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Thalamic afferents emphasize the different functions of macaque precuneate areas

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Title: **THALAMIC AFFERENTS EMPHASIZE THE DIFFERENT FUNCTIONS OF
MACAQUE PRECUNEATE AREAS**

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Running title (max 40 char.): Thalamic input to the macaque precuneus

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Abstract

We studied the thalamic afferents to cortical areas in the precuneus using injections of retrograde fluorescent neuronal tracers in four male macaques (*Macaca fascicularis*). Six injections were within the limits of cytoarchitectural area PGm, one in area 31 and one in area PEci. Precuneate areas shared strong input from the posterior thalamus (lateral posterior nucleus and pulvinar complex) and moderate input from the medial, lateral and intralaminar thalamic regions. Area PGm received strong connections from the subdivisions of the pulvinar linked to association and visual function (the medial and lateral nuclei), whereas areas 31 and PEci received afferents from the oral division of the pulvinar. All 3 cytoarchitectural areas also received input from subdivisions of the lateral thalamus linked to motor function (ventral lateral and ventral anterior nuclei), with area PEci receiving additional input from a subdivision linked to somatosensory function (ventral posterior lateral nucleus). Finally, only PGm received substantial limbic association afferents, mainly via the lateral dorsal nucleus. These results indicate that area PGm integrates information from visual association, motor and limbic regions of the thalamus, in line with a hypothesized role in spatial cognition, including navigation. By comparison, dorsal precuneate areas (31 and PEci) are more involved in sensorimotor functions, being akin to adjacent areas of the dorsal parietal cortex.

INTRODUCTION

The term precuneate gyrus (or precuneus) refers to the region of cortex located on the medial surface of the parietal lobe, which is anatomically circumscribed by the parieto-occipital, suprasplenial and cingulate sulci. Functionally, the precuneus is delimited caudally by visual and visuomotor association cortex (areas V6 and V6A), dorsally by sensorimotor superior parietal areas (PEc, PEci), rostrally by subdivisions of posterior cingulate cortex (area 23 complex), and ventrally by still poorly characterized subdivisions of putative visual cortex (Passarelli et al. 2018). In terms of cytoarchitecture, the precuneus is similarly organized in humans (Vogt and Laureys 2005; Cavanna and Trimble 2006) and monkeys (Morecraft et al. 2004; Vogt et al. 2005), with most of its volume occupied by cytoarchitectural areas PGm and 31.

In humans, the precuneus has been considered a critical component of the brain systems for numerous aspects of cognitive functions, including participation in the network linked to the neural correlates of self-consciousness (Cavanna and Trimble 2006), and has been reported to be particularly affected in cognitive decline linked to aging and dementia (Buckner et al. 2000, 2005; Matsuda 2001; Lustig et al. 2003). According to studies in non-human primates (Olson et al. 1996; Ferraina et al. 1997; Thier and Andersen 1998; Sato et al. 2006, 2010) and humans (Wenderoth et al. 2005; Baumann et al. 2012), the caudal medial cortex, including the precuneate region, emerges as a site where multiple sensorimotor systems interact to guide complex spatial behaviours such as gaze and limb orientation and navigation. Anatomically, there is evidence showing that, in monkeys, the precuneus forms extensive connections with superior and inferior parietal, cingulate and retrosplenial areas of the cortex, whereas connections with the frontal lobe are somewhat de-emphasized when compared to adjacent superior parietal areas (Pandya and Seltzer 1982; Cavada and Goldman-Rakic 1989a, b; Leichnetz 2001; Parvizi et al. 2006; Passarelli et al. 2018). In this aspect, the precuneus appears to be a node, or intermediary structure, in the parieto-medial temporal pathway with access to the hippocampal and parahippocampal formations, suggesting a role in

whole-body movements and spatial memory (Kravitz et al. 2011).

Recent neuroimaging studies in healthy humans have identified functional interactions between the precuneus and the thalamus (Tomasi and Volkow 2011; Zhang and Li 2012; Cunningham et al. 2017) and described changes in the thalamic-precuneate system in disorders of consciousness (Soddu et al. 2010; Fernández-Espejo et al. 2012). Previous studies in macaques (Yeterian and Pandya 1985, 1988; Schmammann and Pandya 1990; Leichnetz 2001) have reported diffuse projections from nearly all regions of the thalamus to the precuneus but these reports were generally based on few cases and descriptions were largely qualitative. Here, we re-examined in more detail the thalamic afferents to the precuneate gyrus in macaques based on a set of localized retrograde tracer injections. These data add to a recent body of quantitative descriptions of the subcortical input to the caudomedial lobe (Buckwalter et al. 2008; Gamberini et al. 2016; Impieri et al. 2018) and allow us to look for specific functional trends within this large cortical region.

MATERIALS AND METHODS

Fluorescent tracers were injected in the precuneate cortex of four adult monkeys (*Macaca fascicularis*, 2.5 – 4.9 kg) (Fig. 1). Details of the location of injections and tracers used are presented in Table 1. Experimental protocols conformed to the guidelines of the Australian Code of Practice for the Care and Use of Animals for Scientific Purposes and the European Union Directive 86/609/EEC and the revised Directive 2010/63/EU. The Monash University Animal Experimentation Ethics Committee and the University of Bologna Bioethics Committee approved the experimental procedures (University of Bologna Permit N° 170/2015-PR, 19/03/2015; Monash University Permits N° MARP2012/082, SOBSA/P/2010/05, SOBSA/P/2010/28).

Table 1 near here

Fig. 1 near here

Surgical procedures

A detailed description of experimental procedures is available in a previous publication, which included data on cortical connections from many of the same animals (Passarelli et al. 2018). Surgeries were performed under aseptic conditions and full anaesthesia, with the heart rate, blood pressure, respiratory depth and body temperature monitored continuously. The protocol for anaesthesia varied between animals, reflecting procedures approved by the two Universities as well as refinements introduced during the project. Animal A9 (injection cases 1 and 2; see Table 1) was pre-treated with injections of atropine (0.04 mg/kg, i.m.) and ketamine hydrochloride (15 mg/kg, i.m.) and, after 30 min, anesthetized with sodium thiopental (8 mg/kg, i.v.), with additional doses administered as required. Animals MF3 and MF4 (injection cases 3, 5, 6, and 8) were pre-medicated with i.m. injections of diazepam (3.0 mg/kg) and atropine (0.2 mg/kg), and anaesthetized 30 min later with alfaxalone (10 mg/kg, i.m.), with supplemental doses (5 mg/kg) administered i.v. as required. Finally, animal MF6 (injection cases 4 and 7) was pre-medicated with i.m. injections of diazepam (1.0 mg/kg) and atropine (0.04 mg/kg), and anesthetised 30 min later with an i.m.

injection of ketamine (2 mg/kg), medetomidine (0.02 mg/kg) and butorphanol (0.1 mg/kg), after which the animal was intubated and maintained with isoflurane (0.5–2%). In all cases hydration was provided by constant i.v. infusion of Hartmann's solution. Dexamethasone (0.3 mg/kg, i.m.) and Norocillin (25 mg/kg, i.m.) were also administered at the start of the procedures. The animals were placed in a stereotaxic frame and a craniotomy was performed over the posterior parietal cortex. In order to reach the mesial surface of the hemisphere, a portion of the posterior parietal cortex of the contralateral hemisphere (which likely included parts of areas PE, PEc, V6A and/or PGm) was retracted or removed by aspiration, the falx cerebri was retracted, and the mesial surface exposed. Injection sites were selected by direct visualization relative to the cingulate and parieto-occipital sulci. The fluorescent tracers were typically injected using a microsyringe that had a glass micropipette attached to its needle. In 2 cases (Table 1), the retrograde fluorescent tracers Fast blue (FB) and Diamidino Yellow (DY) were directly applied into the cortex as crystals (approximately 100-200 μm in diameter), with the aid of blunt tungsten wires (Rosa et al. 2005). Analgesics (ketorolac, 1 mg/kg, i.m., Temgesic, 0.01 mg/kg, i.m. or Carprofen, 4-5 mg/kg, s.c.) were provided postoperatively.

The day after the surgery, the animals had some difficulty in performing reaching and grasping movements with the hand contralateral to the lesion (for example, to get a piece of fruit) and were somewhat slow in putting the opposite foot on the ground; eating and drinking behaviour were as before surgery and the animals did not look in pain or otherwise distressed. Impairments are overall in agreement with the functional properties of the lesioned areas (Battaglini et al. 2002; Fattori et al. 2017; Gamberini et al. 2018). The motor-related symptoms disappeared a few days after surgery.

Tissue Processing

After a survival period of 14 days, the animals were anesthetized (as above, for different cases) and administered a lethal i.v. barbiturate injection. They were then perfused with several litres of

heparinised saline, followed by 4% paraformaldehyde in 0.1M phosphate buffer at pH 7.4 (followed by 5% buffered glycerol in animal A9). The brains were cryoprotected by immersion in graded series of buffered sucrose (10–30%, most animals) or glycerol (10–20%, animal A9). Coronal sections (50 μm in most cases, 60 μm in A9) were obtained using a freezing microtome. One in five sections was left unstained for observation under a fluorescence microscope. Adjacent sections were stained for Nissl substance, myelin (Gallyas 1979) and cytochrome oxidase (Wong-Riley 1979), and a fifth series was left in reserve. All sections were coverslipped with DPX mounting medium after rapid dehydration in ethanol and clearing with xylene.

