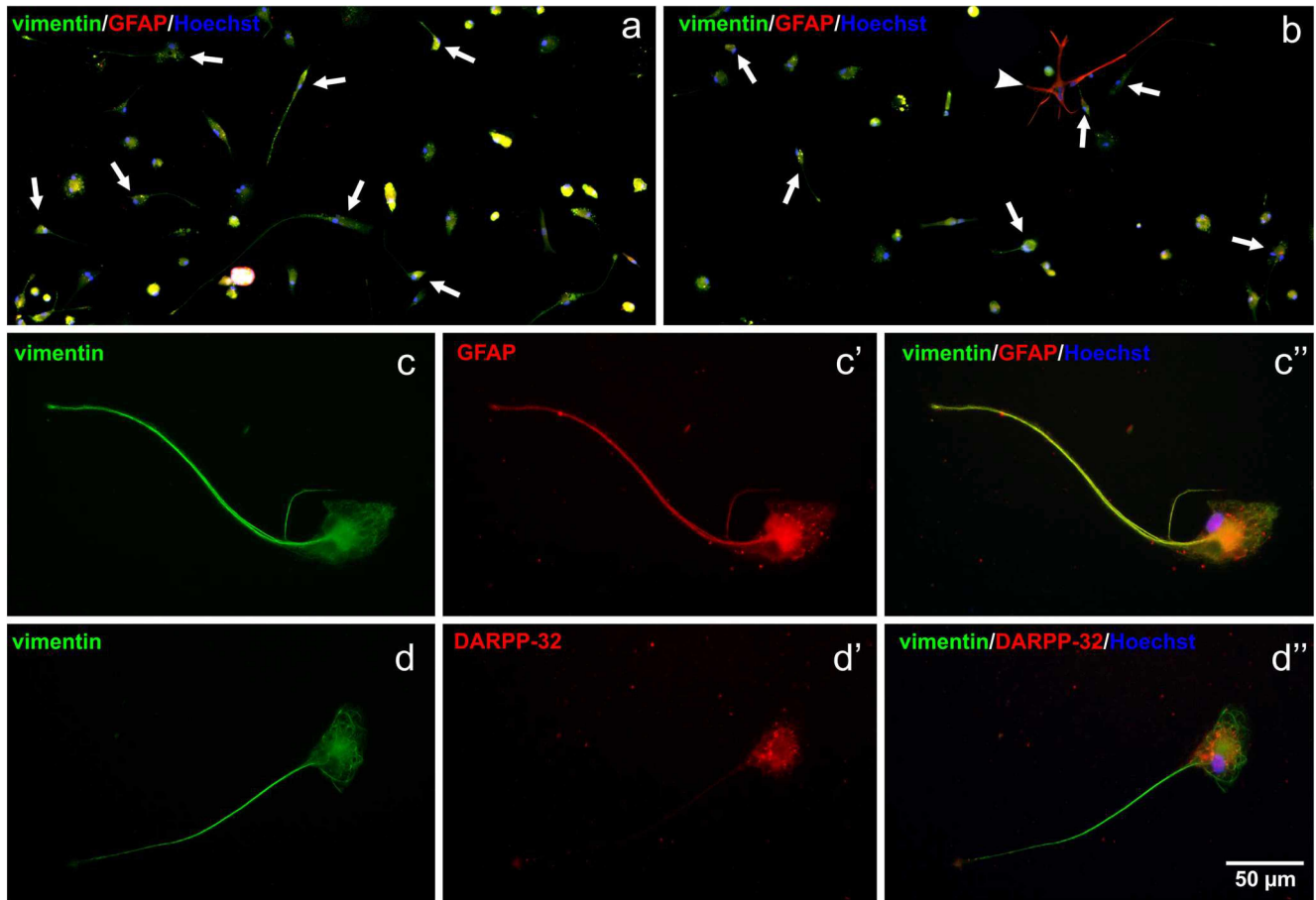


Fig. 3 Immunocytochemical characterization of tanyocyte-like cells. **a, b** Low-magnification images of cultures co-stained for vimentin and GFAP. Several tanyocyte-like cells can be observed, indicated by *arrows*. A group of strongly GFAP-positive non-tanycytes are indicated in **(b)** with an *arrowhead*. **c–c''** Representative image of a typical tanyocyte-like cell

where a double immunofluorescent staining for vimentin and GFAP was performed. **d–d''** A typical tanyocyte-like cell as seen after double-immunofluorescent staining for vimentin and DARPP-32. Composite images of the double staining plus nuclear Hoechst signal are also provided for each case

a temperature known to block active vesicular trafficking pathways. In order to test if internalized molecules were located in an endocytic compartment, we performed immunostaining against Rab5 in cell cultures previously incubated with microspheres at 37 °C. We found that a

significant fraction of the intracellular red fluorescent punctate co-stained with this early endocytic marker (Fig. 5g–j). Thus, tanyocyte-like cells seem to display active endocytic processes under the experimental conditions described here.

