

## REVIEW

# Pheochromocytoma/paraganglioma preclinical models: which to use and why?

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## Abstract

Pheochromocytomas/paragangliomas (PPGLs) are rare neuroendocrine tumours linked to more than 15 susceptibility genes. PPGLs present with very different genotype/phenotype correlations. Certainly, depending on the mutated gene, and the activated intracellular signalling pathways, as well as their metastatic potential, each tumour is immensely different. One of the major challenges in *in vitro* research, whatever the study field, is to choose the best cellular model for that study. Unfortunately, most of the time there is not 'a best' cell model. Thus, in order to avoid observations that could be related to and/or dependent on a specific cell line, researchers often perform the same experiments using different cell lines simultaneously. The situation is even more complicated when there are only very few cell models obtained in different species for a disease. This is the case for PPGLs. In this review, we will describe the characteristics of the different cell lines and of mouse models, trying to understand if there is one that is more appropriate to use, depending on which aspect of the tumours one is trying to investigate.

## Key Words

- ▶ : pheochromocytoma/paraganglioma
- ▶ cell models
- ▶ animal models

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## Introduction

In 2017, the World Health Organization (WHO) of Endocrine Tumours recognised anatomic criteria for PPGL classification (1). Pheochromocytomas (PCCs) are tumours originating from the neural crest-derived chromaffin cells within the adrenal medulla. They commonly produce one or more catecholamines: dopamine (DA), noradrenaline (NA or norepinephrine) or adrenaline (A or epinephrine). The presence of secretory granules at ultrastructure is a diagnostic feature of pheochromocytoma. Paragangliomas (PGLs) derive from extra-adrenal paraganglia cells and can be further classified according to their clinical and biological characteristics. In fact, they can be categorised into two main classes. The first one is Head and Neck (HN)-PGLs. They owe their name to their usual association with branches of the vagus and glossopharyngeal nerves, and typically lack catecholamine secretion (2). They are mostly found at the bifurcation of the common carotid artery, where the carotid body is located, but they may

also form in the jugular, tympanic, vagal, or laryngeal paraganglia (3) or in other locations (4, 5). The second class is Sympathetic PGLs that form in sympathetic paraganglia and are biochemically active, since they secrete catecholamines. Despite showing a ubiquitous distribution, from the skull base to the pelvic floor, approximately 85% develop below the diaphragm (3). The standard treatment of PPGLs is the surgical removal of the tumour (6). Since there is no histological system that is currently endorsed for understanding the biological aggressiveness of these tumours, all PPGLs have metastatic potential and their metastatic nature is determined by the presence or lack of distant metastases in sites where normal paraganglia are not found, more specifically bone and histologically confirmed lymph nodes (1). Unfortunately, there is no available cure for metastatic lesions, and understanding the cancer driver events is crucial for the development of targeted therapies.

## Pheochromocytomas and paragangliomas (PPGLs)

PCC and PPGLs are rare tumours with variable aggressiveness characterised by a large spectrum of hereditary predisposition. Indeed, over one-third of PPGLs are associated with inherited susceptibility genes, which is the highest rate among all tumour types (7). Mutations have been identified in more than 15 genes and, based on gene profiling, PPGLs can be assigned to three different 'clusters' (for recent reviews on the genetic/phenotype correlations of PPGLs see: (8, 9, 10)). PPGLs with mutations in genes encoding the hypoxia-inducible factor (HIF)  $2\alpha$ , von Hippel–Lindau tumour suppressor (VHL), prolyl hydroxylase domain (PHD), fumarate hydratase (FH), and succinate dehydrogenase subunits (SDHx) are included in cluster 1. These tumours are characterised by the activation of pseudohypoxic pathways and by an immature catecholamine-secreting phenotype. Intriguingly, tumours mutated in one of the enzymes involved in the Krebs cycle are more aggressive than others, showing an elevated metastatic potential. Cluster 2 includes PPGLs with mutations in the *RET* proto-oncogene, Neurofibromatosis type 1 (*NF1*) tumour suppressor gene, TransMEMbrane protein (*TMEM127*) gene, Harvey rat sarcoma viral oncogene homolog (*HRAS*) and MYC Associated Factor X (*MAX*) gene. Cluster 2 PPGLs show activated MAPK and mTOR signalling pathways, are mostly benign and exhibit a mature catecholamine phenotype with strong expression of phenylethanolamine *N*-methyltransferase (PNMT), which is the enzyme that converts NA to A. Recently, introducing also a cluster 3 has been suggested, which includes tumours associated with mutations in the Mastermind Like Transcriptional Coactivator 3 (*MAML3*) gene, which is involved in the Wnt signalling pathway. These PPGLs are characterised by a high Ki67 expression, aggressive behaviour and early metastatic dissemination.