Data Analysis

The unstained sections of the hemispheres ipsilateral to the injection sites were scanned for labelled neurons with microscopes under fluorescence illumination (Zeiss Axioscope 2 Plus). For all sections examined, the outer and inner boundaries of the cerebral cortex, the outlines of the injection sites, and the location of labelled cells were mapped at 500-600 μm intervals (1 in 10 sections) using a digitizing system attached to the microscopes (MD3 digitizer and MDPlot software, Accustage). MDplot drawings were aligned with the nearest histological sections in order to assign labelled neurons to thalamic nuclei. The average extent of PGm and locations of injection sites are reported on a reference 3D reconstruction of a macaque brain from the computerized registration of different brains, using the CARET software (<http://www.nitrc.org/projects/caret/>; Van Essen et al. 2001), as described previously (Gamberini et al. 2009; Passarelli et al. 2018).

Assignment of injection sites to cortical areas PGm, PEci and 31 was based on previously described cyto- and myeloarchitectural features (Fig. 2; Pandya and Seltzer 1982; Morecraft et al. 2004; Passarelli et al. 2018). Identification of the thalamic nuclei (Fig. 2) was based on the atlas of Olszewski (1952) and more recent refinements summarized in Mai and Forutan (2012). For identification of the pulvinal subdivisions and the lateral posterior nucleus, which contained the

bulk of labelled cells in our study, the descriptions by Schmammann and Pandya (1990) and Romanski et al. (1997) were useful. Thus, in the rostral pole of the pulvinar, the oral subdivision is characterized by a low cell density. Caudally, the medial and lateral subdivisions are typically present at the same levels; the medial subdivision can be distinguished by a comparatively high cell density and the lateral pulvinar by the numerous fibre bands crossing through it. The lateral posterior nucleus occupies a dorsal and lateral position relative to the medial pulvinar and can be identified by its moderate-sized cells and darker staining. We noted small variations in the boundaries of the different thalamic nuclei across hemispheres, as reported before (Schmammann and Pandya 1990). Table 2 outlines the terminology followed in the present work.

Because the number of labelled neurons depends on many factors, including the type and volume of the injected tracer (e.g., results of our statistical analysis in the following section), **the laminar placement of the injection sites (Jones 2001)**, the distribution of labelling in each nucleus was expressed as a fraction of the total number of recorded neurons in thalamus, case by case, as reported in previous papers (Buckwalter et al. 2008; Gamberini et al. 2016).

Table 2 near here

Fig. 2 near here

RESULTS

We describe the thalamic afferents to the macaque precuneate cortex after 8 retrograde tracer injections. As shown in Figure 1, areas 31 and PEci each received one injection, and area PGm 6 injections. Three of the PGm injections were placed in the dorsorostral part of this area (cases 3-5), and 3 in the ventrocaudal part (cases 6-8), defined according to our recent study (Passarelli et al. 2018). As detailed in this study, the ventrocaudal part of PGm (which has not been sampled in earlier studies) differs from the dorsorostral part in terms of cortico-cortical connections.

The number of labelled neurons in the cortex and thalamus varied between cases (Table 1), likely reflecting both differences in tracer sensitivity and in the extent of injections (Fig. 1); in particular, our analysis showed that the number of observed labelled cells is related to the size of the injection site (Fig. 1, inset c; Pearson's $r = 0.757$, $p < 0.05$). In agreement with previous reports on the thalamocortical connectivity of posterior parietal areas (Gamberini et al. 2016; Impieri et al. 2018), the number of labelled neurons in the thalamus represented a small fraction of the extrinsic (ipsilateral cortical and thalamic) afferents to each area of the precuneate cortex (area PEci: 1.9%; area 31: 2.1%; dorsorostral PGm: $2.8 \pm 2.6\%$; ventrocaudal PGm: $1.6 \pm 1.4\%$).

Areas PGm, 31 and PEci all received substantial input from the associative nuclei of the posterior thalamus and moderate inputs from medial and lateral regions (Mai and Forutan 2012).

Nevertheless, each of the studied areas could be distinguished on the basis of receiving unique connectivity patterns from specific nuclei, as well as the spatial arrangement of labelled cells, as described in the following sections.

Fig. 3 near here

Table 3

Thalamic afferents to areas PEci and 31

Figure 3 illustrates the distribution of labelled cells in the thalamus after injections in areas 31 and PEci in animal A9 (cases 1 and 2, respectively). Results from these injections were similar, with labelled cells mainly concentrated in dorsal portions of the central-posterior thalamus, although differences in their spatial distribution were often observed within the labelled nuclei (see Fig. 3D-G). The proportions of labelled cells in the various nuclei are reported in Table 3. The majority of projecting cells were located in the pulvinar complex, in particular in the oral subdivision (Pul.o), and the lateral posterior nucleus (LP). Afferents from the posterior thalamic nuclei accounted for >50% of the total thalamic input in both cases. Projection cells in LP were found mainly in the ventral half of the nucleus, with cells labelled by the area 31 injection occupying a dorsomedial location relative to those labelled by the PEci injection (Fig. 3D-G). Moderate afferents to both areas arrived from the mediodorsal nucleus (Fig. 3C-G), the ventrolateral nucleus (mainly from the postrema part, VLps, Fig. 3D, E) and the intralaminar region (in particular, the anterior nucleus CL, Fig. 3E). Despite these shared afferents, areas 31 and PEci could also be distinguished by differential input. Thus, the PEci injection labelled many cells in the somatosensory thalamus (ventral posterior lateral nucleus, caudal part, VPLc, Fig. 3D-G; VPL in Table 3), while only sparse cells were found after the area 31 injection. Conversely, the injection in area 31 revealed overall stronger afferents from the intralaminar region (~14% of total thalamic projections, Table 3) with the inclusion of small projections from the paracentral (Pcn) and suprageniculate (SG) nuclei and additional minor afferents from the anterior (AV) and periventricular (Re, Cdc, Cif) nuclei (Fig. 3A, B), as well as from the ventral anterior (VA) nucleus and the anterior (VLo) and medial (VLm) subdivisions of the ventral lateral nucleus (Fig. 3A, VL in Table 3).

Fig. 4 near here

Fig. 5 near here

Thalamic afferents to area PGm

Figures 4-6 show the distribution of thalamic labelling in cases with injections in dorsorostral (Fig.

4, case 4; Fig. 5, cases 3 and 5) and ventrocaudal (Fig. 4, case 7; Fig. 6, cases 6 and 8) parts of PGm. All 6 injections in PGm resulted in similar patterns of labelling in the thalamus, with some variation in the relative density and the origin of projections depending on the location of injection sites (see Table 3 for quantitative analysis).

Similar to the injections in areas 31 and PEci, the LP and the pulvinar complex nuclei contributed the bulk of projections to PGm. Injections in PGm revealed labelled neurons across the medial, lateral and oral subdivisions of the pulvinar (Pul.m, Pul.l, and Pul.o, respectively). However, the numbers of labelled cells in the Pul.o were much lower (on average, <4%) compared to those in the area 31 and PEci cases (>30%). Afferents from the medial pulvinar were numerous (15-31% across cases), and afferents from the lateral division were variable (0.5-33%) and were not detected in injection case 7 (Figs. 4F, 5E-F, 6F, Table 3).

Consistent labelling, albeit consisting of smaller numbers of labelled cells, was found in the mediodorsal (MD) nucleus (1-15%) and the ventrolateral nuclei (6-24%, VLc, VLps in Figs. 4-6). In some cases, afferents originated in the lateral motor VA nucleus; these were somewhat emphasized in two of the cases with ventrocaudal injections (~4%, cases 6 and 8, Table 3). In the anterior region of the thalamus, labelling of variable strength was present in the laterodorsal (LD) nucleus in 5 out of 6 cases. This label was particularly strong (~21%) in case 7 with injection in ventrocaudal PGm. The other nuclei of the anterior complex (AV, AM, AD) contained a few additional labelled cells. Afferents from the intralaminar nuclei (CM/PF, Pcn, Csl, CL, SG, Li) were also present in several cases (Table 3); these tended to be stronger after injections in the dorsorostral part of PGm (on average ~11% in cases 3-5). Finally, the periventricular region contained overall the lowest numbers of labelled cells, with the exception of case 6 with moderate numbers (7.6%), especially in the central latocellular nucleus (Clc).

Fig. 6 near here

General trends in thalamic afferents to the precuneus

Taken together, our results, based on multiple injections in the precuneus and quantitative analyses, indicate that areas PGm, 31 and PEci receive their main input from the posterior thalamus, in different combinations from the lateral posterior and the pulvinar complex, and moderate input from the medial, lateral and intralaminar regions. This information is summarized in Fig. 7. Three main features of the pattern of thalamic input to the precuneus are evident in our data. First, results highlight differential roles for PGm (with strong connectivity with the associative medial and the visual lateral divisions of the pulvinar), and areas 31 and PEci (which receive afferents from the oral division). Second, our injections hint at some degree of topographical organization in the projections from LP and the medial pulvinar, such that more dorsal parts of these nuclei send projections to ventral parts of the precuneus (ventrocaudal PGm: Fig. 6D-F). Conversely, locations in the dorsal precuneus are targeted by central/ventral parts of LP (PEci/31: Fig. 3D-G, dorsorostral PGm: Fig. 5D) and the medial pulvinar (dorsorostral PGm: Fig. 5E). A third feature relates to the lateral group of thalamic nuclei, which contained consistent labelling after injections in all 3 areas. Mai and Forutan (2012) proposed the subdivision of the lateral region into a “motor” and “sensory” domain based on relevant functional properties. In line with this proposal, areas PGm and 31 showed modest labelling (<10% of total labelled cells) in the “motor” thalamus, including the VA and VL nuclei, while numbers of labelled cells after injection in area PEci were balanced in the “sensory” (VPL) and “motor” domain (approximately 8% in each).