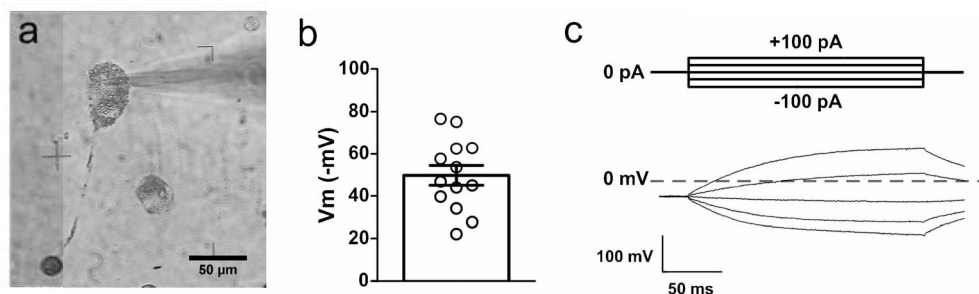


Fig. 4 Electrophysiological characterization of tanyocyte-like cells. **a** Representative image of a typical tanyocyte-like cell accessed by the pipette during recording. **b** Distribution of membrane potential values observed in cultured tanyocyte-like cells. **c** Representative voltage traces in

response to current injection performed on these cells. Recordings were done in current clamp mode applying current injections from –100 to +100 pA in square pulses of 200 ms

