The *Foxb1*-Expressing Neurons of the Ventrolateral Hypothalamic Parvafox Nucleus Project to Defensive Circuits

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ABSTRACT

The parvafox nucleus is an elongated structure that is lodged within the ventrolateral hypothalamus and lies along the optic tract. It comprises axially located parvalbumin (Parv)-positive neurons and a peripheral cuff of *Foxb1*-expressing ones. In the present study, injections of Cre-dependent adenoviral constructs were targeted to the ventrolateral hypothalamus of *Foxb1/Cre* mice to label specifically and map the efferent connections of the *Foxb1*-expressing subpopulation of neurons of the parvafox nucleus. These neurons project more widely than do the Parv-positive ones and implicate a part of the axons known to emanate from the lateral hypothalamus. High labeling densities were found in the dorsolateral and the upper lateral portion of the periaqueductal gray (PAG), the Su3 and PV2 nuclei of the ventrolateral PAG, the cuneiform nucleus, the mesencephalic reticular formation, and the superior colliculus. Intermediate densities of terminals were encountered in the septum, bed nucleus of the stria terminalis, substantia innominata, various thalamic and hypothalamic nuclei, pedunculopontine nucleus, Barrington's nucleus, retrofacial nucleus, and retroambigual nucleus. Scattered terminals were observed in the olfactory bulbs, the prefrontal cortex and the lamina X of the cervical spinal cord. Because the terminals were demonstrated to express the glutamate transporter VGlut2, the projections are presumed to be excitatory. A common denominator of the main target sites of the *Foxb1*-positive axons of the parvafox nucleus appears to be an involvement in the defensive reactions to life-threatening situations. The hypothalamic parvafox nucleus may contribute to the autonomic manifestations that accompany the expression of emotions.

INDEXING TERMS: PAG; anxiety; stress; defensive reaction; aggression; attack area; PV1 nucleus; PV1-Foxb1 nucleus; blood pressure control; respiration; vocalization

The lateral hypothalamus is a large, reticular region of the diencephalon. It is composed of a "confederation of scattered cells" (Peyron et al., 1998), which are involved in several physiological processes, ranging from the control of feeding to that of sleep (Saper, 2004). Interest in this part of the hypothalamus underwent a revival after the discovery therein of abundant scattered neurons expressing the peptide hypocretin/orexin (de Lecea et al., 1998; Sakurai et al., 1998b). Hypocretin/orexin-expressing cells are intermingled with MCH-positive ones (Skofitsch et al., 1985). The two populations play similar roles in the control of food intake and complementary ones in that of the sleep–wake cycle (Barson et al., 2013; de Lecea et al., 1998). The orexigenic function of the two peptides was revealed after intracerebral injections (Sakurai et al., 1998b), and the hypnagogic one after the detection of defective orexin receptors in narcoleptic dogs (Lin et al., 1999). These findings have been confirmed by tracing the axonal projections of the hypocretin/orexin- and MCH-expressing neurons (Peyron et al., 1998; Skofitsch et al., 1985). The excitatory projections

of the hypocretinin/orexin-positive cells to the locus coeruleus (Peyron et al., 1998) highlight their role in arousal and wakefulness (Sutcliffe and de Lecea, 2000). What is remarkable in this unparalleled success story is the unusual way in which the functional roles of the nerve cell aggregates in the lateral hypothalamus were unfolded. In most other cases, the path of discovery begins with the application of classical Nissl-staining techniques, whereby subtle cytoarchitectonic differences are revealed. Later, the connections are mapped using precise stereotactic tracer techniques (see, e.g., Hahn and Swanson, 2015). From these meticulous connectivity studies, the probable functions of individual cytoarchitectonic regions can then be deduced. For example, the connections of the juxtaventromedial region of the lateral hypothalamus point to a prominent role in reproductive behavior (Hahn and Swanson, 2015). Hence, both the "unique" peptide approach and the Nissl-stereotactic tracer injection one have been useful in elucidating the roles of subpopulations of neurons or of subareas in the lateral hypothalamus.

Using a combination of the two approaches, viz. by injecting Cre-dependent tracers into Parv/Cre mice and by monitoring the expression of the nonunique marker parvalbumin (Parv), we have mapped the connections of the Parv-positive string of neurons in the PV1 nucleus (Celio et al., 2013), which is embedded in the medial forebrain bundle (mfb) of the ventrolateral hypothalamus. Later, the existence of a cuff of Foxb1-expressing neurons was revealed around the Parv-expressing ones (Bilella et al., 2014). Taking this circumstance into account, the PV1 nucleus was thenceforth referred to as the parvafox nucleus (Alvarez-Bolado and Celio, 2015). *Foxb1* is a transcription factor gene of the forkhead family. It is expressed predominantly in the cell nucleus and is absent from the dendrites and the axons (Alvarez-Bolado et al., 2000a,b; Kaestner et al., 1993; Radyushkin et al., 2005; Wolf and Ryu, 2013; Zhao et al., 2008). *Foxb1* is expressed not only in neurons of

the parvafox nucleus but also in those of the mammillary nuclei (Zhao et al., 2008). The aim of the present study was to trace the projections of the cuff of *Foxb1*-expressing neurons of the parvafox nucleus after the stereotactic injection into *Foxb1/Cre* mice of Cre-dependent adeno-associated viral constructs that selectively target this group of cells.

The *Foxb1*-expressing neurons were found to project heavily to a number of brain sites that are known to be involved in defensive behavior. The influence exerted by the *Foxb1*-expressing component of the parvafox nucleus on these brain regions may relate to the autonomic concomitants of the defensive responses for which these regions are renowned.

MATERIALS AND METHODS

The animals used in this study were *Foxb1-Cre* mice, in which the Foxb1-immunopositive neurons coexpress the enzyme Cre-recombinase and EGFP (Zhao et al., 2007). The animals were subjected to a 12-hour dark/12-hour light cycle, maintained at a constant temperature (24°C), and fed ad libitum. The study was performed in accordance with the regulations of the Swiss animal experimentation law (permissions 26-10, 27-10, 2013_04_FR, and 2013_05_FR).

Tract-tracing experiments

Sixteen mice, weighing 25–30 g, were anesthetized with a mixture of Ketalar (Parke-Davis, Ann Arbor, MI; 75 mg/kg body weight) and xylazine (Streuli, Uznach, Switzerland; 10 mg/kg body weight). The adenoviral anterograde tracers were injected in a volume of 20 nl from a Hamilton syringe, which was connected to a stereotactic apparatus (Kopf, Tujinga, CA). The injections into the right lateral hypothalamus were made at the following coordinates: rostrocaudal Bregma −1.5 mm, mediolateral −1.4 mm, dorsoventral −5.5 mm. The viral constructs included: AAV2/1.CAG.FLEX.EGFP.WPRE.bGH, which was injected into 10 mice (see Table 2;

Abbreviations

1	layers, tenia tecta	AD	anterodorsal thalamic nucleus
3v	third ventricle	AHA	anterior hypothalamic area, anterior part
4v	fourth ventricle	AHC	anterior hypothalamic area, central
7N	facial nucleus	AHP	anterior hypothalamic area, posterior
10N	dorsal motor nucleus of vagus	AIV	agranular insular cortex, ventral part
12N	hypoglossal nucleus	Amb	ambiguous nucleus
A1	noradrenaline cells	Ambc	ambiguous nucleus, pars compacta
A13	A13 dopamine cells	AmbL	ambiguous nucle, loose
AAD	anterior amygdaloid area, dorsal part	ant	anterior hypophysis
AAV	anterior amygdaloid area, ventral part	AOB	accessory olfactory bulb
ac	anterior commisure	AOD	anterior olfactory area, dorsal part
aca	anterior commissure, anterior	AOE	anterior olfactory area, external part
Acb	accumbens nucleus	AOL	anterior olfactory area, lateral part
Acbc	accumbens nucleus, core	AOM	anterior olfactory nucleus, medial
AcbSh	accumbens nucleus, shell	AOV	anterior olfactory nucleus, ventral
ACo	anterior cortical nucleus	Arc	arcuate nucleus

AVDM	anteroventral thalamic nucleus, dorsomedial		MeAD	medial amygdalar nucleus, anterodorsal
AVPO	anteroventral preoptic nucleus		MeAV	medial amygdalar nucleus, anteroventral
aq	aqueduct		MePD	medial amygdalar nucleus, posterodorsal
Bar	Barrington's nucleus		MePV	medial amygdalar nucleus, posteroventral
Bö	Bötzinger complex		mfb	medial forebrain bundle
BST	bed nucleus of the stria terminalis		mlf	medial longitudinal fascicle
BSTL	bed nucleus of the stria terminalis, lateral division		MnR	medial raphe nucleus
BSTMA	bed nucleus of the stria terminalis, medial division, anterior part		MO	medial orbital cortex
			MPA	medial preoptic area
BSTMPI	bed nucleus of the stria terminalis, medial division, posterointermediate part		MPB	medial parabrachial nucleus
			MPOA	medial preoptic area, anterior
BSTMPL	bed nucleus of the stria terminalis, medial division, posterolateral part		MRe	mammillary recess, 3rd ventricle
			mt	mammilloothalamic tract
BSTV	bed nucleus of the stria terminalis, ventral division		mtg	mammillotegmental tract
C1	adrenaline cells		MVe	medial vestibular nucleus
CC	central canal		MVePC	medial vestibular nucleus, parvicellular part
Ce	central amygdaloid nucleus		NRTh	reticular thalamic nucleus
CeM	central amygdaloid nucleus, medial division		NTS	nucleus of the solitary tract
CGA	central gray, alpha part		Nv	navicular postolfactory nucleus
CGPn	central gray pons		Op	optic nerve layer superior colliculus
CI	inferior colliculus		opt	optic tract
CL	central lateral thalamic nucleus		OV	olfactory part of lateral ventricle
CM	central medial thalamic nucleus		Pa	paraventricular hypothalamic nucleus
CnF	cuneiform nucleus		PAG	periaqueductal gray
CPu	caudate putamen (striatum)		PaLM	paraventricular hypothalamic nucleus lateral magnocellular part
Cu	cuneate nucleus		PC	paracentral thalamic nucleus
DG	dentate gyrus		PCRt	parvicellular reticular nucleus
dlPAG	dorsolateral periaqueductal gray		PCRtA	parvicellular reticular nucleus, alpha part
DM	dorsomedial hypothalamic nucleus		PDTg	posterodorsal tegmental nucleus
DMTg	dorsomedial tegmental area		PF	parafascicular thalamic nucleus
DMV	dorsomedial hypothalamic nucleus, ventral part		PFA	paraformaldehyde
DP	dorsal peduncular cortex		PH	posterior hypothalamic area
DpG	deep gray layer, sup colliculus		PLH	peduncular part lateral hypothalamus
DPGi	dorsal paragigantocellular nucleus		PMD	premammilary nucleus, dorsal
DpMe	deep mesencephalic nucleus		PMV	premammilary nucleus, ventral part
DR	dorsal raphe nucleus		PnC	pontine reticular nucleus, caudal part
DRV	dorsal raphe nucleus, ventral part		Post	posterior hypophysis
DTgC	dorsal tegmental nucleus, central		PPTg	pedunculopontine nucleus
DTgP	dorsal tegmental nucleus, pericentral		Pr	prepositus nucleus
DTT	dorsal taenia tecta		PrBö	pre-Bötzinger complex
dVLM	dorsoventrolateral reticular nucleus		PrL	prelimbic cortex
EPl	external plexiform layer of the olfactory bulb		PSTh	parasubthalamic nucleus
f	fornix		PV	paraventricular thalamic nucleus
Fr	fasciculus retroflexus		PV2	parvalbumin-positive nucleus 2
FrA	frontal association cortex		PVA	paraventricular thalamic nucleus, anterior
g7	genu facial nerve		PVP	paraventricular thalamic nucleus, posterior
gcc	genu corpus callosum		PVT	paraventricular thalamic nucleus
Gem	gemini hypothalamic nucleus		Re	reuniens thalamic nucleus
Gi	gigantocellular reticular nucleus		Rf	fasciculus retroflexus
GlA	glomerular layer of the accessory olfactory bulb		Rh	rhomboid thalamic nucleus
Gr	gracile nucleus		RMg	raphe magnus nucleus
GrO	granular cell layer of the olfactory bulb		RpN	raphe pontis nucleus
HDB	nucleus of the horizontal limb of the diagonal band		RVL	rostroventrolateral medulla reticular nucleus
Hi	hippocampus		SC	superior colliculus
icp	inferior cerebellar peduncle		SHi	septohippocampal nucleus
InG	intermediate gray layer of the superior colliculus		SI	substantia innominata
InWh	intermed white layer sup colliculus		SOlCe	solitary nucleus, central part
IOM	inferior olive, medial nucleus		sp5	spinal trigeminal tract
IRt	intermediate reticular nucleus		Sp5C	spinal trigeminal nucleus, caudal part
LA	lateroanterior hypothalamic nucleus		Sp5I	spinal trigeminal nucleus, interpolar part
LC	locus coeruleus		Su3	supraoculomotor nucleus of the PAG
LDTg	laterodorsal tegmental nucleus		Sub	submedius thalamic nucleus
LH	lateral hypothalamic area		SuM	supramammillary nucleus
LHb	lateral habenular nucleus		SuML	supramammillary nucleus, lateral part
LM	lateral mammillary nucleus		VA	ventral anterior thalamic nucleus
LO	lateral orbital cortex		VDB	nucleus vertical limb diagonal band
LPAG	lateral periaqueductal gray		Virg	virgola
LPB	lateral parabrachial nucleus		vlPAG	ventrolateral PAG
LPGi	lateral paragigantocellular nucleus		VLTe	ventrolateral tegmental nucleus
LPO	lateral preoptic area		VM	ventromedial thalamic nucleus
LRt	lateral reticular nucleus		VMHC	ventromedial hypothalamic nucleus, central
LS	lateral septal nucleus		VMHDM	ventromedial hypothalamic nucleus, dorsomedial
LSD, LSV	lateral septal nucleus, dorsal, ventral parts		VMHVL	ventromedial hypothalamic nucleus, ventrolateral
LSI	lateral septal nucleus, dorsal, resp ventral parts		VO	ventral orbital cortex
LV	lateral ventricle		VP	ventral pallidum
LVe	lateral vestibular nucleus		VRe	ventral reuniens thalamic nucleus
MCPO	magnocellular preoptic nucleus		Vsc	ventrospinocerebellar tract
MDC	mediodorsal thalamic nucleus, central		VTA	ventral tegmental area
MDL	mediodorsal thalamic nucleus, lateral		VTg	ventral tegmental nucleus (Gudden)
MDM	mediodorsal thalamic nucleus, medial		VTM	ventral tuberomammillary nucleus
MdV	medullary reticular nucleus		ZI	zona incerta
ME	median eminence			
MeA	medial amygdalar nucleus, anterior			