### Tumour biology

As reported previously, a common feature of cluster 1 tumours is the activation of HIFs. For this reason, scientists refer to them as pseudohypoxia-driven tumours. Physiologically, in response to low extracellular oxygen levels (i.e. hypoxia), HIF activated pathways are induced. In cluster 1 tumours, HIF pathways are activated regardless of the extracellular oxygen concentration, a condition defined as pseudohypoxia. HIFs are heterodimeric

transcription factors and their inducible components (HIF1 $\alpha$  and HIF2 $\alpha$ ) are very closely regulated by proteasomal degradation, which occurs after their hydroxylation by the enzyme PHD1/2. Mutations in some of the susceptibility genes cause an impairment of the Krebs cycle, resulting in an accumulation of the oncometabolites succinate, fumarate, or 2-hydroxyglutarate, that in turn leads to the inhibition of PHD1/2. Inactivation of PHD1/2 results in a decrease of HIF- $\alpha$  hydroxylation that cannot be degraded any longer (11, 12, 13, 14). Therefore, cluster 1 mutations promote angiogenesis, tumour invasion, and metastasis.

Mutations of genes belonging to cluster 2 lead to activation of the phosphatidylinositol-3-kinase (PI3K)/AKT, mammalian target of rapamycin (mTORC1)/p70S6 kinase (p70S6K), and RAS/RAF/ERK signalling pathways, which in turn promote cell proliferation, survival, cancer development and angiogenesis.

The adrenal gland is the body's only significant source of circulating A. Tumours of cluster 2 are predominantly intra-adrenal and thus secrete A, while the extra-adrenal ones produce NA and/or dopamine. Conversely, cluster 1 tumours, even those occurring in the adrenal gland, which are often associated with *VHL* mutations, do not produce significant A (15).

Tumours mutated for the Cold Shock Domain-containing E1 (*CSDE1*) and the *MAML3* fusion genes belong to cluster 3. *MAML3* mutations lead to over-activation of Wnt and Hedgehog signalling and *MAML3*-mutated PPGLs showed elevated Ki-67 expression, aggressive behaviour and early metastatic spread. *CSDE1* mutations lead to over-activation of  $\beta$ -catenin, a target of Wnt signalling, and favour tumour proliferation, invasion, and metastases. The catecholamine phenotype of these tumours is rather unknown (15).

### Cell models

#### Rat pheochromocytoma (PC12)

The adrenal rat pheochromocytoma (PC12) cell line was originally isolated from a PCC developed in an irradiated rat in 1976 (16). This cell line has the characteristic of precursor cells for both sympathetic neurons and chromaffin cells. PC12 can differentiate towards a neuronal phenotype in response to nerve growth factor (NGF) (17), which causes these cells to extend long processes and become electrically excitable (18). In contrast, dexamethasone (DEX) treatment induces a more endocrine-like phenotype and upregulates catecholamine