Fig. 7 near here

Comparison of precuneate cortex with neighbouring areas

In the final part of our analysis, we compare the thalamic afferents to areas PGm, 31 and PEci with those to neighbouring cortical areas (Fig. 1) using published quantitative data sets. Our analysis

included extrastriate area V6, posterior parietal areas V6A (Gamberini et al. 2016) and PEc (Impieri et al. 2018), and subdivisions of area 23 (Buckwalter et al. 2008). Figure 8 presents a graphic overview of the patterns of thalamic projections to the caudal medial lobe. On the basis of connections with the posterior thalamus, it is possible to identify two main patterns of connectivity oriented as oblique bands, with each of the areas forming the caudomedial lobe retaining distinct connectional features.

A "posterior" connectional pattern is evident for areas V6A, PEc, and PGM. This group shares strong projections from the posterior thalamus with emphasis on projections from the medial pulvinar. Projections from the lateral posterior nucleus become progressively weaker in a rostral direction (in areas PEc and PGM). Additional common projections arrive from the lateral and intralaminar thalamus but these are generally moderate/weak. An "anterior" pattern consists of the posterior cingulate areas (PEci, 31, 23c). Input from the posterior thalamus is still dense but, different to the posterior parietal pattern, projections from the oral pulvinar become evident, emphasizing an associative skeletomotor role.

Lastly, two ventral areas, V6 and 23a/b, are at the limits of these patterns. Area V6 in the ventral-caudal part of the medial lobe is unique in its emphasis on the visual modality with inputs from the lateral and inferior subdivisions of the pulvinar and the lateral geniculate (Gamberini et al. 2016). A second strong contingent of afferents to V6 is noted from the intralaminar (Pcn) and periventricular (Cdc) region; input from the lateral posterior, mediodorsal and the ventral region, typical for posterior parietal areas, is still present but in much lower percentages. Area 23a/b, in the ventrorostral pole of the caudal medial cortex, shows the strongest input from the intralaminar nuclei, followed by inputs from the anterior region, the mediodorsal nucleus and the medial subdivision of pulvinar.

Fig. 8 near here

DISCUSSION

The main conclusion of this tracing study is that the macaque area PGm, defined histologically, integrates information from a set of visual, association, motor and limbic regions of the thalamus. The general lack of input from thalamic structures associated with somatomotor functions (such as the ventral posterior complex and the oral pulvinar; Grieve et al. 2000; Padberg et al. 2009) differentiates PGm from the adjacent superior parietal areas, including PEci and 31, specialized in analyses related to the use of the limbs.

The projections from multiple thalamic regions parallel the rich cortical connectivity of PGm with visual, parietal association, dorsolateral prefrontal and limbic systems (Pandya and Seltzer 1982; Cavada and Goldman-Rakic 1989a, b; Leichnetz 2001; Passarelli et al. 2018). Our earlier study on the cortical connectivity of the area (Passarelli et al. 2018) showed evidence of anatomical heterogeneity: pathways from premotor cortex preferentially targeted dorsorostral PGm, whereas ventrocaudal PGm received preferential connections from extrastriate and posterior cingulate/retrosplenial areas. Similar trends were not obvious in the organization of thalamocortical projections (Table 3 and Fig. 7): dorsorostral PGm showed somewhat denser input from the intralaminar region whereas ventrocaudal PGm received stronger projections from the lateral motor (VL/VA) and anterior limbic nuclei (mainly LD and AV), but differences were generally subtle. In this regard, previous findings rather support our observation here that the pattern of subcortical connectivity is often a relatively poor predictor of architectonic variation and/or regional specialization (Boussaoud et al., 1992; Buckwalter et al, 2008; Gamberini et al., 2016). We did, however, observe variability in some of the thalamic projections to PGm; for example, projections from the limbic lateral dorsal nucleus, the ventral lateral nucleus and the lateral pulvinar targeted strongly specific parts of PGm (Table 3), hinting at a complex pattern of organization that cannot be fully captured by traditional notions of homogeneous cortical and thalamic connections throughout

the borders of this architectural field (Passarelli et al. 2018).

Origins of major thalamic afferents to area PGm

The connections of PGm with the thalamus have been previously studied in macaques using anterograde and retrograde tracer injections in the cortex (Schmahmann and Pandya 1990; Leichnetz 2001; Parvizi et al. 2006; Buckwalter et al. 2008), but those studies generally targeted dorsal portions of this cytoarchitectural area, and robust quantitative data were reported only in one paper (Buckwalter et al. 2008). Despite differences in the type of tracers (retro- or anterograde) or concentrations of fluorescent tracers (e.g., 3-4% for FB and DY in Buckwalter et al. 2008), present and earlier quantitative analyses converge in showing that the most substantial thalamic input to PGm (on average ~60% of total thalamic input) originates in the lateral posterior nucleus and the pulvinar. While most studies include projections from the oral, lateral and medial divisions of the pulvinar, the large, bi-directional tracing study by Leichnetz (2001) only reported connections with the lateral and medial divisions.

The pulvinar complex and the lateral posterior nucleus constitute the numerically predominant projections of the thalamus to the parietal lobe (present findings; Cappe et al., 2007; Gamberini et al., 2016; Impieri et al., 2018; Schmahmann and Pandya, 1990), and thalamic connections likely reflect parietal specializations in cognition and in the planning and control of motor actions (Battaglia-Mayer et al. 2001; Andersen et al. 2014; Hadjimitsakis et al. 2015, 2019; Galletti and Fattori 2018). The LP, as an associative nucleus, provides information on somatic and attentional stimuli useful in the control of movements (Kamishina et al. 2008, 2009; Gamberini et al. 2016). A modulatory effect on these functions could be attained via the rich dopamine innervation of the monkey and human LP (García-Cabezas et al. 2007). The border between the LP and the pulvinar nuclei has been difficult to establish in neurophysiological studies (Acuña et al. 1986, 1990; Cudeiro et al. 1989), and the two are often considered as a functional unit. The medial pulvinar,

much like LP, is viewed as a multimodal associative nucleus: it has extensive connectivity with high-order cortical systems (but not with early sensory areas, Homman-Ludiye and Bourne, 2019), it forms the subcortical component of the attentional network of the brain (Shipp 2003; Saalman et al. 2012) and is involved in visuospatial processing and reward-based decision making (Grieve et al. 2000; Wilke et al. 2013).

In addition to the strong input from the medial division, consistent (and, occasionally, of comparable strength) input to PGm was observed from the lateral pulvinar (on average ~10% of total, Table 3), which has connections with extrastriate and ventral stream areas (Gamberini et al., 2016; Kaas and Lyon, 2007). Projection neurons to PGm were restricted to the dorsal part of this division, similar to what has been reported for the inferior parietal areas (Baleydier and Morel 1992; Hardy and Lynch 1992). This territory overlaps at least partially the field P4, which forms connections with peripheral representations of V2 and V4 (Ungerleider et al. 2014; Gattass et al. 2014). The same dorsal territory of the lateral pulvinar could also be coextensive with the field Pdm (Petersen et al. 1985) where neurons show poor retinotopic organization, but responses are enhanced for stimuli that become the focus of the animal's attention.

Minor thalamic afferents to area PGm

Secondary afferents to PGm arrived from several other thalamic nuclei, notably the VL and VA nuclei (on average, ~12% of total), the MD nucleus and the anterior (superior) and intralaminar nuclei (~ 7% of total projections from each thalamic region). This distribution is in agreement with previous studies (Buckwalter et al., 2008; Schmammann and Pandya, 1990; Yeterian and Pandya, 1985), with small differences noted; for example, in the presence of labelled cells in the LD nucleus.

The ventral nuclei and the mediodorsal nucleus receive input from the cerebellum and the basal ganglia, and send their main output to the frontal cortex (Alexander et al., 1991; Asanuma et al.,

1983; Kultas-Ilinsky et al., 2003; McFarland and Haber, 2002; Mitchell, 2015). Through posterior parts of the VL nucleus (the postrema, VLps, and caudal, VLc, divisions) PGM appears to receive primarily cerebellar influences (Asanuma et al. 1983; Stepniewska et al. 2003) or a balance of cerebellar-pallidal influences, according to other studies (Rouiller et al. 1994). The dorsal-posterior part of VL is the dominant source of input to medial and caudal parietal areas (Schmahmann and Pandya 1990; Gamberini et al. 2016; Impieri et al. 2018); the parietal projection zone is much restricted compared to the wider territory of the lateral thalamic region that sends projections to motor and premotor areas (Matelli and Luppino 1996; Morel et al. 2005; Burman et al. 2014b, a). In addition, the role of VLc/VLps in motor behaviour is unclear given that the majority of responses in this region occur after passive somatosensory stimulation (Vitek et al. 1994) and microstimulation of the dorsal VL has little effect in producing movement (Vitek et al. 1996).

The functions of medial, intralaminar and limbic connections of PGM are less clear. Prevalent theories suggest they might be associated with the arousal and visual awareness systems (Purpura and Schiff 1997), with spatial learning and memory (Aggleton and Nelson 2015) and the temporal propagation of cortical activity (Pergola et al. 2018). Several aspects of these behaviours might be mediated by eye movements, as suggested by the presence of neurons in the medial and central thalamus that show oculomotor, fixation and eye position activations (Schlag and Schlag-Rey 1984; Schlag-Rey and Schlag 1984; Watanabe and Funahashi 2004; Tanaka 2005).

Conclusions

Several lines of evidence in humans, including results from resting-state functional connectivity and morphological analyses (Cauda et al. 2010; Pereira-Pedro and Bruner 2016; Bruner et al. 2017) as well as comparative imaging studies between humans and macaques (Margulies et al. 2009) suggest that the caudomedial lobe contains several functional modules, although the specifics of the extent and roles for each functional entity remain to be fully characterized (Cauda et al. 2010).