Antibody	Structure of immunogen	Manufacturer, catalog No.	Dilution factor	RIID
Antiparvalbumin	Purified carp parvalbumin	SWant, lot 10-11F, PV235	1: 1,000	AB_10000343
Anti-GFP	Jellyfish (Aequorea victoria)	Life Technologies, lot A64551, rabbit polyclonal	1:2,000	AB_221570
Anti-VGluT1	Strep-Tag fusion protein of rat VGLUT 1 (aa 456–560)	Synaptic Systems, 135 403, rabbit polyclonal	1:30,000	AB_887883
Anti-VGluT2	Strep-Tag fusion protein of rat VGLUT 2 (aa 510–582)	Synaptic Systems, 135403, rabbit polyclonal	1:30,000	AB_2315570
Anti-GAD67	Purified mouse monoclonal IgG2a in buffer containing 0.1% sodium azide	Millipore, MAB5406	1:1,000	AB_2278725
TH	TH purified from rat PC12 cells	Immunostar, 22941, lot 1241002, mouse monoclonal	1:3,000	AB_572268
nNOS	Lyophilized whole-serum IgG	Immunostar, rabbit polyclonal, 24287, lot 741001	1:2,000	AB_572256

[1]Well characterized antibodies were acquired from various commercial sources.

194/13, 195/13, 249/13, 9/15, 10/15, 180/15, 182/15, 183/15, 25/16, 26/16), and AAV2/1.CAG.FLEX.Tomato.WPRE.bGH, which was injected into eight mice (see Table 2; 307/13, 392/14, 9/15, 10/15, 131/15, 180/15, 25/16, 26/16). The constructs were developed at the Allen Institute and are commercially available from the Vector Core at the University of Pennsylvania (http://www.med.upenn.edu/gtp/vectorcore/).

Four weeks after the injections, the mice were transcardially perfused first with physiological (0.9%) saline and then with a 4%- solution of paraformaldehyde in 0.1M phosphate buffer, [(PB) pH 7.3]. The brains and the spinal cords were excised, immersed overnight in the same fixative, and then transferred for cryoprotection to a chilled solution of 30% sucrose (4°C). Forty-micrometer-thick coronal, parasagittal and horizontal cryosections were prepared with a freezing microtome (Reichert-Jung, Vienna, Austria) and collected in PB.

Immunohistochemistry

In some tracing experiments, free-floating sections were exposed for 2 days at 4°C to a rabbit anti-GFP antiserum (Life Technologies, Japan; catalog No. A6455, RRID:AB_221570; diluted 1:3,000 in Tris-buffered saline [TBS] containing 0.2% Triton X-100 and 10% bovine serum). The sections were rinsed in TBS and then exposed for 2 hours at room temperature to the anti-rabbit biotinylated secondary antibody (Vector Laboratories, Burlingame, CA; diluted 1:200 in TBS + 10% bovine serum). After rinsing once in TBS and twice in Tris-HCl (pH 8.2), the sections were exposed for 2 hours to Alexa 488-conjugated streptavidin (Jackson Immunoresearch; diluted 1:500 in Tris-HCl, pH 8.2). To identify the region of the parvafox nucleus, all sections were subsequently exposed, in parallel, for 1 day at room temperature to a primary monoclonal antibody against Parv (PV235; SWant, Marly, Switzerland; catalog No. 235, RRID:AB_10000343; diluted 1:1,000 in TBS + 0.2% Triton X-100 + 10% bovine serum) and then for 2 hours to an anti-mouse Cy3 antibody (Jackson Immunoresearch; diluted 1:200 in Tris-HCl, pH 8.2).

To reveal the neurotransmitter used by the terminals, antisera against the glutamate transporters VGlut1 and VGlut2 (Synaptic Systems, Göttingen, Germany; catalog No. 135 403, RRID:AB_887883; Synaptic Systems, catalog No. 135 103 RRID:AB_2315570) and the GABA-synthesizing enzyme GAD67 (Millipore, Darmstadt, Germany; catalog No. MAB5406, RRID:AB_2278725) were employed. To identify the catecholaminergic sites, antibodies against tyrosine hydroxylase (TH; Immunostar, Hudson, WI; catalog No. 22941, RRID:AB_572268) were used. To identify the dLPAG, an antiserum against the neuronal nitric oxide synthase (nNOS; Immunostar; catalog No. 24287, RRID:AB_572256) was employed (Table 1).

Cell nuclei were sporadically revealed by counterstaining with DAPI (diluted 1:2,000 in phosphate-buffered saline) for 5 minutes at room temperature. The sections were mounted and evaluated in a Leica 6000 epifluorescence microscope (equipped with a Hamamatsu C4742-95 camera), a digital slide scanner (Nanozoomer, Hamamatsu), or a Leica TCS SP5 confocal laser microscope.

The images were postprocessed for brightness and contrast in Adobe Photoshop CS5. The image stacks were prepared with ImageJ 1.44p (RRID:SCR_003070), and the figures were collated in the Adobe Illustrator CS6 (RRID:SCR_014198).

TABLE 2.

Distribution of Labeled Axons and Terminals in the Brain After Injection of Cre-Dependent Tracers in the Ventrolateral Hypothalamus of *Foxb1-Cre* Mice[1]

Brain area	Region	Stained fibers	Terminals	Other injections
Telencephalon	Lateral septum, ventral (LSV)		+	
	Lateral septum, intermediary (LSI)		+	
	Accessory olfactory bulb (AOB)		+ +	
	Anterior olfactory area, dorsal part (AOD)		+ +	
	Anterior olfactory area, external part (AOE)		+ +	
	Anterior olfactory area, lateral part (AOL)		+ +	
	Anterior olfactory area, medial part (AOM)		+ +	
	Granular cell layer olfactory bulb (GrO)		+ +	
	Lateral orbital cortex (LO)		+	
	Medial orbital cortex (MO)		+	
	Ventral orbital cortex (VO)		+	
	Nucleus of the stria medullaris (SM)	+	+	
	Substantia innominate (SI)	(+)	(+)	
Thalamus	Anterodorsal thalamic nucleus (AD)		(+)	+ + + 195-13
	Anterodorsal tegmental nucleus (AVDM)		+	
	Centromedial thalamic nucleus (CM)		+	
	Central thalamic nucleus (CL)		+	
	Field of Forel (FF)		+	
	Mediodorsal thalamic nucleus, central (MDC)		+	
	Medial habenular nucleus (MHB)		+ +	
	Paraventricular hypothalamic nucleus lateral magnocellular part (PALM)	+ +	+ + Bilat	
	Parafascicular thalamic nucleus (PF)		+ +	
	Prerubral nucleus (PR)		+	
	Precommissural nucleus (PrC)		+ + Bilat	
	Periventricular fiber system (Pv)	+ + +		
	Paraventricular thalamic nucleus (PV)		+	
	Paraventricular thalamic nucleus, anterior (PVA)			
	Paraventricular thalamic nucleus, posterior (PVP)		+ +	
	Reuniens thalamic nucleus (Re)		+	
	Submedius thalamic nucleus (SUB)		+ +	
	Zona incerta (bilateral) (ZI)	+ +	+ + bilat	
Hypothalamus	Anterior hypothalamic area, anterior and posterior (AHA and AHP)		+	
	Gemini hypothalamic nucleus (Gem)		+ + +	
	Lateroanterior hypothalamic nucleus (LA)	(+)	(+)	
	Lateral hypothalamus (LH)	+ +	+ +	
	Magnocellular preoptic nucleus (MCPO)	+ +	+ +	
	Medial forebrain bundle (mfb)	+ +		
	Parasubthalamic nucleus (PSTh)		+ + +	
	Tuber cinereum (TC)	+	+ +	
	Ventral tuberomammillary nucleus (VTM)		+ +	
Mesencephalon	Cuneiform nucleus (CnF)		+ + +	
	Kölliker-Fuse nucleus (KF)			+ + 9-15 Red 246-15
	Dorsolateral periaqueductal gray (dLPAG)		+ +	
	Lateral periaqueductal gray (lPAG)		+ + +	
	Deep mesencephalic nucleus (DpMe)		+ + +	
	Parvalbumin nucleus 2 (PV2)		+ + +	
	Peduncolopontine tegmental nucleus (PPTg)		+ +	
	Supraoculomotor nucleus (Su3)		+ + +	
	Ventrolateral nucleus (VL)		+	
Superior colliculus	Deep gray layer (DpG)		+ +	
	Intermediate gray layer (InG)		+ +	
	InWh intermediate white layer	+ +		
	Posterior pretectal nucleus, olivary pretectal nucleus (PPT, OPT)	+	+ +	
Tegmentum	Barrington (Bar)		+ +	

TABLE 2. *Continued*

Brain area	Region	Stained fibers	Terminals	Other injections
	Central gray, alpha and beta (Cga and CgB)		+	
	Central gray pons (CgP)		+	
	Dorsal tegmental nucleus, central (DTgC)		++	
	Dorsal tegmental nucleus, pericentral (DTgP)		++	
	Locus coeruleus (LC)		+	
	Laterodorsal tegmental nucleus (LDTg)		++	
	Microcellular tegmental nucleus (MiTg; border)		++	
	Posterodorsal tegmental nucleus (PDTg)		++	
	Pontine reticular nucleus caudal (PnC)	++	+	
	Sublaterodorsal nucleus (SUL)		+	
	Ventral tegmental area (VTA)	++	+	
Metencephalon	Medial lemniscus (ml)	++		
	Dorsal paragigantocellular nucleus (DPGi)		++	
	Deep mesencephalic nucleus (DpMe)	++	+	
	Gigantocellular nucleus (Gi)	+	+	
	Intermediate reticular nucleus (IRt)		++	
	Lateral paragigantocellular nucleus (LpGi)		+	
	Lateral reticular nucleus (LRt)		++	
	Lateral vestibular nucleus (LVe)	+		
	Med reticular nucleus dorsal (MdD)	+	+	
	Medial vestibular nucleus, parvicellular (MVePC)	+		
Myelencephalon	Pontine reticular nucleus (PnC)		(+)	
	Pedunculopontine tegmental nucleus (PpTn)		++	
	Nucleus retroambiguus (Ramb)		++	
	Spinal trigeminal nucleus, interpolar part (Sp5I)	+		

[1]The brain areas in which terminals and/or axons arising from the site of the injections in the lateral hypothalamus have been observed through the caudal and the rostral part of the brains (+) very low density. +, Low density; ++, high density; +++, very high density.

Characterization of the antisera

The monoclonal antiserum against Parv (SWant; catalog No. PV235) recognizes only one band (corresponding to a molecular weight of 12 kD) in Western blots of brain extracts that have been derived from various species, including rats and mice. The pattern of staining in the brain was consistent with previously documented findings (Celio, 1990). No staining is observed in the brains of Parv knockout mice (Schwaller et al., 1999).

The antiserum against GFP (Life Technologies; catalog No. A6455) was raised against the full-length recombinant molecule. Staining was observed only in the parvafox nucleus of the ventrolateral hypothalamus and in the mammillary nuclei. In other parts of the brain of animals that had been injected with a GFP-bearing viral construct for tracing, the pattern of distribution accorded with the expected one.

The specificity of the antiserum against VGlut2 (Synaptic Systems; catalog No. 135 403) was demonstrated by Western blot analysis of homogenized rat brains and/or crude synaptic vesicle in which a single, broad band corresponding to a molecular weight slightly above 60 kD was manifested and by preadsorbing the sections with the GST fusion protein that was used as an immunogen, which abolished all immunoreactivity (Zhou et al., 2007).