synthesis and storage (19, 20). The responsiveness to NGF and DEX by PC12 cells has allowed them to be used as a model of neuronal differentiation and pluripotency. Chromaffin cells and sympathetic neurons derived from the neural crest and the lineages diverge very early, nevertheless, some transdifferentiation can occur (21, 22, 23). These cells are widely utilised as a model to study synthesis, storage, and secretion of catecholamines and regulation of nervous system development. In addition, PC12 cells are also used for the study of the differentiation effects on exocytotic vesicles and, since PC12 cells present some advantages over other cell models for neurosecretion, including chromaffin cells, it is likely that PC12 will continue to be used as a model to study exocytosis (24). PC12 cells are important also for PPGLs studies. Indeed, they express several of the catecholamine biosynthetic enzymes, including tyrosine hydroxylase (TH) along with the enzymes converting L-DOPA to DA and DA to NA, that is, aromatic L-amino acid decarboxylase (AADC) and DA  $\beta$ -hydroxylase (DBH), respectively. However, the final enzyme in the catecholamine biosynthetic pathway, PNMT, which produces A from NA, is not expressed in these cells. Indeed, PC12 cells are noradrenergic (16). Nevertheless, dexamethasone promotes the expression of the catecholamine biosynthetic enzymes, including PNMT (25). Of note, in these cells, functional MAX protein is not expressed. Hopewell and colleagues have identified a homozygous mutation of MAX gene, which is consistent with the aberrant processing of MAX transcripts in these cells. It has been suggested that the loss of MAX expression may have been a selected event either in the development of the rat PCC or during subsequent cell culturing *in vitro* (26).

### Mouse pheochromocytoma cell line (MPC) and mouse tumour tissue cells (MTT)

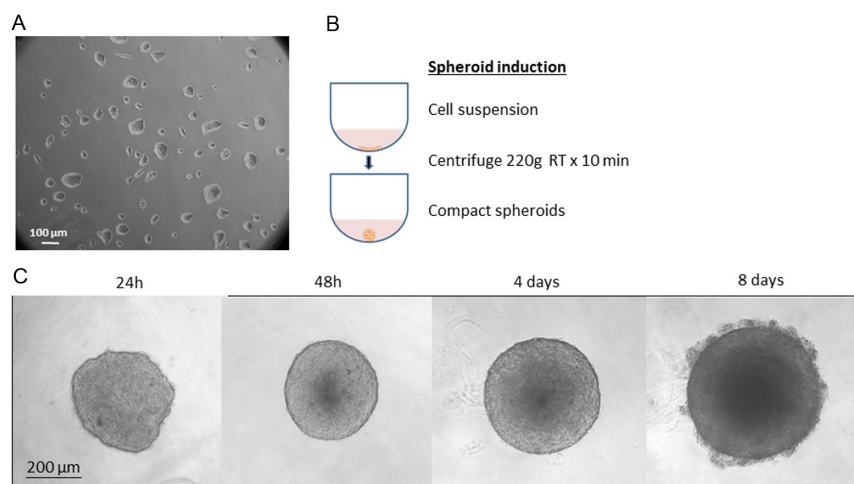
In 1994, Jacks *et al.* constructed a mouse strain with a knock-out (KO) mutation of *Nf1*, the murine counterpart of *NF1* (27). This mutation was associated with the frequent development of PCCs. Tischler *et al.* characterised these tumours and isolated cells cultured *in vitro* (28). A few years later, in the same laboratory, five mouse PCC cell lines (MPC) were stabilised from primary cultures of PCCs arising in mice with a *Nf1* heterozygous KO mutation. The cell line generally employed for the preclinical studies pertaining to human tumours is MPC4/30/PRR. These cells typically showed extensive spontaneous neuronal differentiation and expressed neurofibromin. Nevertheless, subsequent studies demonstrated that the

wild type *Nf1* allele was not present, possibly indicating that the protein produced was functionally defective (29).