Architectural studies in macaque monkeys (Morecraft et al. 2004; Luppino et al. 2005) have described a series of areas in the caudomedial lobe and their corticocortical connections (Cavada and Goldman-Rakic 1989b, a; Parvizi et al. 2006; Gamberini et al. 2016; Passarelli et al. 2018). Our present results and previous findings (Schmahmann and Pandya 1990; Buckwalter et al. 2008; Gamberini et al. 2016; Impieri et al. 2018) show that although the caudomedial lobe is a major site of converging subcortical projections from the posterior thalamus, there is specificity in the projections from the various subdivisions of the pulvinar. Thus, dorsorostral parts of medial parietal cortex (areas 31 and PEci) are targeted by the oral pulvinar; the caudal medial lobe, including the core of the precuneus (PGm), V6A and PEc, much like the caudal inferior parietal cortex and the posterior cingulate cortex, receives substantial projections from the medial pulvinar, in particular from its dorsal part. The origins of projections are different for each region: those directed to parietal cortex stem from lateral and central locations; projections to cingulate cortex also arise from medial locations. The ventral and caudal part of the medial lobe, corresponding to the parieto-occipital area V6, receives its major input from “visual” parts of the pulvinar, i.e., the inferior and lateral subdivisions. In a broad sense, these data conform to a general role for the dorsal caudomedial lobe in the guidance of sensory-spatial behaviour whereas the ventral caudalmost region forms part of the visual networks of the brain. Future studies designed to map the sensory properties of pulvinar populations are necessary to understand in more detail the role of these specific pulvinar-cortical networks in shaping primate and human motor functions and cognition.

ABBREVIATIONS

Thalamic nuclei and regional subdivisions

Superior or Anterior

AD, Anterior Dorsal

AM, Anterior Medial

AV, Anterior Ventral

LD, Lateral Dorsal

Medial

MD, Medial Dorsal

MDdc, Medial Dorsal, densocellular part

MDmc, Medial Dorsal, magnocellular part

MDmc/pc, Medial Dorsal, magnocellular/parvocellular part

MDmf, Medial Dorsal, multiform part

MDpc, Medial Dorsal, parvocellular part

Lateral

VA, Ventral Anterior

VAdc, Ventral Anterior, densocellular part

VAmc, Ventral Anterior, magnocellular part

VApdc, Ventral Anterior, parvocellular/densocellular part

VL, Ventral Lateral

VLc, Ventral Lateral, caudal

VLm, Ventral Lateral, medial

VLo, Ventral Lateral, oral

VLps, Ventral Lateral, postrema

VPI, Ventral Posterior Inferior

VPL, Ventral Posterior Lateral

VPLc, Ventral Posterior Lateral, caudal

VPLo, Ventral Posterior Lateral, oral

VPM, Ventral Posterior Medial

VPMpc, Ventral Posterior Medial, parvocellular

X, area X

Intralaminar

CL, Central Lateral

CM, Centromedian

Csl, Central superior lateral

Li, Limitans

Pcn, Paracentral

PF, Parafascicular

SG, Suprageniculate

Periventricular

Cdc, Central densocellular

Cif, Central inferior

Cim, Central intermediate

Clc, Central latocellular

Cs, Central superior

Pa, Paraventricular

Pt, Paratenial

Re, Reuniens

Posterior

LP, Lateral Posterior

Pul, Pulvinar

Pul.i, Pulvinar, inferior subdivision

Pul.l, Pulvinar, lateral subdivision

Pul.m, Pulvinar, medial subdivision

Pul.o, Pulvinar, oral (anterior) subdivision

GLd, Lateral geniculate body, dorsal division

GMmc, Medial geniculate body, magnocellular division

GMpc, Medial geniculate body, parvocellular division

Other

Hlpc, Lateral habenula, parvocellular division

Hm, Medial habenula

R, Reticular

Sm, Stria medullaris

STN, Subthalamic nucleus

THI, Habenulo-interpedicular tract

Cortical areas

SPL, superior parietal lobule

dr PGm, area PGm, dorsorostral sector

vc PGm, area PGm, ventrocaudal sector

V6, area V6

V6Ad, area V6A, dorsal portion

V6Av, area V6A, ventral portion

PEc, area PEc

PE, area PE

PEci, area PE, cingulate portion

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LEGENDS

Figure 1. Summary of injection site locations.

Top: Medial view of a reconstruction of a macaque left hemisphere showing the location of the injection sites, based on computerized registration of the brains of the different animals relative to the sulcal pattern. Injections in areas 31 (purple), PEci (white), the dorsorostral part of area PGm (cyan) and the ventrocaudal part of area PGm (green) are indicated. The dashed lines indicate the approximate extent of area PGm according to previous studies (Pandya and Seltzer 1982; Cavada and Goldman-Rakic 1989b, a; Morecraft et al. 2004). The pink and yellow shading indicate the revised borders of areas PGm and 31 (Passarelli et al. 2018). White symbols indicate the approximate locations of injection sites in neighbouring areas according to other studies: star, injections in PEc (Impieri et al. 2018); triangles, injections in V6 and V6A (Gamberini et al. 2016); squares, injections in subdivisions of 23 (Buckwalter et al. 2008). For each injection, the core (dark spot) and the halo zone (coloured region around the core) are shown. Dashed lines indicate the average architectonic border of injected (PGm, 31 and PEci) and neighbouring areas. The location of the injection sites in the cortical thickness is shown on single coronal sections. Red lines depict traces of layer 4 and correspond to the boundary between superficial and deep layers.

Bottom: Epifluorescent photographs of coronal sections through the injection sites in 4 cases. Case 1: FB injection of area 31, Case 2: DY injection of area PEci, Case 3: FB injection of the dorsorostral area PGm, Case 8: DY injection of the ventrocaudal area PGm. Scale bar = 1.0 mm.

Insets a and b show examples of FB- and DY-labelled cells (from cases 1 and 2, respectively) in greater magnification. Inset c: the scatter plot shows that the size of the injection site (in mm³) is related to the number of labelled cells in the cortex and thalamus.

Abbreviations: cal, calcarine sulcus; cgs, cingulate sulcus; ips, intraparietal sulcus; pos, parieto-occipital sulcus; D, dorsal; L, lateral; V, ventral; A, anterior.

Figure 2. Cytoarchitecture of injected areas and of thalamic nuclei.

A. Low-power photomicrographs showing the cytoarchitectural criteria used to distinguish areas in the precuneus cortex. Coronal sections were taken at the levels indicated on the inset. Scale bar = 200 μm .

B, C. Low-power photomicrographs of representative coronal sections from animals A9-R (B) and MF6-L (C) showing the cytoarchitectonic subdivisions of thalamic nuclei. Scale bar = 1 mm.

Abbreviations: cc, corpus callosum; M, medial. For the nomenclature of thalamic nuclei see the list of abbreviations.

Figure 3. Thalamic afferents to areas 31 and PEci.

Seven coronal sections (A–G) through the thalamus illustrating the distribution of labelled cells after injections in areas 31 (case 1, black circles) and PEci (case 2, white circles). In the top right of the figure, a medial view of a reconstruction of a macaque left hemisphere showing the location of the injection sites (shown here and in Figures 3-5). The numbers in parentheses, shown here and in Figures 4, 5 and 6, correlate to the approximate coronal level reported by Olszewski (1952). For the nomenclature of thalamic nuclei see the list of abbreviations.

Figure 4. Thalamic afferents to dorsorostral and ventrocaudal sector of PGm.

Six coronal sections from cases 4 and 7 are reported. Black circles represent the labelled cells after injection in the dorsorostral sector of PGm (case 4), and white circles after injection in the ventrocaudal sector of PGm (case 7) in the same hemisphere. For the nomenclature of thalamic nuclei, see the list of abbreviations. Other details and abbreviations as in Figures 1 and 3.

Figure 5. Thalamic afferents to dorsorostral sector of area PGm.

Six coronal sections from cases 3 (black circles) and 5 (white circles) are reported. For the nomenclature of thalamic nuclei, see the list of abbreviations. Other details and abbreviations as in Figures 1 and 3.

Figure 6. Thalamic afferents to ventrocaudal sector of area PGm.

Six coronal sections from cases 6 (black circles) and 8 (white circles) are reported. For the nomenclature of thalamic nuclei, see the list of abbreviations. Other details and abbreviations as in Figures 1 and 3.

Figure 7. Summary of the thalamic afferents after injections in precuneate cortex.

A. Thalamic afferents to areas 31 and PEci in cases 1 and 2, respectively.

B. Average percentages of labelled cells in thalamic nuclei after tracer injections in dorsorostral and ventrocaudal sectors of area PGm. Vertical bar, SD.

Dashed lines represent nuclei where thalamic label was not observed. Gross subdivisions of the thalamus into 6 regions follow the proposal by Mai and Forutan (2012).

For the nomenclature of thalamic nuclei, see list of abbreviations and Table 2.

Figure 8. Patterns of thalamic input to precuneus and neighbouring areas

In A, the "posterior" pattern of connections including areas V6A (population data from Gamberini et al., 2016), PEc (population data from Impieri et al., 2018), and PGm (population data from present study); in B, the "anterior" pattern including areas PEci (case 2 of present study), 31 (case 1 of present study) and 23c (case 4 of Buckwalter et al., 2008). For comparison in C, thalamic afferents after injections in areas V6 (population data from Gamberini et al., 2016) and 23a/b (case 1 of Buckwalter et al., 2008) are represented.

For the nomenclature of thalamic nuclei, see list of abbreviations and Table 2.

Table 1. Injection sites and neuronal tracers employed in the experiments.

Table 2. Correspondence of nomenclature of the thalamic nuclei involved in this study.