The specificity of the antiserum against VGlut1 (Synaptic Systems; catalog No. 135 302) has been demonstrated by Western blot analysis of homogenized rat brains and/or crude synaptic vesicle fractions from rat brain, in which a single, broad band slightly above 60 kD was found, and by preadsorption experiments with the GST fusion protein used as immunogen, in which the immunostaining of rat brain sections was completely eliminated (as for VGlut 2)

The specificity of the antiserum against glutamic acid decarboxylase 67 (GAD67; Millipore; catalog No. MAB5406) has been demonstrated by the reaction with the 67-kD isoform of glutamate decarboxylase (GAD67) of rat, mouse, and human origins (other species not yet tested). No detectable cross-reactivity with GAD65 by Western blot on rat brain lysate compared with blot probed with AB1511 that reacts with both GAD65 and GAD67 was seen.

The specificity of the antiserum against tyrosine hydroxylase (TH; Immunostar; catalog No. 22941) has been demonstrated with standard immunohistochemical methods. The antiserum demonstrates significant labeling of rat catecholamine neuron systems via the indirect immunofluorescent and biotin/avidin–HRP techniques. The antibody recognizes the 34-kD catalytic core of TH.

The antiserum against neuronal nitric oxide synthase (nNOS; Immunostar; catalog No. 24287, lot 741001) was tested by using standard immunohistochemical
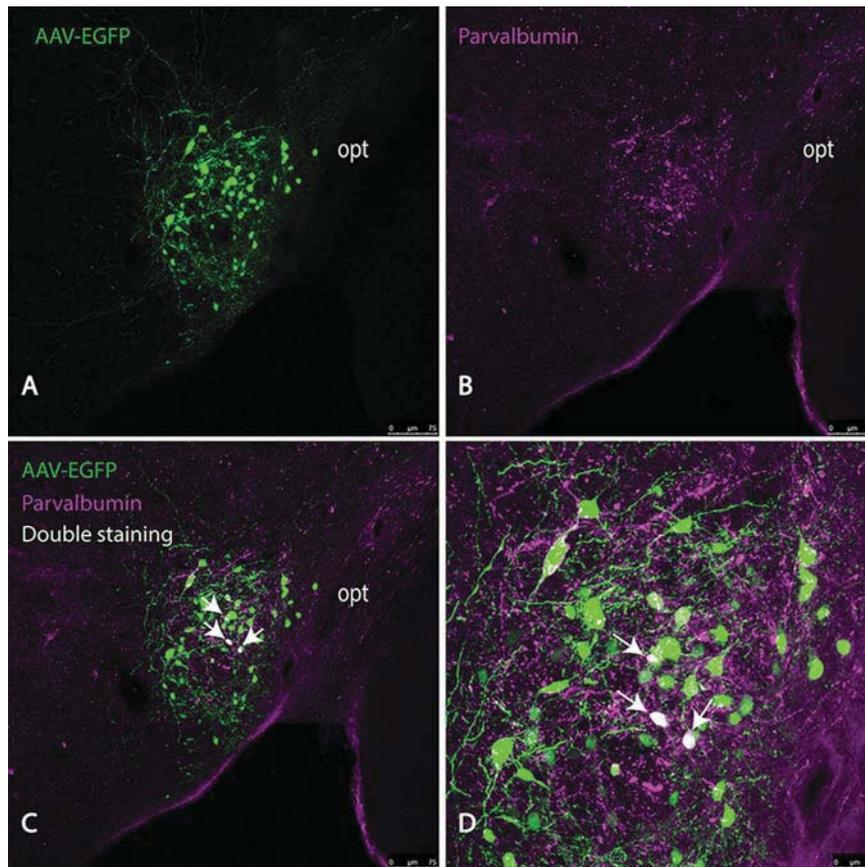
Figure 1. Confocal images of a cluster of *Foxb1/Cre*-expressing neurons in the ventrolateral hypothalamus after a stereotactic injection of the Cre-dependent AAV-EGFP construct. **A:** EGFP is expressed in the perikarya, the dendrites, and the axons of a large number of small cells. **B:** The Parv-immunopositive neurons are located within the bounds of the cluster of AAV-EGFP-labeled cells and are less numerous than these. **C:** A few of the neurons manifest dual coloring (arrows), which indicates that AAV-EGFP and Parv are colocalized. **D:** In this higher magnification view of C, the double staining is clearly visible (injection 195/13). Scale bars = 75 μm in A–C; 25 μm in D.

TABLE 3.

Quantification of Labeled Neurons in Four Injections[1]

	Mouse line	Marker	Region	Estimated population (using the mean section thickness	CE
194/13	Foxb1/Cre	AAV2/1-Cre virus	vLH	1,027.05	0.10
195/13	Foxb1/Cre	AAV2/1-Cre virus	vLH	879.14	0.10
249/13	Foxb1/Cre	AAV2/1-Cre virus	vLH	339.88	0.16
307/13	Foxb1/Cre	AAV2/1-Cre virus	vLH	1,206.34	0.11

[1]The mean total number of cells that took up the AAV2/1-Cre-dependent virus in the lateral hypothalamic parvafox nucleus of the *Foxb1/Cre* mice was estimated by using the optical fractionator. Given that the total number of Foxb1-expressing neurons in the parvafox nucleus has been estimated to be 1,200 (Bilella et al., 2014), almost all were infected with the virus in experiments 194/13 and 307/13.

methods. The antiserum worked properly for rat hypothalamus, striatum, cortex, and spinal cord as judged from indirect immunofluorescent and biotin/avidin-HRP techniques. Western blot analysis of brain homogenates shows the antibody specifically labels a band of approximately 155 kD. Immunolabeling is completely abolished by preadsorption with synthetic human nNOS (1419–1433) at 5 μg/ml of diluted antibody. No cross-reactivity with other forms of NOS was observed.

Cell counting

In four mice, the *Foxb1*/EGFP-positive neurons that took up the viral vectors were counted using the optical fractionator (West et al., 1991). This quantification was performed with in Stereoinvestigator 10.52 (MBF-Bioscience, Williston, VT; RRID:SCR_002526) mounted on a Zeiss photomicroscope (equipped with Hamamatsu Orca 0G5 camera). Counts were made on a uniform random systematic sample of every fourth coronal section. For
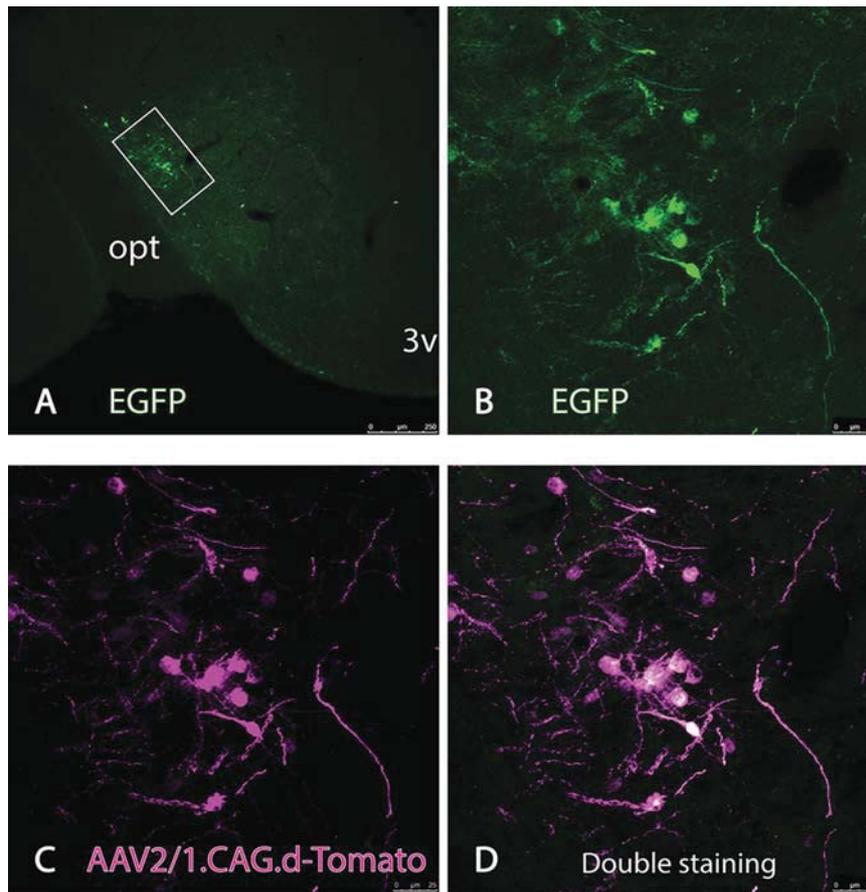
Figure 2. Confocal images of the parvafox nucleus after a stereotactic injection of the AAV-dTomato construct. **A:** Overview of the nucleus after exposure to an antibody against EGFP. The presence of intrinsic EGFP is thereby revealed. **B:** Higher magnification view of A. **C:** dTomato is expressed in the soma, the dendrites, and the axons of the *Foxb1/EGFP*-expressing neurons in the parvafox nucleus. **D:** Dual staining for Foxb1 and EGFP is clearly visible, attesting to the high specificity of the viral vector for the *Foxb1*-expressing neurons (injection 307/13). Scale bars = 250 μm in A; 25 μm in B–D.

the stereological estimations, the region of interest on the sections was first delineated. A counting frame of 70 × 70 μm was placed at 100-μm intervals along the x- and the y-axes of the section. The grid size was adapted to achieve an acceptable coefficient of error (CE; see below). At each counting site, the thickness of the section was estimated to be 30 μm.

The CE expresses the precision of the estimate (Slomianka and West, 2005). An average CE value of less than 0.15 was deemed to be appropriate for the present study. The adopted sampling scheme yielded on average CE of 0.12 (Table 2). To ensure that most of the variance across the animals was attributable to the experimental manipulations and to biological variation (but not to the estimation of the cell number), the CE^2/CV^2 ratio was calculated. Less than 10% of the variance across the animals ($CE^2/CV^2 = 0.085$) originated from the estimation of cell number.

RESULTS
Injection cases

In 13 of the 16 *Foxb1*-EGFP/Cre mice, the injections precisely hit the region of the lateral hypothalamus in which the *Foxb1*-expressing neurons were located (injections 194/13, 195/13, 249/13, 307/13, 392/14, 9/15, 10/15, 131/15, 180/15, 182/15, 183/15, 25/16, 26/16; see Table 2). In the remaining three cases, the injections were slightly too lateral, too medial, or too distal and also hit the mammillary bodies. These three brains served as controls. The most comprehensive analyses were performed on five representative data sets that implicated the parvafox nucleus most exclusively (injections 194/13, 9/15, 10/15, 131/15, 180/15).