MPC is a valid tool for studying genes and signalling pathways that regulate cell growth and differentiation in adrenal medullary neoplasms, and are a unique model for studying the regulation of PNMT expression (30). Indeed, a particularly interesting characteristic of the mouse PCCs is positive staining for PNMT and their ability to produce A. Moreover, MPC is also considered as a useful model for studying neurotransmitter release and neuroendocrine secretion (20).

The less differentiated derivative of MPC, designated as MTT (for mouse tumour tissue-derived) cells, was then established from MPC tumour tissue formed after reinjection of the original cell line into nude mice. These cells are characterised by the development of solid organ metastatic lesions, predominantly in the liver, 4 weeks after injection. The MTT cells maintain a PCC phenotype, as confirmed by measuring intracellular catecholamines, by evaluating the expression of TH and PNMT along with monitoring the presence of dense-core secretory granules. Furthermore, gene expression array analysis on MTT revealed genes that may be crucial for the development of an aggressive and more malignant phenotype of PCC (31). The MPC and MTT lines are complementary for drug testing purposes, in that MTT best reflects aggressive metastases, while MPC is better differentiated and more comparable to slow-growing, hormonally active metastases (32). Another interesting aspect of MPC and MTT is that they do not express HIF2 $\alpha$ , but only HIF1 $\alpha$  (13).

Several cell lines were then derived from MPC and MTT to improve and diversify research studies, with a particular interest in *SDHB* mutations, because mutations in this gene represent a high-risk factor for malignancy and poor prognosis (33, 34). To assess possible associations between *SDHB* gene mutations and invasiveness, Richter and colleagues established an MTT *SDHB* knockdown by viral transduction with lentiviral particles (35). Of note, one of the shortcomings of silencing the *SDHB* subunit in an *Nf1* heterozygous KO mouse is the mixing of the phenotypes of clusters 1 and 2. By exploiting the MPC and MTT characteristic of spontaneously forming clusters in cultures, we generated, for the first time, PCC spheroids with homogeneous size and geometry, as shown in Fig. 1 (36). In contrast to cultures, tumour cell spheroids mimic the tumour mass and better represent the cells' organisation within the tumour, forming a necrotic core, an external zone with proliferating cells and clear margins. This characteristic allows the operator to easily manipulate

**Figure 1**

Cluster and spheroid formation. (A) Representative image of MTT cells that spontaneously form clusters when cultured in monolayer. (B) Induction of spheroids by centrifugation of  $5 \times 10^3$  MTT cells/well for 10 min at 220 *g* in 96-well plate with low adherence round bottom. (C) After 48 h, spheroids assume a compact and rigid shape with clear edges, and they can be easily manipulated. In time, it is possible to observe an increase of spheroids mass, and the formation of a necrotic core surrounded by a proliferating cell crown. All images are representative.

the spheroids and use them for performing experiments. Moreover, chromaffin cell features of PCC cell lines remain unchanged during cultivation under spheroid conditions (37). However, multicellular spheroids have some limitations. Although their spherical architecture is quite similar to the 'Zellballen' pattern of PCCs, every spheroid has central necrosis, which is absent in most PCCs or present only focally, especially in aggressive tumours. The presence of this core could be a confusing factor for a clear distinction between pseudohypoxic cluster 1 and cluster 2. Nevertheless, it has been shown that three-dimensional tumour cell spheroids provide an excellent *in vitro* model to study the influence of hypoxia under conditions close to the *in vivo* situation (37). Indeed, currently not much is known about spheroids maintained in hypoxic conditions and surely some further studies should be performed to take advantage of this research field. Another advantage of using spheroids is the excellent proven suitability of this model for drug screenings (38, 39).