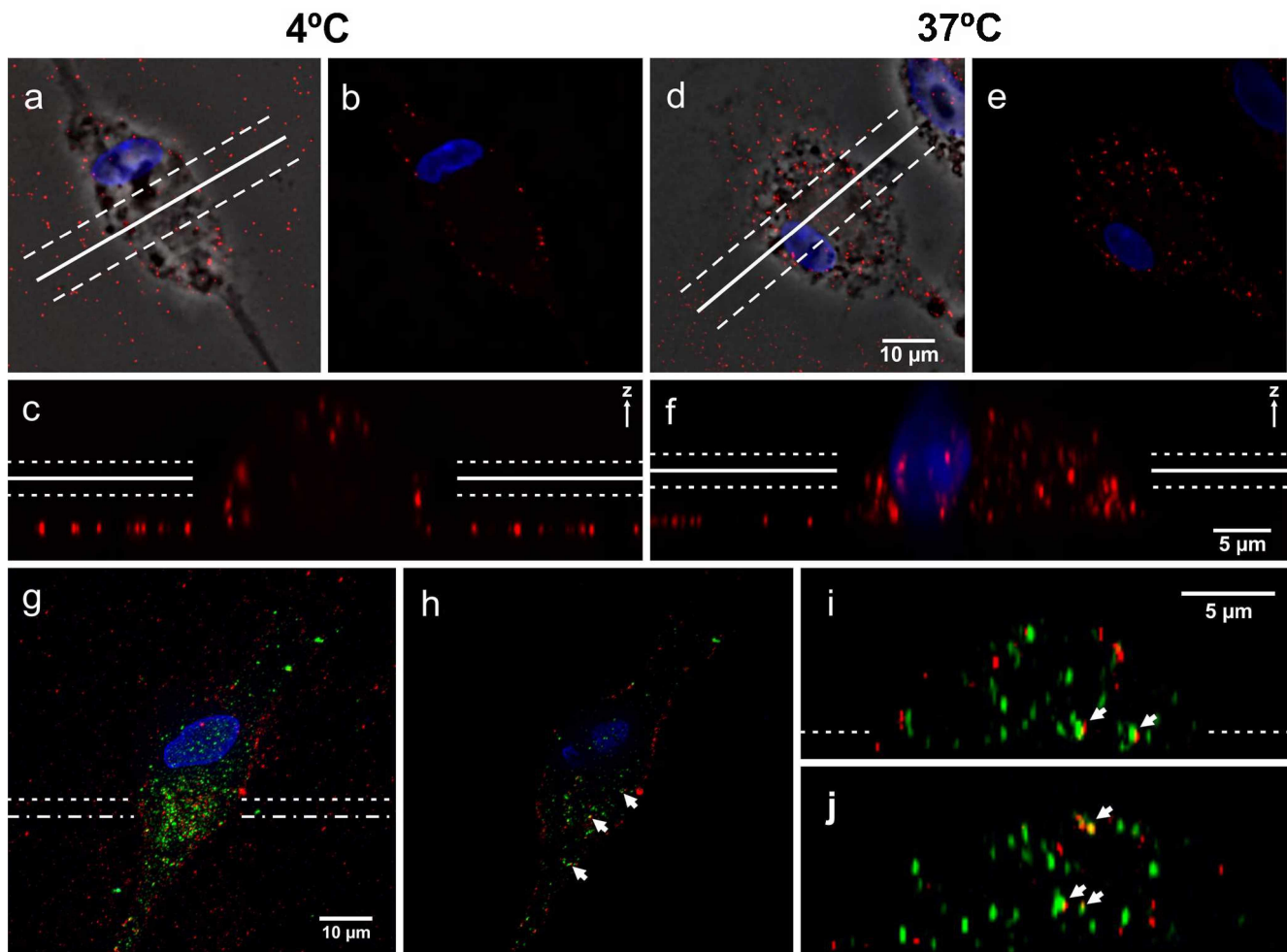


Fig. 5 Endocytic activity of tanyocyte-like cells. Cultured tanyocyte-like cells were incubated in the presence of red fluorescent microspheres for 15 min at 4 °C or 37 °C, fixed and a fluorescence image stack was acquired at 200-nm intervals and deconvolved. **a, d** Maximum intensity projection of the stacks for the 4 °C and 37 °C conditions, respectively. Corresponding phase contrast and Hoechst nuclear stain images are superimposed to better delineate the cell contour. **b, e** Maximum intensity projection of the same stacks but only including a 2- μ m range starting 2 μ m away from the glass slide, so to only include the bulk of the cytoplasm of each cell. The cell incubated at 4 °C only displays a punctate red fluorescent signal at the periphery, while the cell incubated at 37 °C also shows abundant intracellular signal. For these same cells, **c, f** show a 10-

μ m-wide orthogonal maximum intensity projection of planes passing through the middle portion of the cell body. Again, only the cell incubated at 37 °C show an extensive red fluorescent signal that corresponds to intracellular locations. Corresponding center slice and projection range for **(b)** and **(e)** are indicated in **(c)** and **(f)** with *continuous* and *dashed lines*, while the same is done for **(c)** and **(f)**, **(a)** and **(b)**. **g–j** show a similar experiment performed at 37 °C with fluospheres (*red*) in which an anti-Rab5 staining (*green*) was performed afterwards. **g** Maximum intensity projection. **h** Single image from the stack, at the level indicated by *dashed line* in **(i)**. **i, j** Orthogonal views at the levels indicated in **(g)** with *dashed* and *dot-dashed lines*, respectively. *Arrows* indicate fluospheres associated with Rab5 labeling

Discussion

Here, we used a combination of rigorous sample collection, based on neuroanatomical criteria as well as on the age of the animals, together with specific and defined culture conditions, in order to culture cells that resemble in vivo tanyocytes in terms of structural features, molecular markers and biological activity. Thus, we provide a simple and reproducible method for the primary culture of rat hypothalamic tanyocyte-like cells. To our knowledge, this is the first in vitro system in which tanyocyte-like cells can be individually identified and used to investigate some aspects of the cell biology of tanyocytes that

require morphologically conserved cells, such as the mechanisms mediating endocytosis or transcytosis.

The age of the animals at which we obtained the tissue for cultures was based on previous studies (Chauvet et al. 1996; Prevot et al. 2003) and seemed to be a critical aspect of the procedure. Thus, despite tanyocyte cultures having been described from earlier stages (Gabrion et al. 1988, 1998; Jordan et al. 1990), the P10 stage has resulted in the most successful and reproducible cultures in our hands. Here, we confirmed that the anatomical organization and the immunostaining patterns of hypothalamic tanyocytes of 10-day-old pups were very similar to those seen in adult rats. Similarly