Labeled cell bodies were restricted to the ventrolateral hypothalamus (Fig. 1A), namely, to the region in which the parvafox nucleus is located, as revealed by
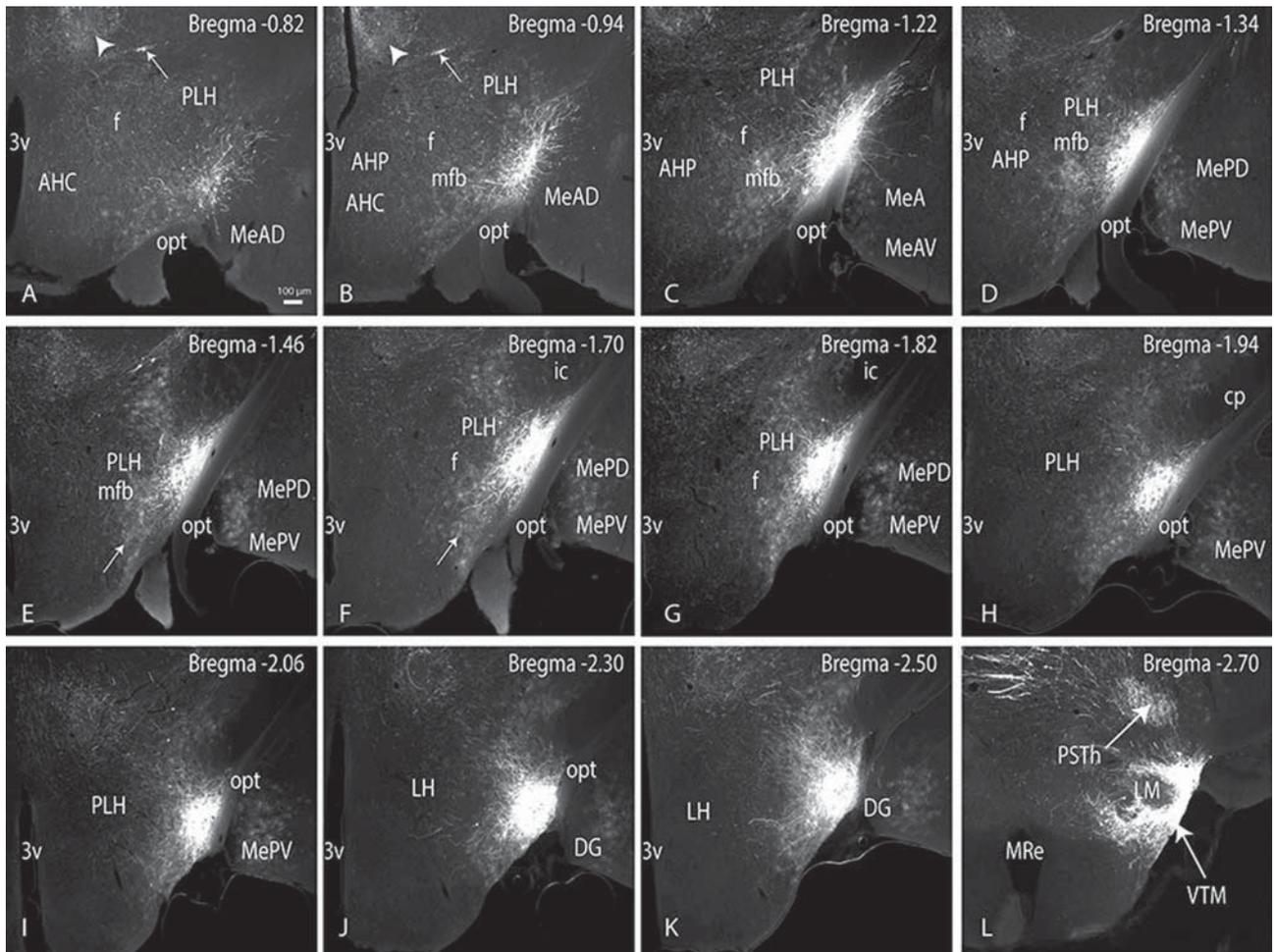
Figure 3. Epifluorescent images of 12 consecutive coronal sections through the parvafox nucleus, which was cut in a rostral (A)-to-caudal (L) direction [Bregma $\sim$ –0.82 mm (A) to Bregma $\sim$ –2.70 mm (L)]. The Cre-dependent AAV-EGFP has been taken up by the perikarya and fills the dendrites and the axons of the *Foxb1/Cre*-expressing cells. **A,B:** Parvafox nucleus, proximal portion. Only few *Foxb1*/Cre-expressing neurons are visible. The axons from the nucleus run throughout the lateral hypothalamus and reach the zona incerta (arrow) and the ventral thalamic nucleus (arrowhead) giving rise to descending and ascending bundle of fibers, respectively (Bregma $\sim$ –0.82 mm, Bregma $\sim$ –0.94 mm). **C–E:** The Foxb1-Cre-expressing neurons send axons that spread in all directions; fibers pass through the peduncular part of the lateral hypothalamus (PLH), and most of them flow into the medial forebrain bundle reaching several brain regions (Bregma $\sim$ –1.22 mm, Bregma $\sim$ –1.34 mm, Bregma $\sim$ –1.46 mm). **F,G:** Middle portion of the parvafox nucleus. The neurons send projections in all directions; bundles of fibers run around the internal capsule, and others run in the same direction of the nucleus (Bregma $\sim$ –1.70 mm, Bregma $\sim$ –1.82 mm). **H–K:** Four consecutive sections through the most distal portion of the parvafox nucleus. The *Foxb1*/Cre-expressing neurons send projections in different directions (Bregma $\sim$ –1.94 mm, Bregma $\sim$ –2.50 mm). **L:** This panel shows only the dense bundle of fibers that run around the lateral mammillary nucleus (LM), throughout the ventral tuberomammillary nucleus, giving rise to the descending projections, although a part of axons pass through the parasubthalamic nucleus (PSTh; Bregma $\sim$ –2.70 mm; injection 194/13; the same animals as in Figs. 1, 5, 11, and 13). Scale bar = 100 μm.

immunostaining for Parv (Fig. 1B) and GFP in Foxb1-Cre-GFP mice. In a few cases, the virus spread to astrocytes, but they fluoresced only very weakly (Fig. 3D,E). In some experiments, *Foxb1*-expressing cells of the mammillary nuclei were colabeled. The pathways emanating from these nuclei, namely, the mammillothalamic and the mammillotegmental tracts, were readily identified; so, too, were the corresponding terminal fields in the anterior thalamic nuclei (see Figs. 7D, 8A,B) and in

the ventral and the dorsal tegmental nuclei (see Fig. 9). These specimens served as controls in the present study. Some of the Foxb1-immunopositive cells are known also to express Parv (Bilella et al., 2014). This finding was confirmed in the present study (see merged image in Fig. 1C,D). In 13 of the 16 injections, virtually all of the *Foxb1*-expressing neurons that are known to occur in the parvafox nucleus (Bilella et al., 2014) were infected with the virus (see Table 3). An example of this
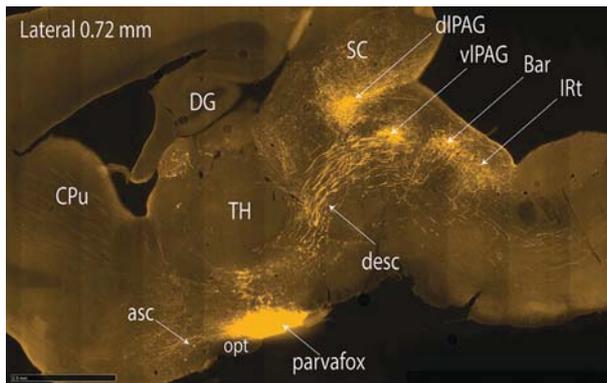
Figure 4. Axonal projections deriving from the Foxb1 neurons of the parvafox nucleus. Parasagittal sections of specimen 131-15. The injection site in the ventrolateral hypothalamus visualizes selectively the Foxb1-Cre-expressing neurons. From this Foxb1 aggregate of neurons, bundles of axons leave in various directions: ascending (asc), thalamic (TH), intrahypothalamic, and descending (desc). Various terminal fields, particularly those in the midbrain, are indicated.

can be seen in Figure 2, in which all of the neurons injected with the adenovirus AAV-tomato construct expressed the EGFP component (indicating the expression of *Foxb1*/Cre) as well.

Injection site

A representative injection site is shown in Figure 3. It depicts AAV-laden, *Foxb1*-expressing neurons and their interlacing dendrites. Together, these formed a dense, oval cluster with its long axis oriented parallel to the optic tract, running from Bregma –0.824 mm to Bregma –2.70 mm. The dendrites of the neurons were disposed in the coronal plane (clearly visible in Fig. 3B,C). Axons exited the labeled *Foxb1*-expressing neurons in various directions that have been described in detail by Nieuwenhuys as characteristic for the "hypothalamic aggression area" (Roeling et al., 1993, 1994a; see summary Fig. 16), namely, ascending (dorsal, lateral, and ventromedial), thalamic (dorsal and ventral), intrahypothalamic, and descending (dorsal, lateral, and medial).

Morphology of labeled axons

Morphologically, the axons exhibited varicosities, branches, and distinct terminal arborizations with patent "boutons terminaux," In the finest branches, the labeled axons displayed varicosities suggestive of synapses, the presence of which was confirmed by immunostaining for VGlut2 (Fig. 5). The density of the varicosities differed according to the area of passage. Indeed, many of the traversed brain areas could be innervated in this way.
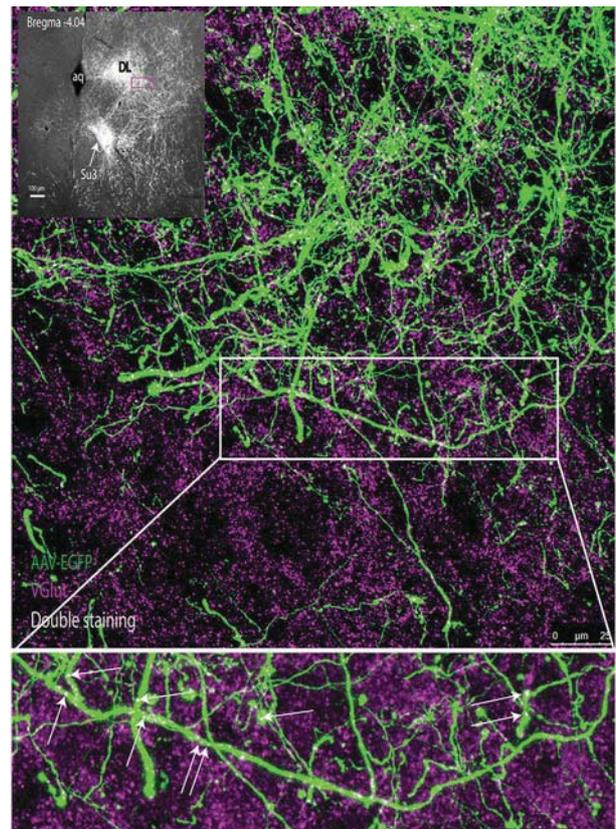

Figure 5. Confocal image illustrating the distribution of the vescicular glutamate transporter 2 (VGluT-2) in the glutamatergic, *Foxb1*-expressing terminals of the dorsolateral periaqueductal gray. The black and white inset indicates the region from which the image was taken. The lower **inset** depicts at higher magnification the small white spots within the immunopositive axonal endings (arrows). They are indicative of the merger of violet and green coloring and indirectly of the presence of the VGlut-2 transporter (Bregma –4.04 mm, injection 194/13). Scale bar = 25 μm.

Summary of results

The Foxb1-expressing neurons of the parvafox nucleus project to the telencephalon, diencephalon (thalamus and hypothalamus), midbrain, and hindbrain (Figs. 4, 6; see also summary Fig. 16). The strongest projection is to the periaqueductal gray (PAG), and the rest of the targets are sparsely labeled in comparison.

Ipsilateral vs. contralateral projections

The largest contingent of projections and terminals was observed ipsilaterally. However, in some regions, a contralateral mirror image of the terminals was seen (see below). The PAG and thalamic staining (e.g., the "virgolas"; see Fig. 8B) were exceptional in this respect, insofar as staining was observed only ipsilaterally. In what follows, we present the results systematically from rostral to caudal.

Figure 6. Schematic representation of a sagittal section of the mouse brain upon which the tracts leaving the Foxb1-expressing neurons in the lateral hypothalamus are projected (adapted from Roeling et al., 1993, 1994b). Three axonal tracts project rostrally, three distally, and two to the thalamus and hypothalamus. From these tracts, eight pathways are generated, which follow different courses and target different organs: 1) rostrodorsal, 2) rostrolateral, 3) rostromedial, 4) medial thalamic, 5) lateral thalamic, 6) distal dorsal, 7) laterodistal, and 8) medio-distal forebrain bundle. The anatomical organization of the ventral medulla is modified from the schematic drawing by Jänig (2006). The abbreviation listed here represent the structures that receive the terminals: AD, anterodorsal thalamic nucleus; AMBc, nucleus ambiguus, compact; AMBL, nucleus ambiguus, loose; AOL, anterior olfactory area, lateral part; AOM, anterior olfactory nucleus, medial; Bar, Barrington's nucleus; Bö, Bötzinger complex; CnF, cuneiform nucleus; dlPAG, dorsolateral periaqueductal gray; Gem, gemini hypothalamic nucleus; Gi, gigantocellular formation; HDB, nucleus of the horizontal limb of the diagonal band; IRt, intermediate reticular nucleus; LDTg, leterodorsal tegmental nucleus; lPAG, lateral periaqueductal gray; LRt, lateral reticular nucleus; LSI, lateral septal nucleus; LSV, lateral septal nucleus, ventral part; PFC, prefrontal cortex; PHA, posterior hypothalamic area; PPTg, posterior tegmental nucleus; PSTh, parasubthalamic nucleus; PV2, parvalbumin nucleus 2; PVT, paraventricular thalamic nucleus; RAMB, retroambiguus; Re, reuniens thalamic nucleus; rVLM, rostral ventrolateral medulla; SC, superior colliculus; Si, substantia innominata; Su3, suproculomotor nucleus; Zi, zona incerta.

Telencephalic projections

The ascending tract followed the mfb and consisted of a diffuse contingent of fine axons traversing the medial and the lateral preoptic area and, after reaching the diagonal band, proceeding to the ventral part of the lateral septal nucleus (Fig. 7). Rostrally, they ascended to the anterior amygdaloid nucleus, ran in the bed nucleus of the stria terminalis, and climbed to the lateral septum (Fig. 7). A few terminals were observed in the medial and lateral portions of the orbital cortex (Fig. 6), the olfactory nuclei (Fig. 6), and the olfactory bulb, lateral to the olfactory ventricle (see summary Fig. 16A, 16B).

Diencephalic projections

Stemming from the cells that were infected with the viral construct, the labeled axons passed into the mfb, ascended to the reuniens thalamic nuclei, and terminated in the dorsal thalamic nucleus (medial and lateral portions; Fig. 8; see also summary Fig. 16C,D,E,F). In the thalamic tract, considerable numbers of fibers run dorsal to the fornix to attain the paraventricular nucleus of the hypothalamus (Fig. 8C). Axons crossed the supraoptic decussation to reach the contralateral parvafox nucleus in the lateroventral hypothalamus. Contralaterally projecting axons were observed also in the supramammillary decussation.

Midbrain projections

Fibers from the parvafox nucleus coursing caudally went through the mfb, ran lateral to the mammillary bodies in the ventral tuberomammillary nucleus (VTM; Fig. 3K,L), traced a wide path through the ventral tegmental region, surrounded the fasciculus retroflexus, rose in dorsal direction in the posterior hypothalamus, and attained the PAG by passing through the medial zona incerta. Labeled axons reached the midbrain through three different pathways: 1) the dorsal (periventricular) fiber system, 2) the lateral (central tegmental) tract, and 3) the mfb.