Table 3. Summary of the projections in thalamic nuclei after injections in areas 31, PEci and PGm of the medial hemisphere.

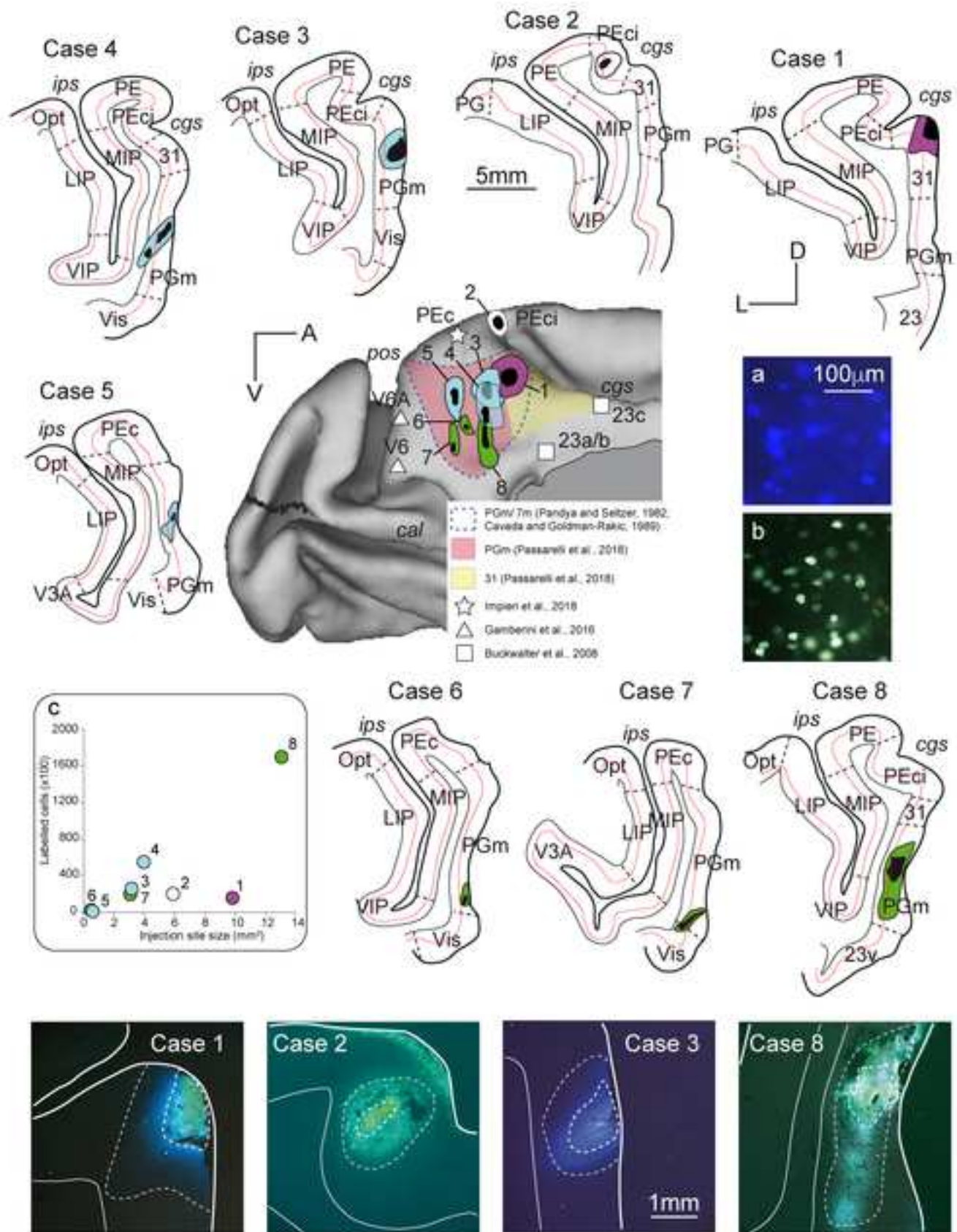


Figure 1_revised