to other central nervous system cells, the aptitude of tanyocyte-like cells to be cultured *in vitro* rapidly decreases with the age of the donor tissue (Chauvet et al. 1996). Indeed, we were unable to culture tanyocyte-like cells from adult rodents (unpublished observations). The reason why dispersed tanyocyte-like cells obtained from adult brains poorly survive the culture procedure is unclear but presumably involves the fact that mature tanyocytes are more susceptible to damage during dissociation, as has been already noted for glial and neuronal cell types (Chauvet et al. 1996). Although tanyocytes could also be cultured from ME/ARC tissue obtained from late embryonic stages, these cell cultures are less enriched in tanyocyte-like cells, likely due to a combination of facts including that the neuroanatomical limits to dissect the ME/ARC are harder to distinguish at prenatal stages and that the mitotic potential of other glial cells is higher at these ages (Rice and Barone 2000). In addition, tanyocytes from embryonic brains may not behave as mature tanyocytes since the ME functionality fully develops post-natally (Peruzzo et al. 2000). In balance, ME/ARC tissue from 10-day-old pups provides an excellent source of tanyocytes as they remain plastic enough to resist dissociation as well as further *in vitro* survival and also they better resemble mature tanyocyte physiology.

Here, we used a defined medium for culturing hypothalamic tanyocytes, a strategy that is in clear contrast to most of the previously described protocols that used culture media supplemented with animal serum. Chemically defined media allow the reduction of variability in the procedure, since all components of the system are specifically known. The defined medium was composed of DMEM:F12 supplemented with serum albumin, insulin, transferrin and gentamycin. As reported previously (Gabrion et al. 1998), thrombin was added to the culture medium from day 3. Thrombin is a serine protease involved in the coagulation cascade that improves the viability of ependymal cells without any effect on oligodendrocytes, macrophages and fibroblasts proliferation and has been described to act through the activation of the protease-activated receptor 1 (Perraud et al. 1987; Tritschler et al. 2007). Transferrin, serum albumin and insulin are common supplements to serum-free media and have been widely used in ependymal serum-free cultures (Gabrion et al. 1998; Grondona et al. 2013; Laabich et al. 1989; Tritschler et al. 2007; Weibel et al. 1984, 1986). Notably, culture conditions seem to mainly favor the survival of tanyocyte-like cells, since neurons, microglia and ciliated ependymal cells were not present in the cell cultures. However, culture conditions also seem to abrogate both the proliferative capacity of tanyocytes, since the number of tanyocyte-like cells was not increased during the cultures (see Chiasson et al. 1999) and their progenitor capabilities and neurogenic potential (Robins et al. 2013), since no neurospheres developed. We hypothesize that the lack of key growth factors in the serum-free medium blunted the proliferative potential of any cell type present in the culture and might

be responsible for the relatively low surface density of cells achieved. An intriguing possibility is that this low surface density is what fosters the differentiation of single tanyocyte-like cells as seen in our culture conditions. Thus, the current method seems to be a reproducible strategy to culture isolated tanyocyte-like cells that were already present in the *in vivo* context. Notably, ~70–80% of the cultured cells did not display a tanyocyte-like morphology. These non-tanyocyte-like cells, which scarcely divide or interact with other cells, likely represented a heterogeneous population of quiescent glial cells, which may include other subtypes of ependymal cells. The presence of other cells, different from tanyocytes, is inherent to the culture system and restricts its use to methodological approaches that involve assessments in individual cells, such as imaging techniques or electrophysiological recordings.

At the electrophysiological level, we found that tanyocyte-like cells showed a steady resting membrane potential and lacked spontaneous or evoked electrical excitability. We obtained a mean value of -50 mV, which is similar to the one previously informed for cultured tanyocytes. Hild and co-workers were the first to measure resting membrane voltage in cultured rat ependymal cells, reporting values between -40 and -60 mV (Hild et al. 1965). Later on, Jarvis and Andrew reported that rat tanyocytes from brain slices displayed a membrane potential ~ -80 mV and presumed that the less negative values obtained in cultured tanyocytes were due to membrane damage (Jarvis and Andrew 1988). Based on our recording conditions with minimum leak current and series resistance, we suggest that the difference of the membrane potential of tanyocytes recorded in brain slices and in culture is related to intrinsic aspects of each experimental model. One plausible explanation is that gap junction complexes, which closely connect tanyocytes *in vivo*, function as hemichannels in culture conditions and partially contribute to depolarize them, since these hemichannels can display non-selective cationic currents under some conditions (Bennett et al. 2003). Moreover, we found no electrical spike-like events in tanyocyte-like cells, whether spontaneous or in response to an intracellular injection of current. Thus, the electrical properties of *in vitro* tanyocyte-like cells agree with previous observations performed in this type of glial cell both *in vitro* and *ex vivo* and are clearly distinct from the ones expected for a neuronal phenotype.