Periventricular axons

The axons of the periventricular system formed thick bundles, which swept mediodorsally at premammillary levels in a periventricular location and then bent dorso-caudally, running alongside the lateral and ventral aspect of the central gray matter to the central tegmental field of the midbrain (Fig. 9).

Figure 7. Coronal sections through the septum in a proximal (Bregma 1.54 mm; A) to distal (Bregma 0.62 mm; D) sequence. A low-density bundle of fibers from the parvafox nucleus passes through the navicular postolfactory nucleus (NV; **A**), the septal nuclei (A,B), and the nucleus of the vertical limb (**B**) headed for the frontal regions. **C:** High magnification of the region of the diagonal band, showing a rich collection of scattered axons and probable endings in this region. **D:** Projections in the thalamic region, in the anterior paraventricular thalamic nucleus (PVA), in the central medial thalamic nucleus (CM), and in the reuniens thalamic nucleus (Re; injection 249/13; the same animal as in Fig. 4). Scale bar = 100 μm in A (applies to A,B); 500 μm for C,D.

Central tegmental tract

The central tegmental tract conveyed axons that ran laterodorsally in the midbrain and innervated the deep gray and intermediate white layers of the superior colliculus (see Fig. 13) as well as the dorsolateral tegmental nucleus (Fig. 9).

mfb

The third major descending pathway is the mfb, which penetrates the ventral tegmental area. In the supra-mammillary decussation, some of the fibers switched to the contralateral side. From this region, axons coursed ventrodorsally in the midline and passed between the oculomotor region and the interstitial nucleus of Cajal into the ventromedial portion of the central gray matter to innervate the Su3 region. Some of the axons remaining in the mfb could be traced caudally in the pontine tegmentum and formed part of the ventromedial and the ventrolateral brainstem fiber system.

PAG projections

The *Foxb1*-expressing neurons projected mainly to five different areas that lay within or close to the PAG,

namely, to a wedge-shaped region that encompassed the dorsolateral and the upper lateral columns (dL; L in Figs. 10A–G, 11), to a PAG-skirting column that lay ventral to the aqueduct (Su3 in Fig. 10D,E), to the deep mesencephalic nucleus (DpMe; Fig. 10E–G), to the cuneiform nucleus (CnF; Fig. 10H,I), and to the deep layers of the superior colliculus. At rostral levels of the PAG, the projections were largely restricted to a wedge-shaped segment that occupied the dorsolateral and the upper portions of its lateral columns (Figs. 10A–C, 11), mainly peripherally. This dense collection of terminal fields, of which the upper half was coincident with nNOS immunoreactivity (Fig. 11), extended from Bregma −2.80 mm rostrally to Bregma −5.00 mm distally (Fig. 10A–G) and was therefore the most prominent terminal field in the PAG (see also scheme in Fig. 12). At the midlevels of the PAG, both the dorsolateral/ upper lateral columns and the Su3 were labeled (Fig. 10C–E), and the projection from the parvafox nucleus was distinctly bicolumnar. With the addition of the deep mesencephalic nucleus (DpMe; Fig. 10E–G), which lies outside the PAG, the projection became tri-columnar. At the other caudal levels of the PAG, the densest labeling coincided with the cluster of other Parv-
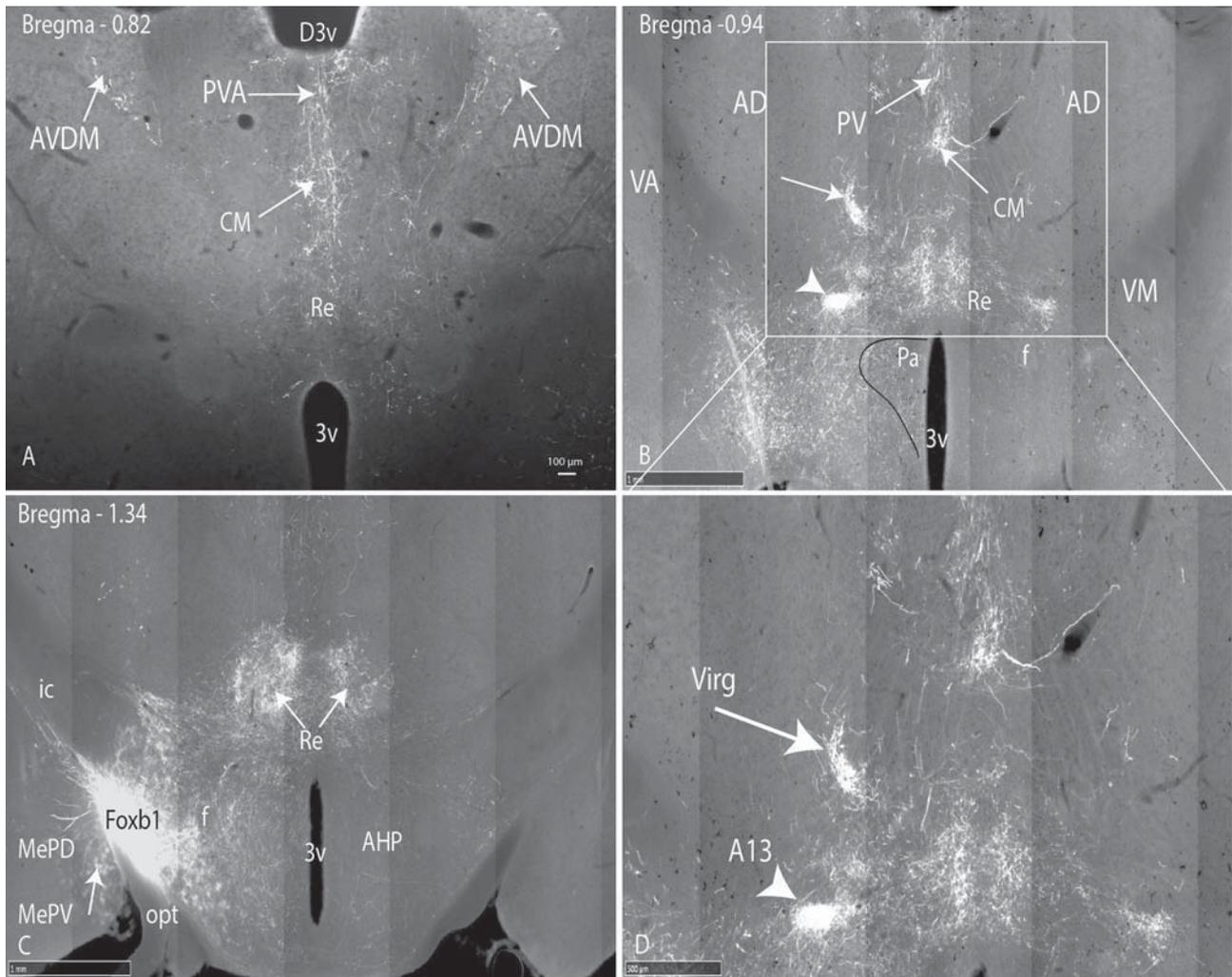
Figure 8. Epifluorescent images of coronal sections through the thalamus illustrating the projections thereto from the parvafox nucleus. **A:** At level Bregma −0.82 the axons from the *Foxb1*-expressing neurons project to the anterior paraventricular thalamic nucleus (PVA), the central medial thalamic nucleus (CM), and the reuniens thalamic nucleus (Re). The anterodorsal and anteroventral thalamic nuclei (AD, AVDM), receive a few terminals. **B:** Hypothalamic and thalamic projections (Bregma −0.94) are rich in the rostral part of the lateral hypothalamus. A dense, oval, ispilaterally stronger ending field is recognizable at the level of the zona incerta and may represent the A13 cell group (arrowhead). Slightly dorsally, a second, strictly ipsilateral, dense terminal field is probably restricted to the PC nucleus of the thalamus. We denote this small region as the "virgola field" because of its inverted comma shape (arrow; injection 249/13). **C:** Coronal section through thalamus and hypothalamus at level Bregma −1.34. The strong, overexposed labeling of the Foxb1 neurons in close proximity to the optic tract (ot) is readily recognizable. The rich terminal field in the reuniens nucleus is visible as well as axons passing through and terminals endings in the region of the zona incerta. The lateral hypothalamus is richly innervated by terminals deriving from Foxb1 axons whereas endings are rare or absent in the ventromedial hypothalamic nucleus. Note particularly the presence of endings in the lateral part of the periventricular nucleus of the hypothalamus. Axons pass also through the region of the cortical amygdala, in which some marked astrocytes also can be observed. **D:** Higher magnification of the thalamic labeling in B, with the two, aforementioned, strong and dense but spatially restricted terminal fields in the A13 region and in the "virgola field." Scale bar = 100 μm.

expressing neurons located in the LDTG, viz. in the so-called PV2 nucleus (Celio et al., 2013) and in the cuneiform nucleus (see also Fig. 10H,I).

The Foxb1 projections to the PAG reached their targets mainly ipsilaterally (Table 2), although in the Su3 a few contralateral ones were also seen (Fig. 10). The various terminal fields of the Foxb1 axons in the PAG are schematized in Figure 12 (and see also summary

Fig. 16G,H). The projection to the PAG is glutamatergic, which was revealed by immunostaining for various neurotransmitter transporters (particularly VGlut2-positive ones; Fig. 5).

Hindbrain projections

A small number of axons forming part of the periventricular projection to the PAG went on caudally as part
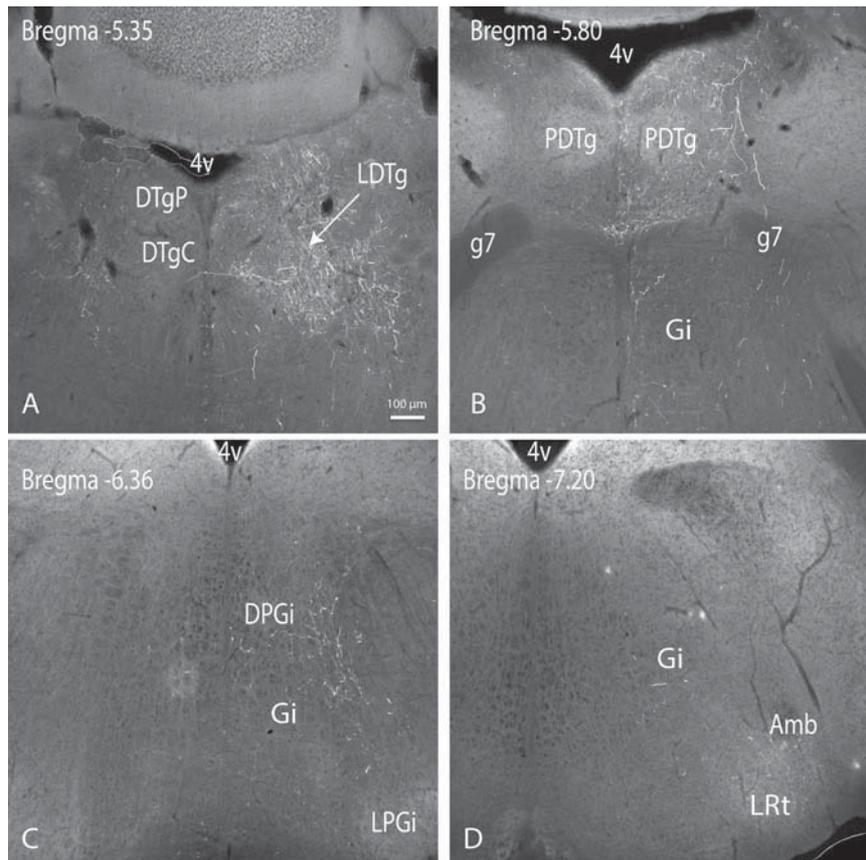
Figure 9. Epifluorescent images of coronal sections through the hindbrain, illustrating the projections thereto of axons emanating from the hypothalamic *Foxb1*-expressing neurons. The fibers traverse the dorsolateral tegmental nucleus (LDTg; Bregma ∼ –5.35 mm; **A**), pass into the pons (Bregma ∼ –5.80 mm; **B**), and continue their course through the dorsal paragigantocellular nucleus (DPGi; Bregma: ∼ –6.36 mm; **C**). Projections are visible also in the gigantocellular reticular (Gi) and the lateral reticular (LRt) nuclei (**D**; injection 194/13). Scale bar = 100 μm.

of the dorsal brainstem fiber system. They terminated at three different hindbrain levels: 1) in the pontine tegmentum (including vestibular nuclei, laterodorsal tegmental nucleus, pedunculopontine nucleus, cuneiform nucleus, and Barrington's nucleus), 2) in the reticular substance (gigantocellular, intermediate, and lateral reticular nuclei), and 3) in the medulla oblongata Bötzinger complex, peri- and retroambigual regions; (see summary Fig. 14, 15, 16)).

Medulla oblongata
Some terminals ended in the retrofacial region of the ventrolateral medulla (Figs. 6, 14), in a region rich in vertically oriented TH-positive axons but devoid of TH-positive neurons, which may correspond to the Bötzinger complex, involved in the regulation of expiration (Fig. 14). The rostral ventrolateral medulla, a vasomotor center, also received terminal endings from the Foxb1-expressing neural population of the parvafox nucleus (Fig. 15A,A′,B,B′). At these levels the compact

portion of the nucleus ambiguus was surrounded, particularly ventrally, by labeled axons and terminals, but none was observed within the boundaries of the nucleus. In the loose part of the ambiguous and in the retroambigual region, on the other hand, many axons terminated profusely in regions known to harbor neurons that control the cardiovascular system and the larynx (vocalization; Fig. 15C,C′)

Spinal projections
The Rexed layer X of the upper cervical segments received scattered terminals.