### Progenitor cells derived from a human pheochromocytoma (hPheo1)

Previous reports have shown that it is possible to immortalise cells, with minimal alteration of the phenotype, by introducing human telomerase reverse transcriptase (hTERT) into human cells (40, 41). In 2013, Ghayee *et al.* applied this technology in an attempt to immortalise endocrine tumours and to develop a cell line from a human PCC. In the end, they developed a neuroendocrine progenitor cell line called hPheo1, which should be useful in dissecting the molecular pathways that influence the growth and differentiation leading to PCC (42). The characterisation of these hPheo1 cells showed that genes associated with catecholamine synthesis were

highly expressed in the tumour tissue of origin, but most of them were downregulated in hPheo1 cells. Nevertheless, when hPheo1 cells were treated with NGF, they were able to develop neurites and to express chromogranin A and PNMT. Despite the lack of hormone production, these cells might be useful for studying signalling pathways controlling growth and metastasis as well as for studying mechanisms of tumour development (43).

### Immortalised chromaffin cells (imCC)

In 2013, Letouzé *et al.* generated an immortalised mouse chromaffin cell line (imCC) harbouring a complete defect in SDH. They created genetically modified mice in which the endogenous *SDHB* exon 2 was flanked by LoxP sites. Subsequently, chromaffin cells were isolated from the adrenal medulla, and deleted for *SDHB* by Cre-mediated recombination (12). This model was created in order to investigate the mechanisms linking SDH deficiency with tumour onset, and it is used to study the metabolic plasticity of SDH-deficient cells. Later on, Morin *et al.* demonstrated that, in these SDHB-deficient cells, hypermethylation and pseudohypoxia act synergistically leading to the acquisition of metastatic traits (44).

### Rat SDH-deficient RS0 cells

Recently, Powers *et al.* (45) developed two cell lines of SDH-deficient chromaffin cells from rats with a heterozygous germline *SDHB* mutation. Heterozygous *Sdhb*<sup>+/-</sup> founder rats were generated with various deletions in the rat *SDHB* gene: offspring were irradiated and maintained until they were killed. Necropsies were performed and tissue from five PCCs that appeared to be

viable was injected subcutaneously into NOD scid gamma (NSG) mice by using a previously described protocol (46). To generate primary cell cultures, minced xenograft tissue was dissociated in collagenase followed by trypsin (30). The utilisation of xenografts to develop cell lines served to expand tumour cell populations, because the usually small primary tumours did not provide a sufficient number of cells for adequate tests of growth conditions. Two cell lines designated RS1/2 (for *Sdhb* haploinsufficient), and RS0 (for Rat *Sdhb* null) originated from cultures of xenografts. The first RS1/2 cell line had lost the mutated *SDHB* allele but retained the wild type allele. These cells show some succinate accumulation consistent with haploinsufficiency, but they cannot be considered fully SDH-deficient. RS1/2 serves either as a control for studies of RS0 or as a potential tool for studying patients' tumours that may be haploinsufficient for SDH, having undetermined driver mutations. On the contrary, the RS0 cell line is SDH-deficient. The genome, transcriptome and metabolome of RS0 closely resemble those of *SDHB*-mutated human PPGLs, representing the only real cell model deriving from cluster 1 tumours. The features that RS0 xenografts share with SDH-deficient human PPGLs include identical histology, loss of *SDHB* protein with retention of *SDHA*, and expression of the neuroendocrine markers such as TH and chromogranin A. Moreover, in RS0, cytoplasmic vacuoles and sparse, often tiny, secretory granules are present. In contrast, RS1/2 cells have larger secretory granules and lack cytoplasmic vacuoles. Interestingly, RS0 cells grown in a serum-free medium form tumour cell spheroids spontaneously, which are able to grow in suspension.