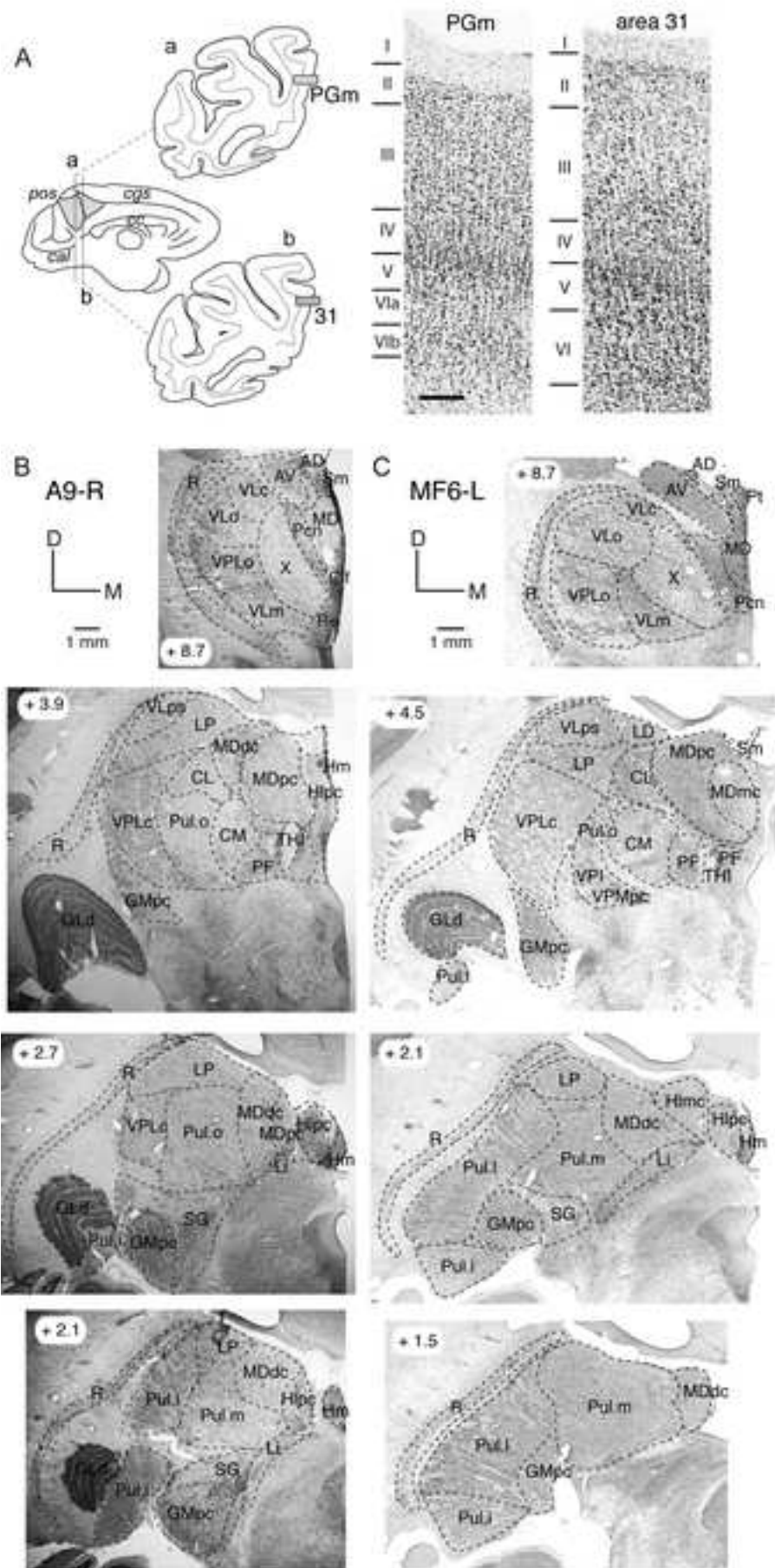


Figure 2_revised

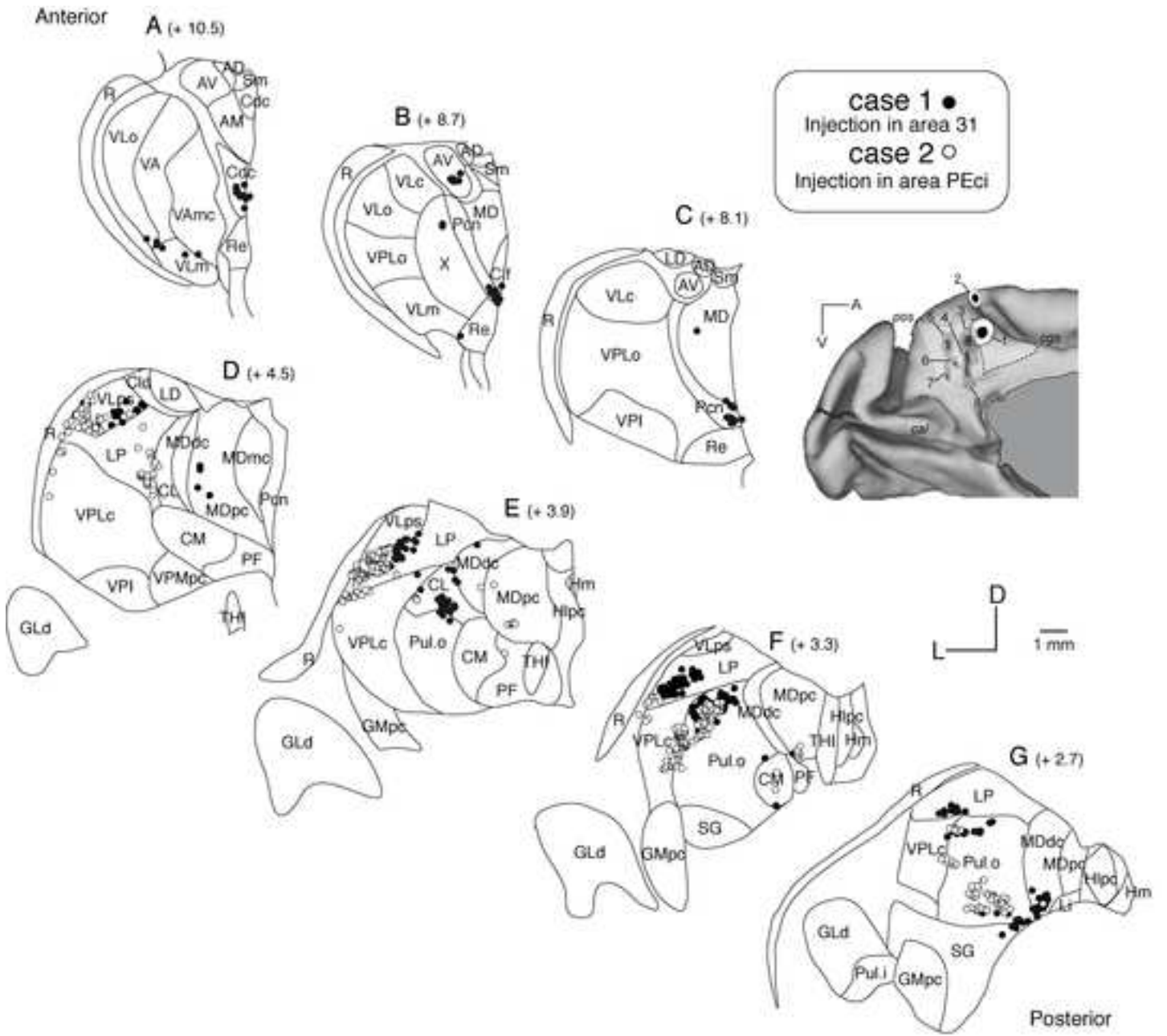


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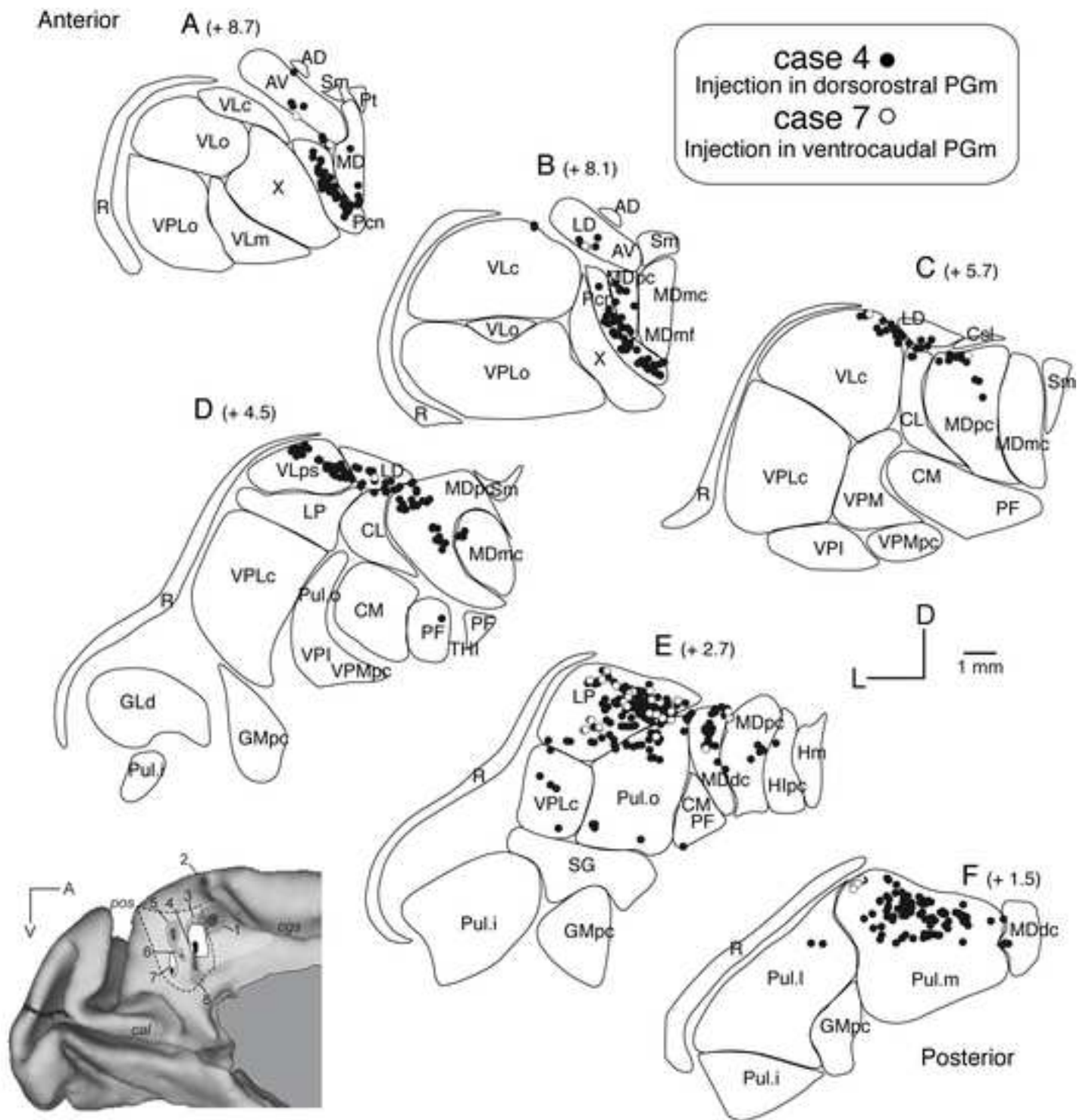