DISCUSSION
The parvafox nucleus projects widely
The projections of the subpopulation of *Foxb1*-expressing neurons in the lateral hypothalamic parvafox nucleus radiate in diverse directions. They extend rostrally to the olfactory bulbs, the orbitofrontal cortex,
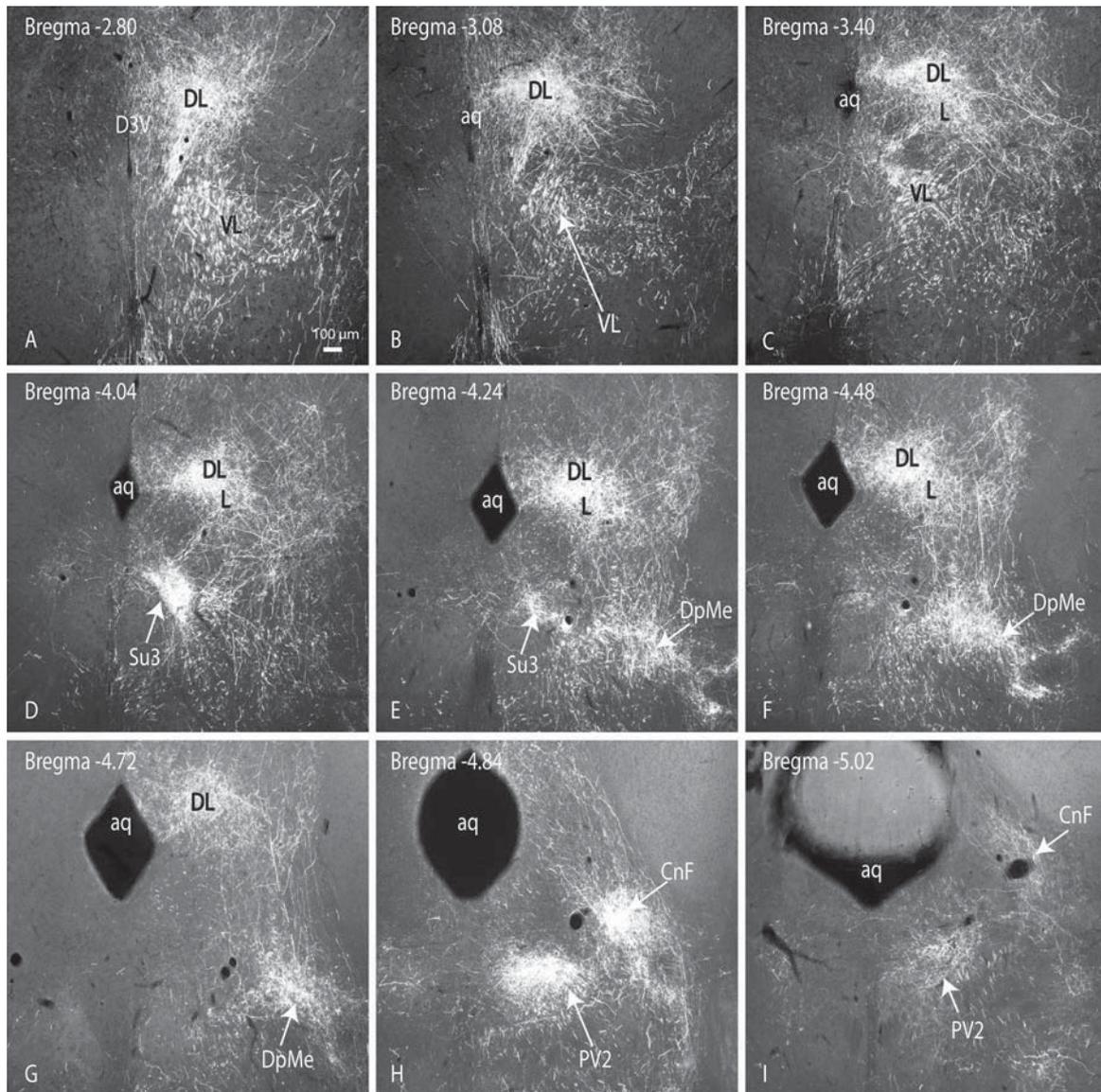
Figure 10. A–I: Epifluorescent images of nine consecutive coronal sections through the PAG, which was cut in a proximal (A)-to-distal (I) direction (E–G: Bregma ~ –4.20 mm to Bregma ~ –4.72 mm; H,I: Bregma ~ –5.00 mm). The images illustrate the projections of the lateral hypothalamic Foxb1-expressing neurons to the PAG. The coarse, AAV-EGFP-labeled axons radiate dorsally toward the rostral PAG and arborize in its dorsolateral (DL) and upper lateral portions (L), as well as in the deep layers of the superior colliculus (see Fig. 8). More caudally (D), the projections to the Su3 become visible. At Bregma ~ –4.04 mm (D), the axons separate into two bundles, which target the dorsolateral and the supraoculomotor (Su3) regions (E–G) and also populate the deep mesencephalic nucleus (DpMe). The terminal field in the dorsolateral region vanishes at level H and is replaced by one in the cuneiform nucleus (CnF; H,I). The ventrolateral terminal field is interrupted at levels F and G and reappears at level H, namely, in the PV2 nucleus (injection 194/13; the same animals as in Figs. 4, 11, and 12). Scale bar = 100 μm.

the septum, and the BNST; dorsally to the thalamus and the rostrolateral hypothalamus; and caudally to the superior colliculus, the PAG, the cuneiform nucleus, the pedunculopontine tegmental nucleus, the mesencephalic reticular formation, the gigantocellular nucleus, the retrofacial nucleus, and the retroambigual nucleus (Fig. 17). The extensive Foxb-expressing projections to many cell groups in the neural axis form the neural substrate for a variety of possible autonomic and behavioral functions. Compared with the projections of the Parv-immunopositive neurons in the parvafox nucleus (Celio et al., 2013), those of the Foxb1-expressing ones spread more widely.

Taking the findings of other investigators on the connections of the lateral hypothalamus as a whole (Berk and Finkelstein, 1982; Holstege, 1987; Hosoya and
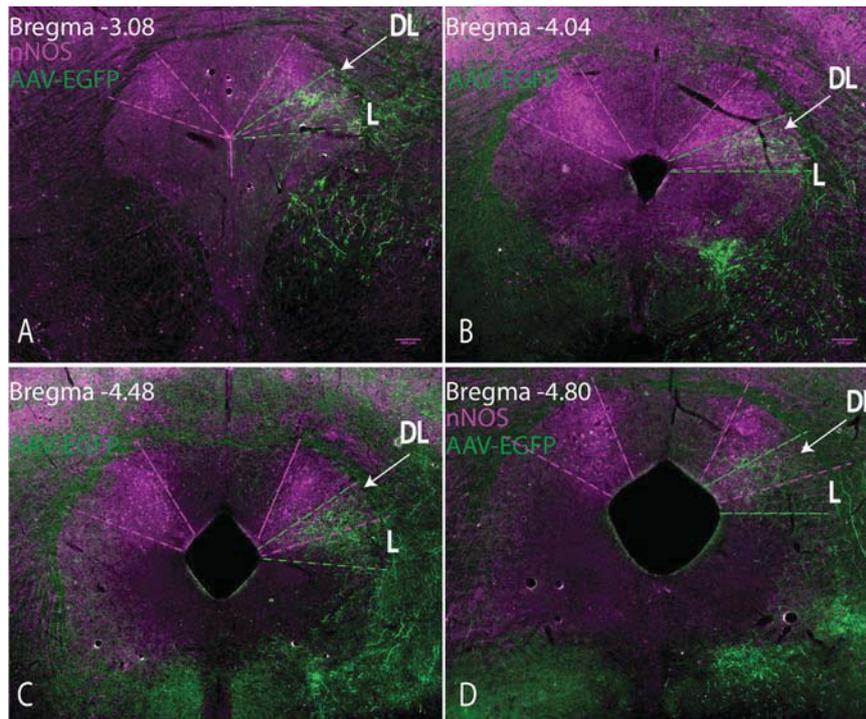
Figure 11. A–D: Epifluorescent images of four consecutive coronal sections through the PAG, which was cut in a proximal (A)-to-distal (D) direction (A,I: Bregma ∼ −3.08 mm to Bregma ∼ −4.80) of animal 25/16 injected in the parvafox of Foxb1-Cre mice stained for nNOS. The image illustrates the partial overlap of the projection from the lateral hypothalamic Foxb1-expressing neurons with neurons expressing the nitric oxide synthase (nNOS). The AAV-EGFP-labeled axons radiate dorsally toward the rostral PAG and arborize in its dorsolateral (DL) and upper lateral portions. The nNOS-positive neurons immunolabeled with a magenta fluorescent marker define the DLPAG. The magenta dashed lines indicate the approximate boundaries of the nNOS-positive cells, and the green dashed lines the boundaries of the terminal fields. The region included between the two lines in the middle corresponds to the region in which nNOS staining and the Foxb1-terminals coexist (injection 25/16). Scale bar = 100 μm.

Matsushita, 1981; Kita and Oomura, 1982; Saper et al., 1979; Veening et al., 1987; Villalobos and Ferssiwi, 1987a,b; Wolf and Sutin, 1966), the projections of the Foxb1-expressing cells are more confined. Most striking is the absence of projections to the amygdala, the parabrachial region, the locus coeruleus, the raphe nuclei, the dorsal vagus, and the solitary nucleus, which are the prominent target of other neuronal subfields in the lateral hypothalamus (Hahn and Swanson, 2010, 2012, 2015), even of the population of hypocretin/orexin (Peyron et al., 1998)- and melanin-concentrating hormone (MCH; Skofitsch et al., 1985)-positive neurons.

However, the output of the parvafox nucleus and of other regions of the lateral hypothalamus shares in common projections to the PAG. In all reports relating to the outputs of the lateral hypothalamus, the PAG is consistently one of the targets (Berk and Finkelstein, 1982; Hahn and Swanson, 2010, 2012, 2015; Holstege, 1987; Hosoya and Matsushita, 1981; Pelosi et al., 2006; Saper et al., 1979; Veening et al., 1987; Villalobos and Ferssiwi, 1987b).

The projection of the Foxb1-expressing neurons in the parvafox nucleus is special in that it extends to both the dorsolateral (dL) and the upper half of the lateral (L) PAG: it does not respect the boundaries known from the classical columns (Bandler et al., 2000). The dLPAG and LPAG are known to play a role in mediating the defensive responses to interoceptive stimuli (e.g., visceral pain) as well as to social and existential threats (e.g., confrontation with a predator and hypoxia; Bandler and Keay, 1996; Canteras and Graeff, 2014; Gross and Canteras, 2012). The responses involve a coordinated regulation of cardiovascular and respiratory activities (Bandler et al., 2000; Chaitoff et al., 2012; Dampney et al., 2013; Iigaya et al., 2010; Johnson et al., 2012). The dLPAG comprises partially overlapping subpopulations of NADPH/nNOS-, calbindin D28k (Barbaresi et al., 2013; Paxinos et al., 2009)- and calretinin-expressing neurons. This PAG column is also the most effective stimulation (or lesioning) site for the enhancement (or suppression) of lordotic behavior and is the target of a heavy contingent of descending inputs from
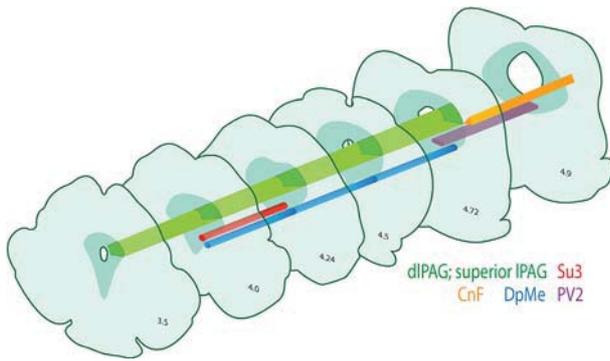
Figure 12. 3-D reconstruction of the terminal fields arising from the Foxb1-expressing neurons in the midbrain. The mouse PAG is represented in different rostrocaudal levels from Bregma −3.5 mm to Bregma −4.9 mm. In green is pictured the column of endings that reaches the dorsolateral (dlPAG) and the lateral periaqueductal gray (lPAG) from Bregma −3.5 mm to Bregma −4.72 mm. The red column shows the terminal fields to the supraoculomotor nucleus (Su3; Bregma −4.0 mm, Bregma −4.24 mm). The column of endings depicted in orange represents the terminals to the cuneiform nucleus (CnF; from Bregma −4.72 mm to −4.9 mm), the blue column the terminals at the level of the deep mesencephalic nucleus (DpMe), and the violet column the terminals on the PV2 (from Bregma −4.5 mm to −4.9 mm). Drawings modified from VanderHorst and Ulfhake (2006).

the hypothalamic ventromedial nucleus (Daniels et al., 1999). It also appears to harbor cells that are involved in vocalization (Larson, 1991). The lateral column of the PAG is also involved in predatory hunting and foraging behaviors (Comoli et al., 2003; Miranda-Paiva et al., 2003; Sukikara et al., 2006).