Tanyocytes do not constitute a homogeneous cell population in the rat, in which four types of them have been distinguished (Rodríguez et al. 2005). Such morphological diversity of hypothalamic tanyocytes *in vivo* likely reflects differences in their functions. Thus, tanyocytes are classified as $\alpha 1$, $\alpha 2$, $\beta 1$ or $\beta 2$ depending on their location, morphology, molecular markers, ultrastructure and functions. The $\alpha 1$ tanyocytes are located in the middle portion of the ventricle and project their process to the ventromedial nucleus, while the $\alpha 2$ tanyocytes are observed ventrally and project into the ARC. The $\beta 1$ and $\beta 2$ tanyocytes are, respectively, located at the lateral or medial portion of the

floor of the third ventricle and send their projections ventrolaterally ($\beta 1$) or ventrally ($\beta 2$) to the plexus of portal fenestrated capillaries located in the external layer of the ME. The tancyte-like cells cultured in the described conditions display several features that resemble those of *in vivo* tancytes. At a morphological level, tancyte-like cells displayed a medium size cell body and a single and long process that ends in a footplate that bears an overall resemblance to that seen *in vivo*, although lacking the extensive branching usually observed. In terms of the molecular markers for tancytes, cultured tancyte-like cells displayed vimentin and DARPP-32 immunoreactive signals, as well as weak GFAP immunoreactivity. Vimentin labeling is a common feature to all ependymal cells, including tancytes (Chauvet et al. 1998). GFAP expression in tancytes has been previously demonstrated (Bascó et al. 1981; de Vitry et al. 1981); however, the immunoreactive signal for GFAP in tancytes is much lower than in astrocytes (Chauvet et al. 1995, 1996; Meister et al. 1988) and, consequently, tancytes have sometimes been regarded as negative for this marker (Chauvet et al. 1998; de Vries et al. 2014). More recently, GFAP expression in mice tancyte subtypes has been mapped and mainly found in dorsal $\alpha 2$ and some $\alpha 1$ subtypes, while little or no signal was seen in ventral $\alpha 2$ or β subtypes (Robins et al. 2013). Thus, our finding of weak GFAP staining in cultured tancyte-like cells together with their strong vimentin immunoreactivity is in agreement with their origin from the mediobasal hypothalamus and suggests that they are more ventral tancyte subtypes, namely $\alpha 2$ or β . Importantly, DARPP-32 is only expressed in the hypothalamic tancytes of the adult rat brain (Meister et al. 1988; Ouimet et al. 1984). Thus, the presence of DARPP-32 in cultured tancyte-like cells strongly indicates that they are indeed tancytes.

Notably, *in vitro* tancyte-like cells contained numerous and large lipid inclusions. Interestingly, *in vivo*, only $\beta 1$ tancytes contained large lipid droplets, which are in close association with the smooth endoplasmic reticulum (Rodríguez et al. 1979). These inclusions are thought to be involved in the synthesis of prostaglandins, which in turn regulates GnRH release in the ME (Rodríguez et al. 2005). Although it is unclear if the presence of the lipid droplets in all tancyte-like cells is due to the culture system or if the culture conditions favor the survival of some subtype/s of tancytes, the fact that our tissue collection procedure mainly involves the medio-basal hypothalamus would agree with the idea that the presently reported cultures are enriched in β tancytes.

Scanning electron microscopy analysis allowed the observation of key morphological features of *in vitro* tancyte-like cells that can be compared with previous characterizations of tancytes *in situ*. Thus, the cell body of the *in vitro* tancyte-like cells displayed prominent blebs or spherical protrusions, which were described in the apical surface of tancytes *in vivo*, by both scanning and transmission electron