Figure 4_revised

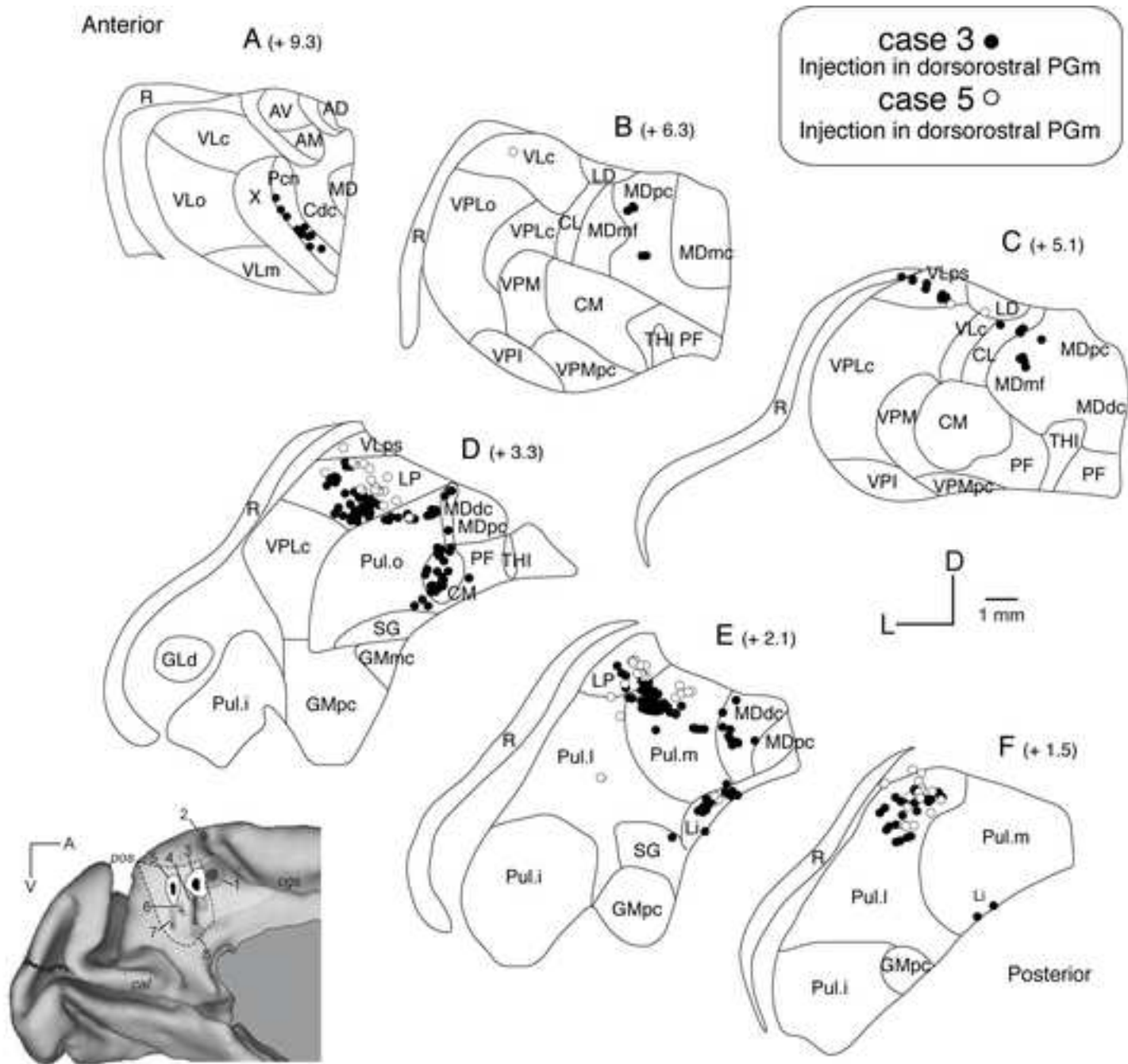


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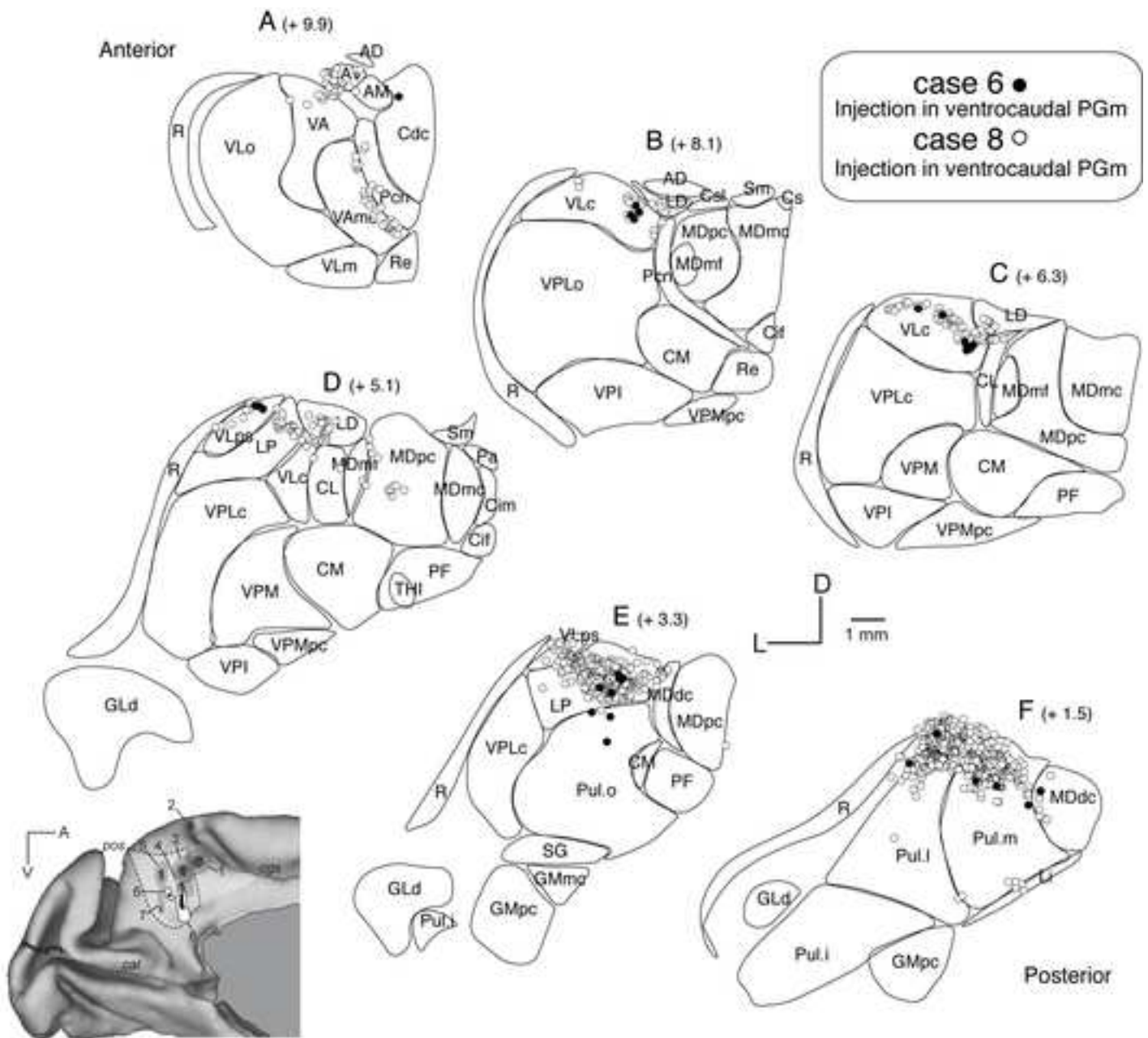


Figure 6_revised

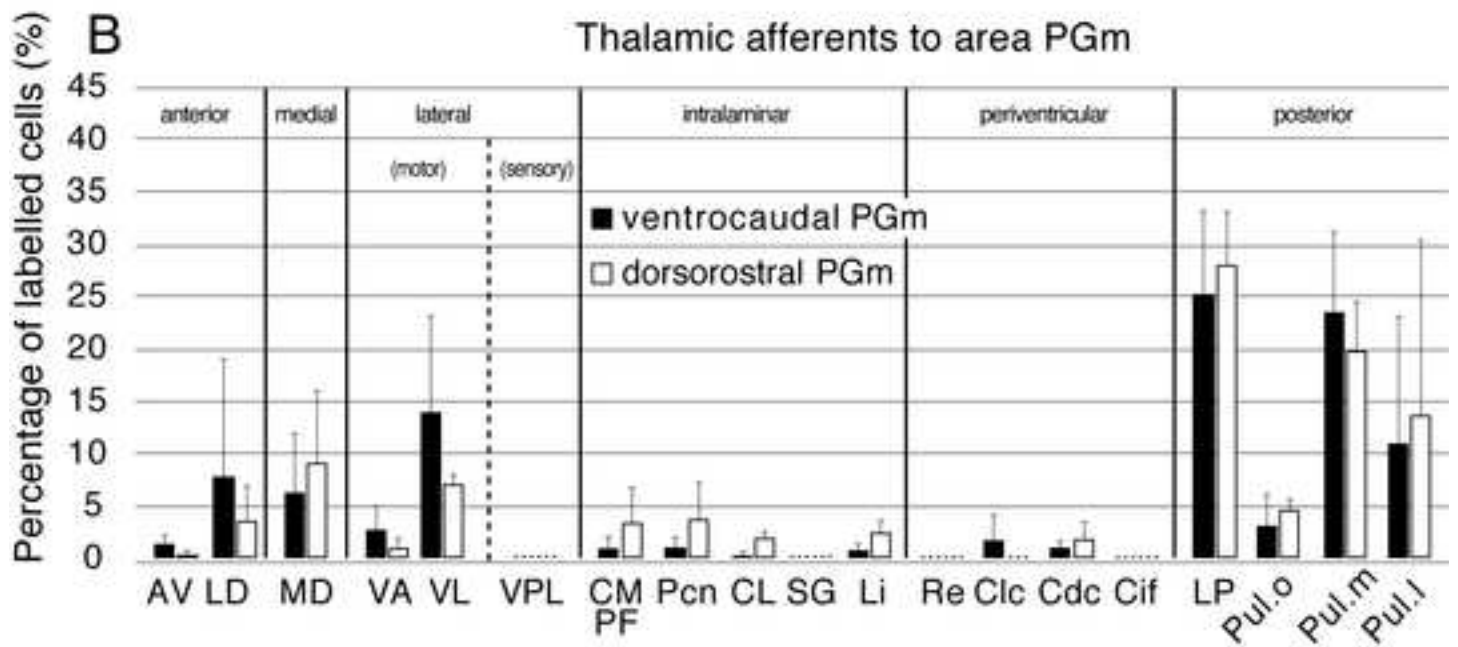
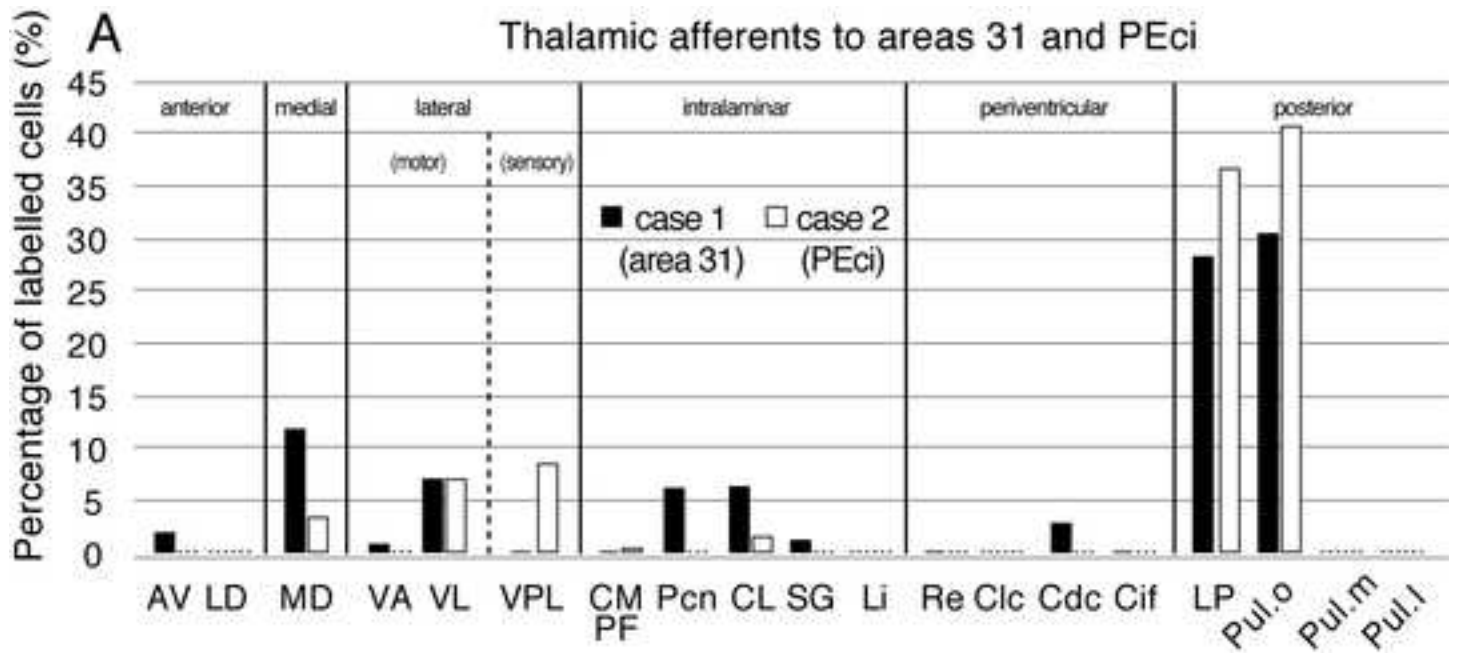