The LPAG sends projections to the lateral hypothalamus, especially at the tuberal levels, dorsolaterally to the fornix, in the region known to contain the MCH- and orexin-positive neurons (Bittencourt et al., 1992; de Lecea et al., 1998; Sakurai et al., 1998a). These two neural populations are involved in the control of locomotion (Krout et al., 2003) which is related to the food and water intake (De Parada et al., 2000). Therefore, on the basis of this physiological process, MCH and orexin are categorized as orexigenic neuropeptides (Qu et al., 1996; Sakurai et al., 1998a). However, the function of orexin and MCH is not limited to food intake. Orexin is also involved in arousal and foraging (Chemelli et al., 1999; Hara et al., 2001; Nishino et al., 2000): activation of the orexin-expressing neurons increases the probability of transition from sleep to wakefulness (Adamantidis et al., 2007).

The Su3 region is the only target that the Foxb1-expressing and Parv-immunopositive neurons (Celio et al., 2013) in the parvafox nucleus share in common.
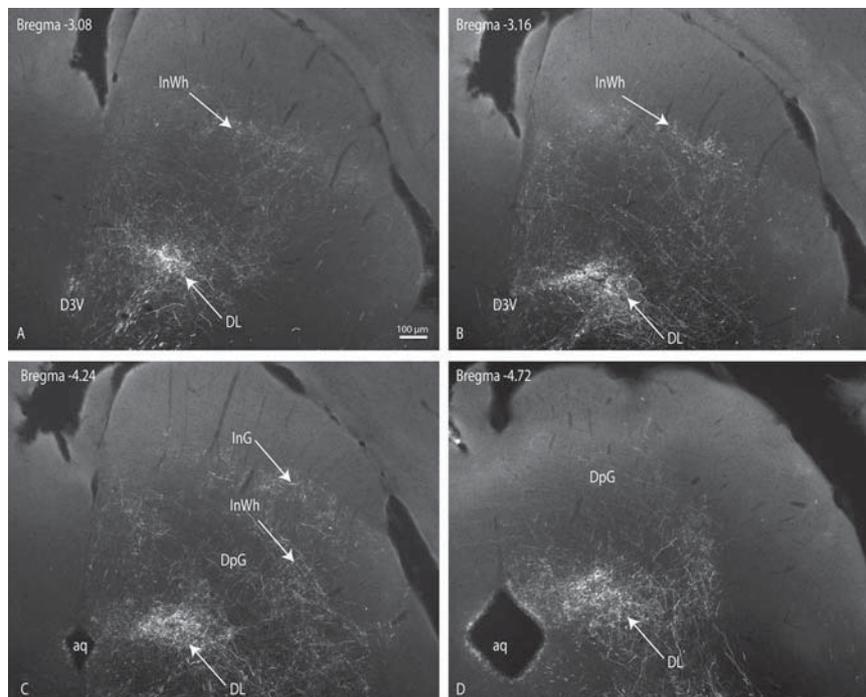
Figure 13. Epifluorescent images of coronal sections through the superior colliculus in a proximal (Bregma ∼ −3.08 mm; A) to distal (Bregma ∼ −4.72 mm; D) direction. A conspicuous bundle of fibers arborizes in the intermediate white layer (InWh). **A,B:** Proceeding caudally, at Bregma level ∼ −4. 24 mm, the fibers reach the intermediate layer of the superior colliculus (InG), the InWh, and the deep layer of the superior colliculus (DpG; **C,D**; injection 194/13; the same animals as in Figs. 1, 3, 5, and 11). Scale bar = 100 μm.
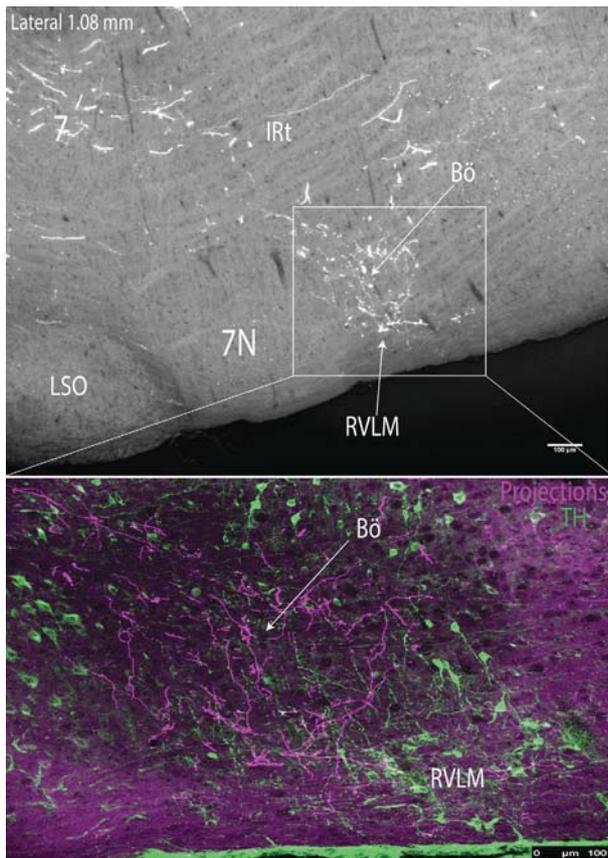
Figure 14. Epifluorescent image of parasagittal section trough the hindbrain (lateral 1.08 mm). The image illustrates the projections of the lateral hypothalamic Foxb1-expressing neurons to the rostral ventrolateral medulla. The **inset** shows an enlarged image of the projections (magenta) probably innervating the Bötzinger complex. In green the tyrosine hydroxylase-positive neurons are shown. Confocal image (injection 131/15). Scale bar = 100 μm.

No other recent publications relating to the hypothalamus report projections in the Su3, not even after stereotactic injections into targeted areas that lie slightly more medial to the parvafox nucleus, such as the hypothalamic tuberal nucleus (Canteras, 1992), the subfornical region (Goto et al., 2005) and the juxtaventromedial portion of the lateral hypothalamus (Hahn and Swanson, 2010, 2012, 2015). In the classical study conducted by Berk and Finkelstein (1982), which involved autoradiographic tracing with ^{3}H-leucine, terminals in a region that lay close to the Su3 were documented (injection M15) but were not commented on as being in any way noteworthy.

The Su3 receives inputs from the linear nucleus of the ventral tegmental region (Del-Fava et al., 2007), the intergeniculate leaflet (Morin and Blanchard, 2005), various oculomotor-related sites (Buttner-Ennever et al.,

2002), and the deep cerebellar nuclei (Gonzalo-Ruiz et al., 1990). A subpopulation of Su3 neurons (Van Bockstaele and Aston-Jones, 1992) projects to the rostral ventrolateral medulla (RVLM), a region that controls cardiovascular activity (Van Bockstaele et al., 1989, 1991) and to the nucleus ambiguus (Ennis et al., 1997). The neurons of the Su3 are activated when a predator is confronted and during hunting (Comoli et al., 2003).

The context-avoidance behavior of freezing rats and their response to predator-associated odors are accompanied by a marked elevation of the expression levels of c-Fos in the cuneiform nucleus (Allen et al., 1996; Carrive et al., 1997; Dielenberg et al., 2001b; Walker and Winn, 2007), which also plays a role in the cardiovascular and respiratory changes associated with defensive reactions (e.g., increases in heart rate and in blood pressure; Korte et al., 1992; Netzer et al., 2011). The cuneiform nucleus receives inputs from the anterior hypothalamic area and the dorsal premammillary nucleus (Motta et al., 2009) as well as from the lateral hypothalamic LHAjvd and LHAjvv (Hahn and Swanson, 2015), although those from the latter two regions are more scarce than those from the Foxb1-expressing neurons of the parvafox nucleus (Bernard et al., 1989; Netzer et al., 2011).

The superior colliculus is a laminated midbrain structure that acts to organize gaze movements, playing a role in orientation movements of the head and the eye in response to various stimuli (Dean et al., 1989). The superficial layers receive retinal information (May, 2006), and the deeper layers receive and integrate information from numerous sensory modalities (Drager and Hubel, 1975; May, 2006; McHaffie et al., 1989; Stein et al., 2009). The presence of a predator is registered in the lower visual field, outputs from which activate the medial deep layer of the superior colliculus, thereby triggering a signal-avoidance response. Appetitive stimuli are detected in the upper visual field, outputs from which activate the lateral deep layers of the superior colliculus (Comoli et al., 2012), whereby a signal-approach reaction is triggered (e.g., predatory hunting; Furigo et al., 2010). The lateral superior colliculus plays a role in the circuitry for sense-guided orientation decisions (Boehnke and Munoz, 2008; Dean et al., 1989; Felsen and Mainen, 2008) and integrates relevant sensory information for prey detection (Boehnke and Munoz, 2008).

The superior colliculus receives information from the ventromedial hypothalamic nucleus (VMH), the dlPAG, the vlPAG (Furigo et al., 2010; Mantyh, 1982; Wiberg, 1992), and the premammillary nucleus (PMV; Canteras and Swanson, 1992). It projects directly to the dlPAG (Rhoades et al., 1989) and, via this route, to the
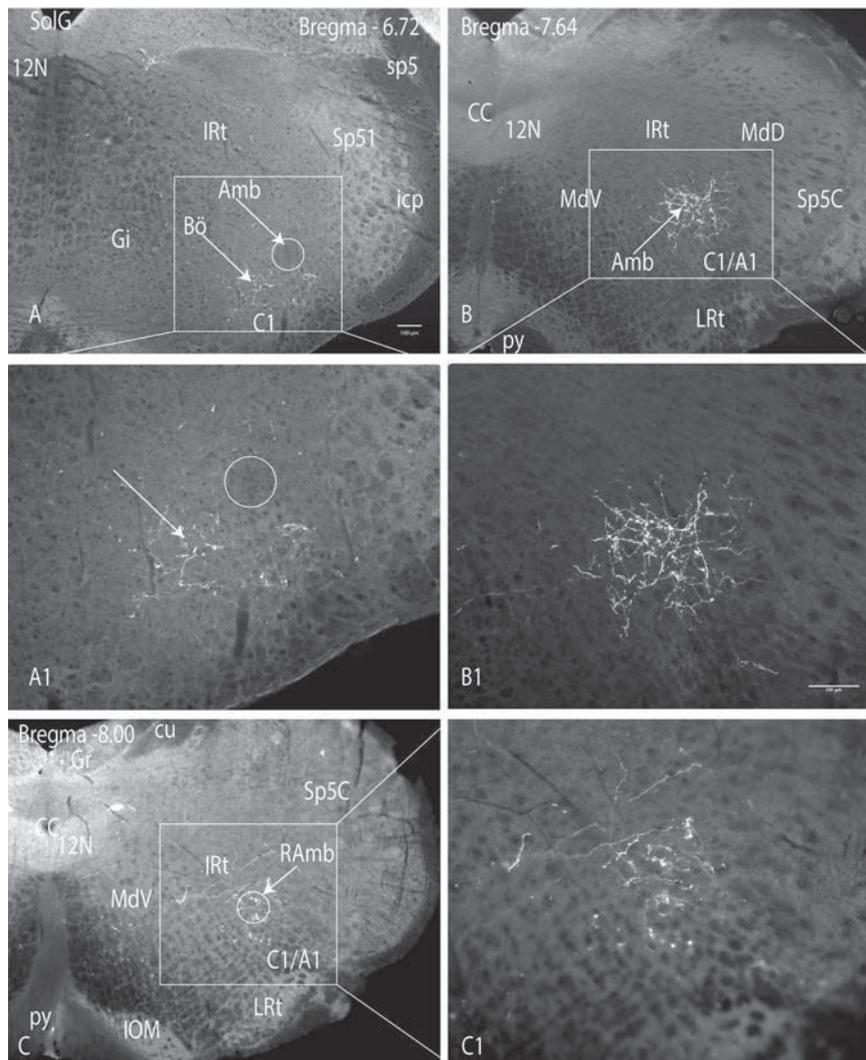
Figure 15. Epifluorescent images of a coronal section through the hindbrain. **A:** In the medulla oblongata, sparse tomato-positive fibers and terminals occupy a region ventral to the ambiguus nucleus (Amb), corresponding to the Bötzinger (Bö) complex (Bregma −6.72 mm). **A1:** Higher magnification the immunopositive axonal endings (arrows) in the Bö region. **B:** More caudally, the axons arising from the Foxb1-expressing neurons of the parvafox nucleus reach the region of the loose ambiguus (Bregma −7.64 mm). **B1:** Higher magnification of the rich ramification of immunopositive axonal endings in this region. **C:** At level Bregma −8.00, endings are found in the retroambigual region. **C1:** Higher magnification of the cluster of endings encircled in C. Scale bar = 100 μm.

cuneiform nucleus (Redgrave et al., 1988). The dlPAG and adjacent portions of the superior colliculus may thus form a part of a defense mechanism in the midbrain. The superior colliculus projects also to the lPAG (Furigo et al., 2010), which seems to be a nodal part of the circuitry involved in the decision-making process between different behavioral responses such as hunting or foraging (Sukikara et al., 2006).