microscopy (Jarvis and Andrew 1988; Paull et al. 1977; Rinne 1966; Rodríguez et al. 1979; Wittkowski 1998). These protrusions facing the third ventricle had been prominently observed in ventral tancytes, namely β , which further supports the notion that our cultures are enriched in this tancyte subtype. Membrane blebbing is not only seen in *in vivo* tancytes but also in any cell type during apoptosis. Since *in vitro* tancyte-like cells fail to show any other morphological feature associated with apoptosis (e.g., cell shrinkage or nuclear fragmentation), the presence of blebs seems to be more suggestive of endocytic activity (Rodríguez et al. 2005). The presence of numerous acidic vesicles, which are presumably lysosomes or late endosomes, in the cell body of *in vitro* tancyte-like cells is also consistent with the existence of an endocytic machinery. In line with these morphological features, tancyte-like cells displayed uptake of extracellular material when the cell cultures were exposed to fluorescent beads. Furthermore, the presence of immunostaining against Rab5 associated with the intracellularly incorporated beads strongly suggests that this uptake indeed took place by endocytosis. Both clathrin- and caveolin-mediated endocytosis have been reported as the main mechanisms for the active uptake of small-sized (20–50 nm) polystyrene beads *in vitro*, depending on cell lineage (dos Santos et al. 2011; Rejman et al. 2004; Smith et al. 2012). Notably, these two endocytic mechanisms are not considered to be equally present in all tancyte subtypes (Peruzzo et al. 2004; Rodríguez et al. 2005). Thus, the described *in vitro* system could be valuable to gain insight into the molecular mechanisms mediating the endocytic and transcytotic properties of these cells. Future studies using specific markers for each tancyte subtype (see Goodman and Hajihosseini 2015) as well as functional evaluations may help to better characterize the nature of the cells reported here.

Acknowledgements This work was supported by grants of the National Agency of Scientific and Technological Promotion of Argentina to MP (PICT2011-2142) and to JR (PICT2013-1145).

We would like to thank Dr. Maria José Bellini, Lic. Micaela García and Dr. Camila Scorticati for kindly providing some of the antibodies for this work.

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

References

- Abd-el-Basset EM, Ahmed I, Kalnins VI, Fedoroff S (1992) Immunoelectron microscopical localization of vimentin and glial fibrillary acidic protein in mouse astrocytes and their precursor cells in culture. *Glia* 6:149–153. doi:10.1002/glia.440060209