Figure 7_revised

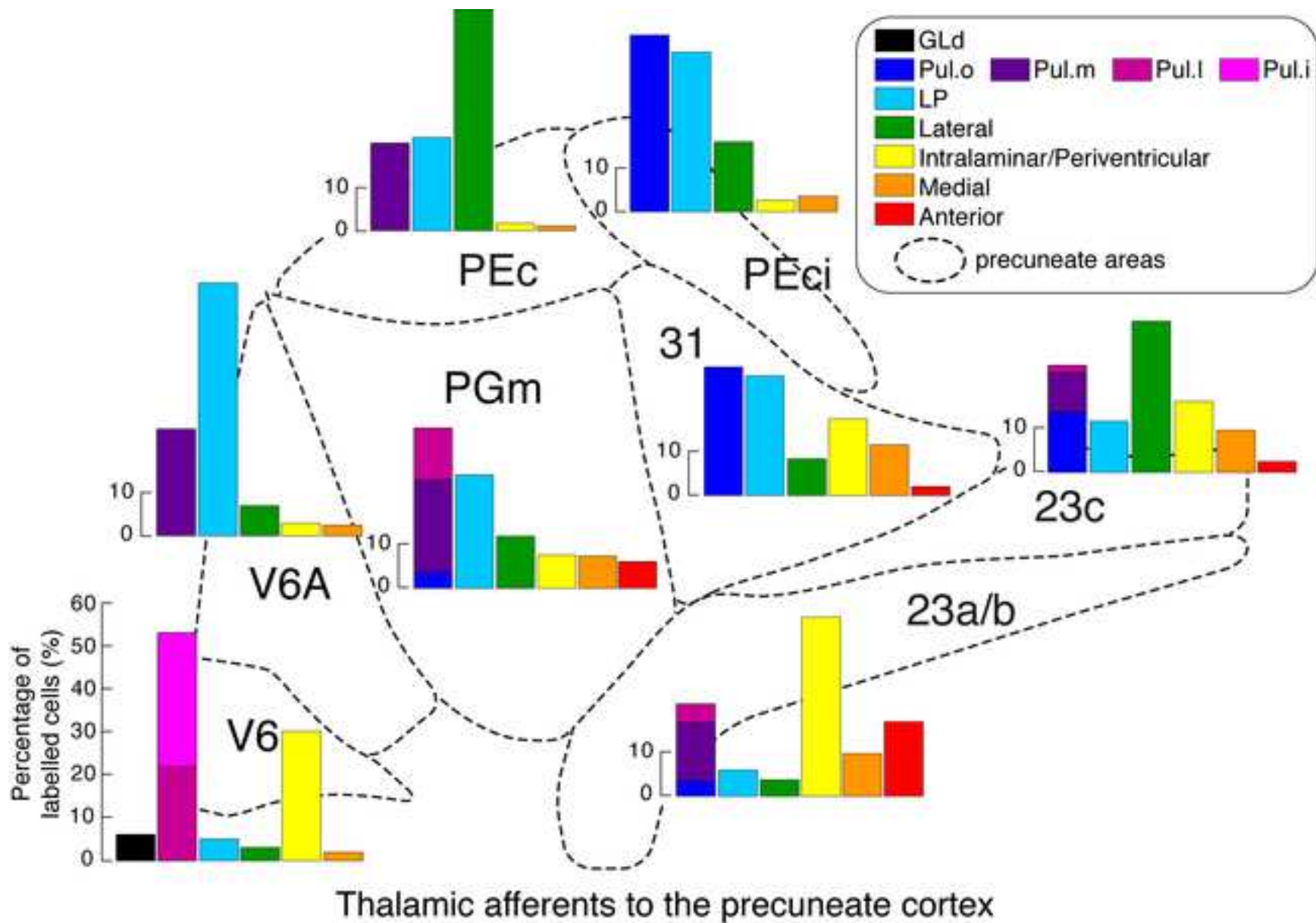


Figure 8_revised

Table 1. Injection sites and neuronal tracers employed in the experiments

Case	Animal-hemisphere ^a	Injected area	Tracer ^b	Amount and concentration of tracer	Number of cortical ^c /thalamic labelled cells
1	A9-R	31	FB	0.2 μ l, 1%	15581/332
2	A9-R	PEci	DY	1 crystal	19949/386
3	MF3-R	dorsorostral PGm	FB	1 crystal	25535/453
4	MF6-L	dorsorostral PGm	FB	0.35 μ l, 1.5%	53398/1742
5	MF3-R	dorsorostral PGm	FR	0.25 μ l, 15%	1281/92
6	MF4-R	ventrocaudal PGm	FR	0.4 μ l, 15%	2496/92
7	MF6-L	ventrocaudal PGm	DY	0.35 μ l, 1.5%	19674/150
8	MF4-R	ventrocaudal PGm	DY	1 μ l, 1.5%	166982/2906

^aR, right; L, left.

^bTracer abbreviations: FB, Fast Blue, Polysciences Europe; DY, Diamidino Yellow, Sigma Aldrich; FR, Fluoro Ruby, Invitrogen – Molecular Probe.

^cExtrinsic ipsilateral connections.

Table 2. Correspondence of nomenclature of the thalamic nuclei involved in this study

Thalamic regions (Mai & Forutan, 2012)	Olszewski (1952)	Present study	Thalamic regions (Mai & Forutan, 2012)	Olszewski (1952)	Present study
Anterior (or superior)	AD	AD	Intralaminar	Cn. Md.	CM
	AV	AV		Pf	PF
	AM	AM		Csl	Csl
	LD	LD		Pcn	Pcn
Medial	MD	MD	Periventricular	Cl	CL
	MDmc	MD		SG	SG
	MDpc	MD		Li	Li
	MDmf	MD		Re	Re
	MDdc	MD		Pt	Pt
Lateral (Motor)	VA	VA	Posterior	Pa	Pa
	VAdc	VA		Cle	Cle
	VAmc	VA		Cdc	Cdc
	VApC	VA		Cif	Cif
	VApC/dc	VA		Cs	Cs
	VLo	VA		LP	LP
	VLc	VL		Pul.o	Pul.o
	VLps	VL		Pul.m	Pul.m
	VLm	VL		Pul.l	Pul.l
	X	VL		Pul.i	Pul.i
	VPLo	VL		GLd	GLd
Lateral (Sensory)	VPLc	VPL	GMmc	GMmc	
	VPM	VPM	GMpc	GMpc	
	VPMpc	VPM			
	VPI	VPI			

Table 3. Summary of the projections in thalamic nuclei after injections in areas 31, PEci and PGm.

Injected area		Area 31	PEci	dorsorostral PGm			ventrocaudal PGm		
Case		1	2	3	4	5	6	7	8
Thalamic regions									
Anterior	AD	-	-	-	-	-	-	-	*
	AV	2.1	-	-	0.7	-	2.2	0.7	0.8
	AM	-	-	-	0.9	-	-	1.3	*
	LD	-	-	0.9	7.2	2.2	-	20.7	2.3
	Total	2.1	0.0	0.9	8.8	2.2	2.2	22.7	3.1
Medial	MD	12.0	3.6	11.3	14.6	1.1	3.3	12.7	2.3
	Total	12.0	3.6	11.3	14.6	1.1	3.3	12.7	2.3
Lateral	VA	1.2	-	1.8	0.9	-	4.3	-	3.5
	VL	7.2	7.3	7.7	6.8	6.5	23.9	12.0	5.6
	VPL	*	8.8	-	*	-	-	-	-
	Total	8.4	16.1	9.5	7.7	6.5	28.2	12.0	9.1
Intralaminar	CM/PF	*	0.8	6.8	2.7	-	2.2	-	*
	Csl	-	-	-	*	1.1	-	0.7	*
	Pcn	6.3	-	4.0	6.8	-	-	2.0	*
	CL	6.6	1.8	2.0	1.0	2.2	-	-	0.6
	SG	1.5	-	*	*	-	-	-	*
	Li	-	-	3.5	1.0	2.2	1.1	-	1.0
	Total	14.4	2.6	16.3	11.5	5.5	3.3	2.7	1.6
Periventricular	Re	*	-	-	-	-	2.2	-	-
	Cle	-	-	-	-	-	4.3	-	*
	Cdc	3.0	-	1.6	3.4	-	1.1	1.3	-
	Cif	*	-	-	-	-	-	-	-
	Total	3.0	0.0	1.6	3.4	0.0	7.6	1.3	*
Posterior	LP	28.3	36.8	21.8	31.6	30.4	16.3	32.0	27.2
	Pul.o	30.4	40.9	5.5	3.6	4.3	6.5	1.3	1.0
	Pul.m	-	-	25.2	16.8	17.4	23.9	15.3	31.0
	Pul.l	-	-	7.5	0.5	32.6	8.7	-	24.0
	Total	58.7	77.7	60.0	52.5	84.7	55.4	48.6	83.2
Other			*	*					
Thalamus total	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	

Relative percentages of labelled cells are listed for each nucleus and for each group of nuclei (in bold). Dash represents no observed labelling. *<0.5% of total projections.