## Mouse models

For research purposes, animal preclinical models are urgently needed, especially because PCC is a rare condition with a limited availability of malignant tissue. The development and proper characterisation of animal models will determine their usefulness as a model for preclinical trials (47). In the past 30 years, many genetically engineered and allografted mouse models have been generated to investigate the mechanisms of PPGL tumourigenesis and test new therapeutic strategies. Among them, only cluster 2-related predisposed models have been successful. In fact, genetically engineered mice with *SDHB* mutations have failed to develop SDH-deficient PPGLs. Indeed, at present, cluster 1 mouse models, that is, KOs for *SDHx*, *FH*, or *VHL* genes, are not

available. The recent progress facilitated by the TALEN, CRISPR/Cas9 or induced pluripotent stem cell (iPSCs) technologies, will promote, most likely, an acceleration of such models in the near future. Regarding SDH-deficient mouse models, the first homozygous KO of *SDHD* was lethal at an embryonic stage (48). Several years later, a second *SDHD* KO mouse model was generated. Two conditional *SDHD* KO were reported, but none of these models resulted in PPGL development (49). Two different teams published negative results regarding engineering a *SDHB*-deficient mouse model (50, 51). The failure to develop a pure *SDHB* KO model suggests a cellular lethality of *SDHB* complete KO in mice. The generation of a predisposed SDH-related model of PPGLs probably requires a second hit that would allow PGL tumourigenesis in rodents.

However, grafted models are available to study cluster 1-related tumours. In a grafted model, human (xenograft) or animal (xeno-/allograft) cancer cells are transplanted either under the skin (ectopic) or into the organ of tumour origin (orthotopic) using immunodeficient rodents, such as athymic nude mice or severely compromised immunodeficient (SCID) mice. Regarding the engrafted mice, Giubellino *et al.* developed a MTT derivative, known as MTT-luc and an additional cell line expressing green fluorescent protein (GFP) and firefly luciferase obtained by transducing the cells with a pre-packaged lentiviral construct. It is intended to be used for *in vivo* bioluminescence imaging of tumour deposits. *In vivo*, implantation of tumour cells transfected or transduced with the reported gene such as luciferase or GFP allows sequential monitoring of tumour growth within the viscera by measuring these photon signals. This technology is reshaping efficacy evaluations and drug-target algorithms in drug animal testing for several tumour types (52). Indeed, allografts with murine cell lines remain the only feasible option to study, *in vivo*, the behaviour of such tumours.

As described previously, recently Powers and co-workers have developed xenograft models from PCCs that arose in rats with a heterozygous germline mutation in *SDHB*. Two distinct, serially transplantable, PCC xenograft models, which have been designated as RS0 and RS1/2, were derived at 84 and 74 weeks, respectively, from macro PCCs that arose in irradiated rats. RS0 xenografts histologically show sharply defined 'Zellballen' architecture, slightly clear cells and prominent blood vessels closely resembling human PGLs, while RS1/2 shows more diffuse growth. RS0 closely recapitulates the genotype and phenotype of hereditary *SDHB*-mutated

human PPGLs and appears to be a promising model for preclinical studies of these tumours (45).

Lussey-Lepoutre reported a detailed overview of existing rodent models carrying the susceptibility genes involved in human PPGLs and their contribution to the improvement of PCC experimental research (53). Briefly, in both the *c-mos* and MEN2B models, multiple PCCs are associated with thyroid C-cell proliferations, as in human MEN2A and MEN2B. However, some lines of *c-mos* mice exclusively develop PCCs or exclusively C-cell tumours, possibly due to differences in the transgene integration site. Interestingly, *Nf1* heterozygous KO mice, which frequently develop PCCs (28), express high levels of wild type RET (54), suggesting that an increase of the downstream RET activated signalling pathways predispose mice to developing PCC. Indeed, patients harbouring RET gain of function mutations develop MEN 2 syndrome.

PCCs in *Pten* KO mice show consistent deletions in mouse chromosome 4 (55), which is homologous to human chromosome 1p, the most frequent deletion in human PCCs. *ErbB2* and *B-Raf* proto-oncogenes mutated mice are also able to develop PCCs (56, 57), as well as those mutated in *Rb*, *Trp53*, *VHL*, and *Ink4a* tumour suppressor genes (58, 59).