- Altman J, Bayer SA (1978) Development of the diencephalon in the rat. III. Ontogeny of the specialized ventricular linings of the hypothalamic third ventricle. *J Comp Neurol* 182:995–1015
- Balland E, Dam J, Langlet F, Caron E, Steculorum S, Messina A, Rasika S, Falluel-Morel A, Anouar Y, Dehouck B, Trinquet E, Jockers R, Bouret SG, Prévot V (2014) Hypothalamic tanycytes are an ERK-gated conduit for leptin into the brain. *Cell Metab* 19:293–301. doi:10.1016/j.cmet.2013.12.015
- Bascó E, Woodhams PL, Hajós F, Balázs R (1981) Immunocytochemical demonstration of glial fibrillary acidic protein in mouse tanycytes. *Anat Embryol (Berl)* 162:217–222
- Bennett MVL, Contreras JE, Bukauskas FF, Sáez JC (2003) New roles for astrocytes: gap junction hemichannels have something to communicate. *Trends Neurosci* 26:610–617. doi:10.1016/j.tins.2003.09.008
- Chauvet N, Parmentier ML, Alonso G (1995) Transected axons of adult hypothalamo-neurohypophysial neurons regenerate along tanycytic processes. *J Neurosci Res* 41:129–144. doi:10.1002/jnr.490410115
- Chauvet N, Privat A, Alonso G (1996) Aged median eminence glial cell cultures promote survival and neurite outgrowth of cocultured neurons. *Glia* 18:211–223. doi:10.1002/(SICI)1098-1136(199611)18:3<211::AID-GLIA5>3.0.CO;2-1
- Chauvet N, Prieto M, Alonso G (1998) Tanycytes present in the adult rat mediobasal hypothalamus support the regeneration of monoaminergic axons. *Exp Neurol* 151:1–13. doi:10.1006/exnr.1998.6784
- Chiasson BJ, Tropepe V, Morshead CM, van der Kooy D (1999) Adult mammalian forebrain ependymal and subependymal cells demonstrate proliferative potential, but only subependymal cells have neural stem cell characteristics. *J Neurosci* 19:4462–4471
- Ciesielski-Treska J, Goetschy JF, Ulrich G, Aunis D (1988) Acquisition of vimentin in astrocytes cultured from postnatal rat brain. *J Neurocytol* 17:79–86
- De Seranno S, Estrella C, Loyens A, Cornea A, Ojeda SR, Beauvillain J-C, Prévot V (2004) Vascular endothelial cells promote acute plasticity in ependymogial cells of the neuroendocrine brain. *J Neurosci* 24:10353–10363. doi:10.1523/JNEUROSCI.3228-04.2004
- de Vitry F, Picart R, Jacque C, Tixier-Vidal A (1981) Glial fibrillary acidic protein. A cellular marker of tanycytes in the mouse hypothalamus. *Dev Neurosci* 4:457–460
- de Vries EM, Kwakkel J, Eggels L, Kalsbeek A, Barrett P, Fliers E, Boelen A (2014) NF κ B signaling is essential for the lipopolysaccharide-induced increase of type 2 deiodinase in tanycytes. *Endocrinology* 155:2000–2008. doi:10.1210/en.2013-2018
- dos Santos T, Varela J, Lynch I, Salvati A, Dawson KA (2011) Effects of transport inhibitors on the cellular uptake of carboxylated polystyrene nanoparticles in different cell lines. *PLoS ONE* 6, e24438. doi:10.1371/journal.pone.0024438
- Fekete C, Mihály E, Herscovici S, Salas J, Tu H, Larsen PR, Lechan RM (2000) DARPP-32 and CREB are present in type 2 iodothyronine deiodinase-producing tanycytes: implications for the regulation of type 2 deiodinase activity. *Brain Res* 862:154–161
- Flament-Durand J, Brion JP (1985) Tanycytes: morphology and functions: a review. *Int Rev Cytol* 96:121–155
- Gabrior J, Peraldi S, Faivre-Bauman A, Klotz C, Ghandour MS, Paulin D, Assenmacher I, Tixier-Vidal A (1988) Characterization of ependymal cells in hypothalamic and choroidal primary cultures. *Neuroscience* 24:993–1007
- Gabrior JB, Herbuté S, Bouillé C, Maurel D, Kuchler-Bopp S, Laabich A, Delaunoy JP (1998) Ependymal and choroidal cells in culture: characterization and functional differentiation. *Microsc Res Tech* 41:124–157. doi:10.1002/(SICI)1097-0029(19980415)41:2<124::AID-JEMT3>3.0.CO;2-U
- García MA, Millán C, Balmaceda-Aguilera C, Castro T, Pastor P, Montecinos H, Reinicke K, Zúñiga F, Vera JC, Oñate SA, Nualart F (2003) Hypothalamic ependymal-glial cells express the glucose transporter GLUT2, a protein involved in glucose sensing. *J Neurochem* 86:709–724
- Goodman T, Hajihosseini MK (2015) Hypothalamic tanycytes—masters and servants of metabolic, neuroendocrine, and neurogenic functions. *Front Neurosci* 9:387. doi:10.3389/fnins.2015.00387
- Grondona JM, Granados-Durán P, Fernández-Llebrez P, López-Ávalos MD (2013) A simple method to obtain pure cultures of multiciliated ependymal cells from adult rodents. *Histochem Cell Biol* 139:205–220. doi:10.1007/s00418-012-1008-2
- Hild W, Takenaka T, Walker F (1965) Electrophysiological properties of ependymal cells from the mammalian brain in tissue culture. *Exp Neurol* 11:493–501
- Jarvis CR, Andrew RD (1988) Correlated electrophysiology and morphology of the ependyma in rat hypothalamus. *J Neurosci* 8:3691–3702
- Jordan FL, Rieke GK, Hughes BW, Thomas WE (1990) Morphological diversity of ependymal cells in tissue culture. *Brain Res Bull* 25:159–163
- Laabich A, Sensenbrenner M, Delaunoy JP (1989) Monolayer cultures of ependymal cells on porous bottom dishes. A tool for transport studies across the brain cerebrospinal barrier. *Neurosci Lett* 103:157–161
- Lazcano I, Cabral A, Uribe RM, Jaimes-Hoy L, Perello M, Joseph-Bravo P, Sánchez-Jaramillo E, Charli J-L (2015) Fasting enhances pyroglutamate II activity in tanycytes of the mediobasal hypothalamus of male adult rats. *Endocrinology* 156:2713–2723. doi:10.1210/en.2014-1885
- Meister B, Hökfelt T, Tsuruo Y, Hemmings H, Ouimet C, Greengard P, Goldstein M (1988) DARPP-32, a dopamine- and cyclic AMP-regulated phosphoprotein in tanycytes of the mediobasal hypothalamus: distribution and relation to dopamine and luteinizing hormone-releasing hormone neurons and other glial elements. *Neuroscience* 27:607–622
- Ouimet CC, Miller PE, Hemmings HC, Walaas SI, Greengard P (1984) DARPP-32, a dopamine- and adenosine 3':5'-monophosphate-regulated phosphoprotein enriched in dopamine-innervated brain regions. III. Immunocytochemical localization. *J Neurosci* 4:111–124
- Paull WK, Martin H, Scott DE (1977) Scanning electron microscopy of the third ventricular floor of the rat. *J Comp Neurol* 175:301–310. doi:10.1002/cne.901750305
- Perraud F, Besnard F, Sensenbrenner M, Labourdette G (1987) Thrombin is a potent mitogen for rat astroblasts but not for oligodendroblasts and neuroblasts in primary culture. *Int J Dev Neurosci* 5:181–188
- Peruzzo B, Pastor FE, Blázquez JL, Schöbitz K, Peláez B, Amat P, Rodríguez EM (2000) A second look at the barriers of the medial basal hypothalamus. *Exp Brain Res* 132:10–26
- Peruzzo B, Pastor FE, Blázquez JL, Amat P, Rodríguez EM (2004) Polarized endocytosis and transcytosis in the hypothalamic tanycytes of the rat. *Cell Tissue Res* 317:147–164. doi:10.1007/s00441-004-0899-1
- Prévot V, Cornea A, Mungenast A, Smiley G, Ojeda SR (2003) Activation of erbB-1 signaling in tanycytes of the median eminence stimulates transforming growth factor beta1 release via prostaglandin E2 production and induces cell plasticity. *J Neurosci* 23:10622–10632
- Rejman J, Oberle V, Zuhorn IS, Hoekstra D (2004) Size-dependent internalization of particles via the pathways of clathrin- and caveolae-mediated endocytosis. *Biochem J* 377:159–169. doi:10.1042/BJ20031253
- Rice D, Barone S (2000) Critical periods of vulnerability for the developing nervous system: evidence from humans and animal models. *Environ Health Perspect* 108(Suppl 3):511–533
- Rinne UK (1966) Ultrastructure of the median eminence of the rat. *Z Zellforsch Mikrosk Anat* 74:98–122
- Robins SC, Stewart I, McNay DE, Taylor V, Giachino C, Goetz M, Ninkovic J, Briancon N, Maratos-Flier E, Flier JS, Kokoeva MV,