### Which model for which tumour?

For preclinical models, it is important to represent the selected tumour cluster as closely as possible, because each cluster is associated with distinctive signalling pathways and metabolic characteristics. These specific features represent potential drug targets (60, 61, 62) or might be responsible for different responses to drug treatments (63).

The generation of an increasing number of animal and cellular models of PPGL has been seen in the past 30 years. Based on cluster classification, cluster 1-cell models only recently became available, with the development of the RSO cell line. Before that, only cluster 2 cell and mouse models were available. RSO is, in fact, a new xenograft and cell line model of SDH-deficient PCC from rats with a heterozygous germline *SDHB* mutation. The genome, transcriptome and metabolome of this model closely resemble those of *SDHB*-mutated human PPGLs. Nevertheless, further experimental data using this model are needed to understand its importance in preclinical PPGL research. Even if Tischler and colleagues have always done their utmost to characterise their models fully and openly, it should be noted and acknowledged

that MPC, MTT and RSO cell lines all required irradiation to induce tumours in the original animals. This approach introduces unknown elements regarding the translation of any results to the human situation.

About tumour cell spheroids, since necrosis is an adverse factor correlated with risk of metastasis (64), a 3D cell model is a good model for the study of PCC metastasis because of the necrotic core in the middle of the spheroids. On the other hand, the necrotic core can represent a limitation for the study of PCC because it creates an artificial hypoxic environment on the central cells within the spheroid, thereby potentially muddying the distinction between pseudohypoxic cluster 1 and cluster 2. For this reason, monolayers may be equivalent or superior for some types of studies.

Given the fact that PC12 and MPC cells are both chromaffin cells, although they belong to different species, these models are somehow comparable. Morphologically, PC12 and MPC are different in size, shape and show distinct adherence and clumping behaviours, which also vary in response to different culture conditions. In fact, they respond dissimilarly to growth substrates and to nerve growth factor. MPC cells genetically and biochemically resemble human PPGLs. For example, they express substantial levels of the A synthesising enzyme, PNMT, and they do not significantly respond to NGF. In contrast, PC12 are noradrenergic and become excitable following NGF treatment. A surprising finding is that MPC cell lines express high levels of the receptor tyrosine kinase RET, which is involved in the pathogenesis of human PCCs, in hereditary MEN2. They also respond to the RET-activating ligand by exhibiting RET phosphorylation, neurite outgrowth, decreased proliferation, and altered expression of catecholamine biosynthetic enzyme. Consequently, cells undergo neuronal differentiation similar to what has been observed in NGF-treated PC12. Usually, RET is minimally expressed by normal mouse chromaffin cells and its high expression in MPC cells suggests possible relationships between two previously unrelated tumour syndromes, neurofibromatosis and MEN2 (65). Moreover, MPC and their metastatic derivative MTT cells naturally lack *HIF2 $\alpha$* . On the contrary, the rat PC12 cell lines express both *HIF1 $\alpha$*  and *HIF2 $\alpha$*  (13). These data indicate that PC12 cell lines could be utilised to study chromaffin cell or sympathetic neuron biology, while MPC cell lines are a useful model for the study of adrenergic tumours. The major cell line characteristics are summarised in Table 1.

In addition to the fact that the *in vitro* conditions are very different from those found *in vivo*, making correlations among data obtained by different research

**Table 1** Schematic representation of the different cell line characteristics.

Cell	Origin	Genotype and cluster	Catecholamine secretion	Forming clusters/spheroids
PC12	Rat pheochromocytoma	MAX chromosomal rearrangement or translocation, cluster 2	Dopamine, noradrenaline	–
MPC	Mouse pheochromocytoma	Heterozygous NF1 knock out, cluster 2	Noradrenaline, adrenaline	+
MTT	Metastatic mouse pheochromocytoma	Heterozygous NF1 knock out, cluster 2	Noradrenaline, adrenaline	+
hPheo1	Human pheochromocytoma	hTERT immortalised, cluster 2	Not secreting	–
imCC	Mouse	SDHB knock out, cluster 1	Not known	–
RS 1/2	Rat pheochromocytoma	Heterozygous SDHB knock out, cluster 1	As xenograft: dopamine, noradrenaline	+/-
RS0	Rat pheochromocytoma	SDHB knock out, cluster 1	As xenograft: dopamine, adrenaline	+

groups, even if acquired using the same cell lines, is not easy for several reasons. Indeed, the same cell types in different laboratories are often maintained in dissimilar culture conditions, including medium composition, O<sub>2</sub> or CO<sub>2</sub> levels and nutrient concentration (such as diverse percentages of serum or factors added to the medium). Another complication is the passage number. In fact, it is good laboratory practice to work with early cell passages in order to obtain reproducible effects under treatments and avoid the accumulation of possible additional mutations occurring over time. Cryopreservation of early passages, whenever possible, is recommended. As an example, PC12 cells were developed almost 50 years ago and have been maintained in different culture media by some laboratories. Therefore, the ability to induce PNMT may vary according to passage number and the way cultures have been maintained.

Moreover, the appropriate cell of origin is of primary importance (66). Another variable to be considered when working with cell lines is that modifying the gene expression pattern, for example with the silencing of MTT cells with shRNA anti-SDHB, leads to combining the characteristics of the different clusters muddying the distinction between cluster 1 and cluster 2. However, at the moment the cell models described are the only ones available and they allow us to study metabolic functions of PCC/PGL. Nevertheless, interesting future perspectives consist in reintroducing in RS0 cells, by transfection, the wt *SDHB* subunit, in order to clarify the metabolic features and functions of intracellular pathways in the tumourigenesis processes.

## Conclusions

Tumours do not exist as a homogeneous population of malignant cells. Rather a tumour is characterised by a

changing microenvironment, resulting from an interplay between a heterogeneous population of malignant cells and their assorted support of various tumour-associated cells, including macrophages, fibroblasts, pericytes, endothelial and immune cells. The multitude reciprocal interactions between tumour cells and the tumour microenvironment (TME) allow minor populations of tumour cells to evade apoptosis and to develop resistance. Moreover, the TME is rich in cytokines and growth factors, which are secreted by either tumour cells or stromal cells and contribute to aberrant growth, angiogenesis, metastasis and drug resistance.

Given these considerations, the importance of studying and targeting the microenvironment as a possible anti-tumour therapy is increasingly clear. Notably, MPC or MTT spheroid stability provides an opportunity for a future generation of multicellular spheroids consisting of PCC cells, endothelial cells and/or fibroblasts that provide a closer model to the *in vivo* situation within the tumour microenvironment. Despite several attempts, in contrast with MPC and MTT, PC12 cells are unable to form spheroids, even using different methods for spheroid generation (37). In addition, the new isolated RS0 cells, in serum-free medium, spontaneously form spheroid-like masses. However, the possibility of handling these spheres and their stability in culture must be further evaluated. Besides spheroids, a future perspective for PCC studies includes the possibility of developing organoids, which better resemble the human adrenal gland, and that could provide the opportunity to understand the role of the adrenal cortex in modulating PCC onset.

With regard to imCC and hPheo1 cells, although they seem to be promising models, all the data present in the literature are limited to those coming from the research groups that have isolated the two cell lines. In this case,

further studies are necessary to make a proper comparison with the other cell lines.

About the *in vivo* models, knock-in and knock-out mice are useful models to investigate the pathogenesis of human tumours including their metastatic potential, as well as subcutaneous or tail vein injection of tumour cells (allograft or xenograft) (67). The development of these models is particularly challenging given the diverse causes and manifestations of tumours.

Considering the dissimilarities among the different models, it is still necessary to be very careful in drawing general conclusions from the results obtained from any single cell line.

#### Declaration of interest

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