- Placzek M (2013) α -Tanycytes of the adult hypothalamic third ventricle include distinct populations of FGF-responsive neural progenitors. *Nat Commun* 4:2049. doi:10.1038/ncomms3049
- Rodríguez EM, González CB, Delannoy L (1979) Cellular organization of the lateral and postinfundibular regions of the median eminence in the rat. *Cell Tissue Res* 201:377–408
- Rodríguez EM, Blázquez JL, Pastor FE, Peláez B, Peña P, Peruzzo B, Amat P (2005) Hypothalamic tanycytes: a key component of brain-endocrine interaction. *Int Rev Cytol* 247:89–164. doi:10.1016/S0074-7696(05)47003-5
- Rodríguez EM, Blázquez JL, Guerra M (2010) The design of barriers in the hypothalamus allows the median eminence and the arcuate nucleus to enjoy private milieus: the former opens to the portal blood and the latter to the cerebrospinal fluid. *Peptides* 31:757–776. doi:10.1016/j.peptides.2010.01.003
- Smith PJ, Giroud M, Wiggins HL, Gower F, Thorley JA, Stolpe B, Mazzolini J, Dyson RJ, Rappoport JZ (2012) Cellular entry of nanoparticles via serum sensitive clathrin-mediated endocytosis, and plasma membrane permeabilization. *Int J Nanomedicine* 7:2045–2055. doi:10.2147/IJN.S29334
- Tritschler F, Murín R, Birk B, Berger J, Rapp M, Hamprecht B, Verleysdonk S (2007) Thrombin causes the enrichment of rat brain primary cultures with ependymal cells via protease-activated receptor 1. *Neurochem Res* 32:1028–1035. doi:10.1007/s11064-006-9267-8
- Weibel M, Pettmann B, Daune G, Labourdette G, Sensenbrenner M (1984) Chemically defined medium for rat astroglial cells in primary culture. *Int J Dev Neurosci* 2:355–366. doi:10.1016/0736-5748(84)90072-8
- Weibel M, Pettmann B, Artault JC, Sensenbrenner M, Labourdette G (1986) Primary culture of rat ependymal cells in serum-free defined medium. *Brain Res* 390:199–209
- Wittkowski W (1998) Tanycytes and pituicytes: morphological and functional aspects of neuroglial interaction. *Microsc Res Tech* 41:29–42. doi:10.1002/(SICI)1097-0029(19980401)41:1<29::AID-JEMT4>3.0.CO;2-P