Taken together, these data support the "sentinel hypothesis" (Merker, 1980) cited by R. Rieck et al. (1986). This postulate suggests that the superior colliculus, in particular, the medial deep layer, could function as a sentinel for dangerous stimuli and be involved in the behavioral responses that comprise the strategies of defense in a threatening situation (Carvalho-Netto et al., 2006; Dean et al., 1988), whereas the lateral deep layer controls orientation of pursuit-like movements and influences arousal, motivation, motor learning planning, and motor output which are critical for foraging and hunting behaviors (Furigo et al., 2010).

Other projections that may be implicated in defensive mechanism
Projections to the vestibular nuclei

The vestibular nuclei are involved in the control and regulation of blood pressure during movement and
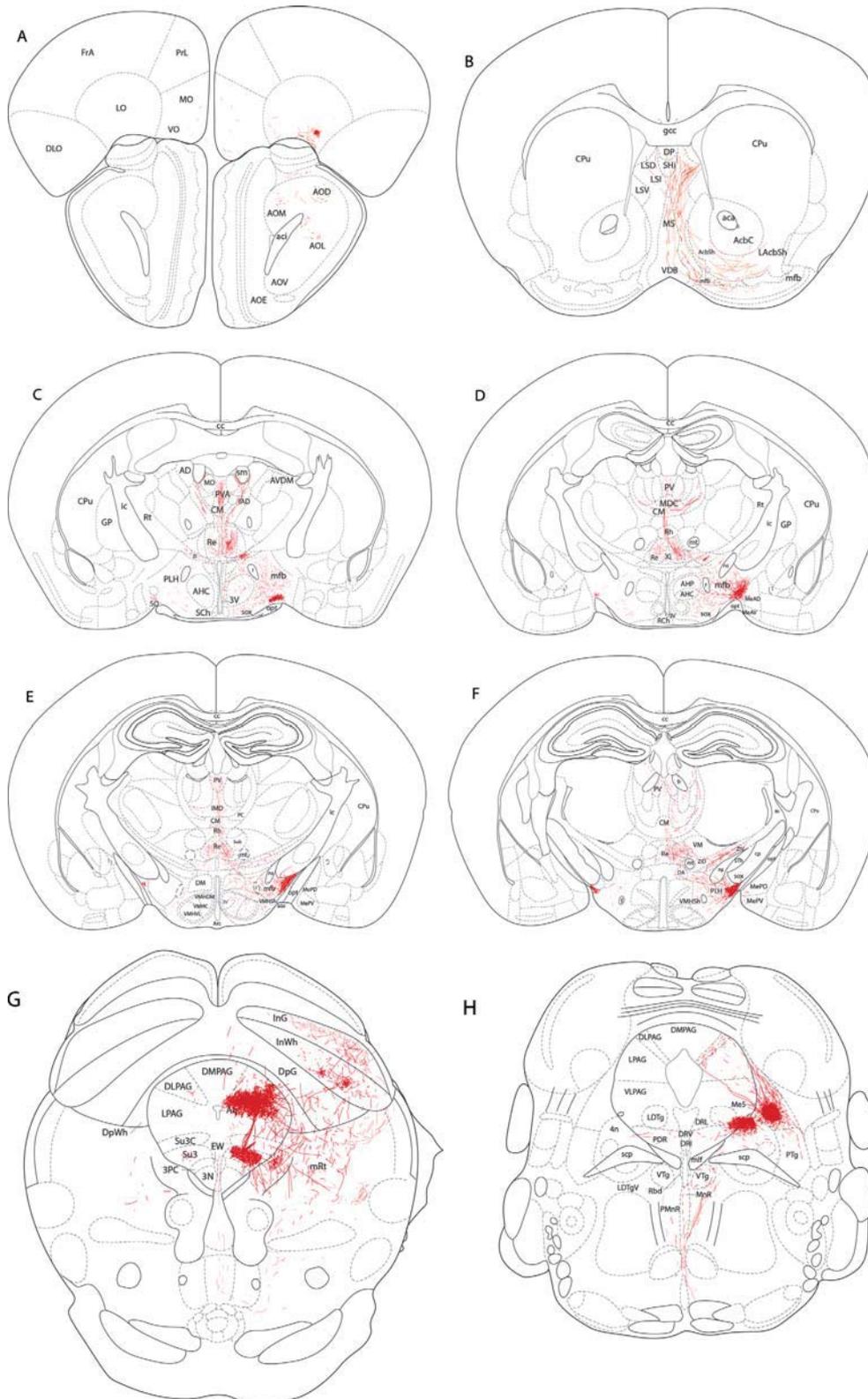
**Figure 16. I–H:** Series of coronal drawings at 8 hindbrain levels of the mouse brain in which the projections of the hypothalamic Foxb1 neurons are represented. The source of all these projections the virus-infected Foxb1 neurons seen in drawings D–F. The main targets in the dorsolateral and upper lateral PAG, as well as the Su3 and PV2 nuclei, are depicted in G and H. Note the absence of terminals in the amygdala, parabrachial nuclei, and solitary tract. Drawings modified from Franklin and Paxinos (2008).
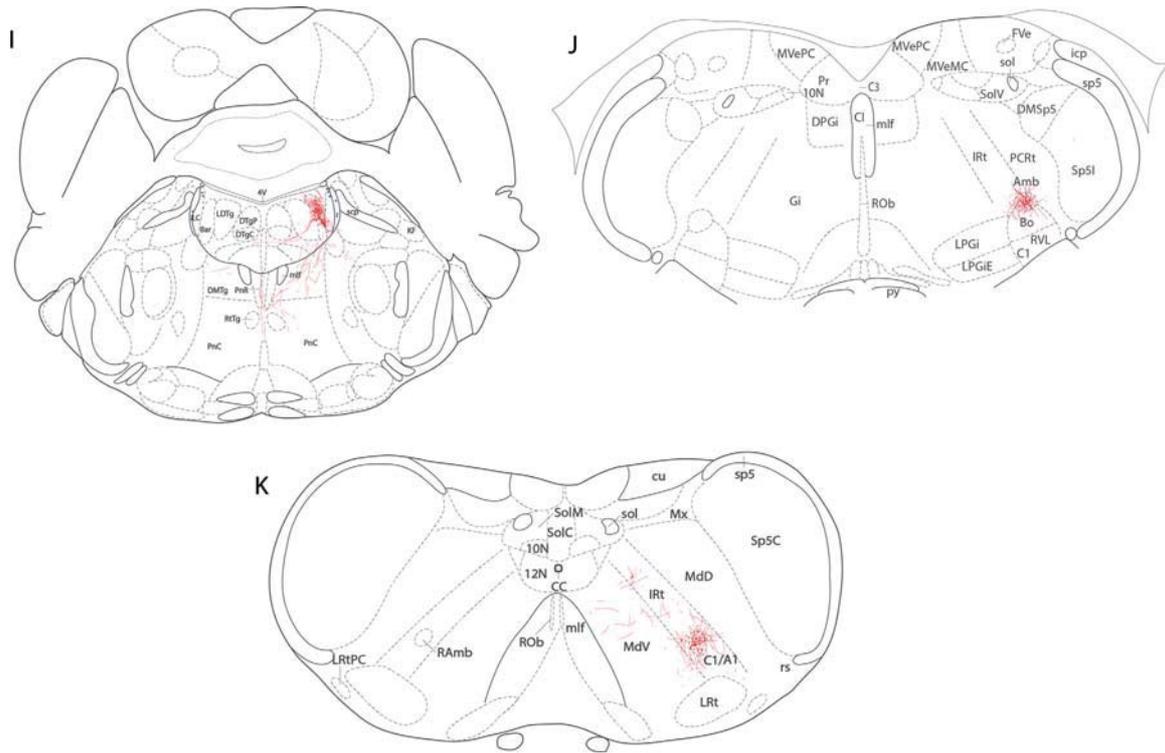
Figure 16. A–K: Series of coronal drawings at 10 levels of the mouse brain in which the projections of the hypothalamic Foxb1 neurons are represented. Drawings modified from Franklin and Paxinos (2008).

changes in posture via their influence on the sympathetic nervous system (Tang and Gernandt, 1969). Outputs from the cervical spinal tract project to the autonomic region of the vestibular nuclei. The reticular formation, including the lateral reticular nucleus (LRt), projects directly to the vestibular nuclei (Jian et al., 2005) and receives inputs from visceral and somatosensory receptors (Ammons, 1987; Blair, 1987; Drew et al., 1996; Fields et al., 1977; Peterson et al., 1976; Schulz et al., 1983; Thies, 1985; Yates and Stocker, 1998) as well as from the raphe nucleus (Blair and Evans, 1991; Knuepfer and Holt, 1991; Lumb and Morrison, 1984), whereby visceral and somatosensory information is integrated.

Projections to the gigantocellular and the lateral reticular nuclei

The gigantocellular reticular formation plays an important role in behavioral arousal (Boissard et al., 2002; Morest, 1961; Morest and Sutin, 1961; Steriade et al., 1984) as well as in the modulation of pain (Mason, 2005; Nagata et al., 2003) and motor activity (Fay and Norgren, 1997; Hattox et al., 2002; Robinson et al., 1994). It receives projections from diverse sources, including the nucleus of the solitary tract, the preoculomotor

nucleus, the Su3, the PAG, the lateral hypothalamus, the periventricular nucleus, and the medial prefrontal cortex (Van Bockstaele et al., 1989). The main targets of the gigantocellular nucleus are the locus coeruleus and the spinal cord (from the gigantocellualr reticular formation; Hermann et al., 2003; Holstege and Kuypers, 1982; Liang et al., 2013).

The LRt is a precerebellar agglomeration of mossy fibers, which relays information to the cerebellar cortex. It receives a massive contingent of afferents from the spinal cord (Brodal, 1949; Kunzle, 1973; Morin et al., 1966) as well as descending fibers from the cerebral cortex (Brodal et al., 1967; Kunzle and Wiesendanger, 1974), the red nucleus (Mizuno et al., 1973), and the superior colliculus (Mizuno et al., 1973). The lateral reticular formation projects a massive bundle of fibers to the cerebellum, which processes the inputs it receives from the different areas. Consequently, only filtered information appertaining to control of movements is delivered to the cerebellum (Alstermark and Ekerot, 2013).

Projections to the olfactory bulb

Common stimuli that trigger specific behavioral responses are the presence of a predator or prey. In both instances, sentience of the presence is registered
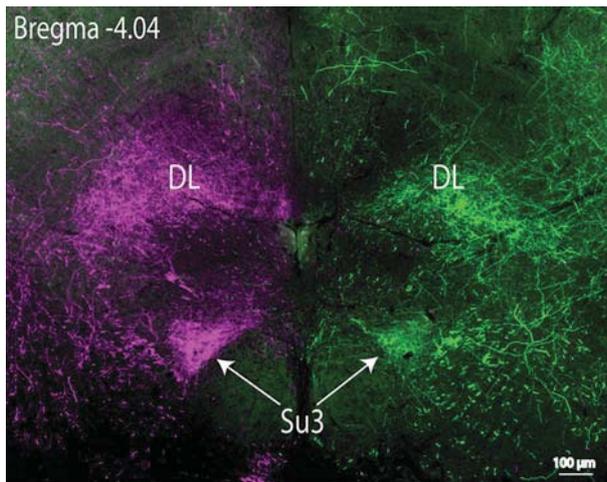
Figure 17. Epifluorescent image of projections in the PAG at Bregma level −4.04 mm after a bilateral injection of two different viral tracers, one bearing dTomato and the other EGFP in the parvafox nucleus. The projections are symmetrical (injection 9/15). Scale bar = 100 μm.

in the visual or the olfactory system. An odor is detected by the olfactory system, from which a bundle of fibers runs consecutively to the amygdala, the ventromedial hypothalamic nucleus (VMH), the lateral hypothalamus and the dlPAG, which together form a specific circuitry (Canteras, 2002; Dielenberg et al., 2001b; Motta et al., 2009).

Projections to the prefrontal cortex

The prefrontal cortex mediates the autonomic responses to emotional situations. Its projections to the central nervous system influence cognitive processes and decision making. It is implicated in the control of hearth rate, blood pressure, respiration, and gastric motility (Diorio et al., 1993; Feldman and Conforti, 1985; Gabbott et al., 2005; Van Eden and Buijs, 2000). The prefrontal cortex plays a pivotal role in mediating the interactions between the central and the autonomic nervous systems. The ventromedial portion projects to the hypothalamus (PVN, DMH, LH) and the dorsomedial one directly to the autonomic centers in the brainstem and spinal cord. The former portion projects to the vlPAG and the latter to the dlPAG (Van Eden and Buijs, 2000).

Projections to the lateral septum

The lateral septum is a sexually dimorphic nucleus, which is the target of vasopressin-expressing innervations. It is heavily interconnected with sites in the lateral hypothalamus that are implicated in the control of rewarding, stimulus-driven states (de Vries et al.,

1981). It is also involved in coping with stress (Koolhaas et al., 1998) as well as in regulating social behavior and anxiety (Veenema and Neumann, 2008). Activation of the CRF2 receptor-expressing subpopulation of neurons in the lateral septum generates anxiety (Anthony et al., 2014).

Projections to the midline nuclei of the thalamus

The thalamic midline nuclei (PV, PVA, CM, IAM, and Rh) harbor neurons that project to the nucleus accumbens, the medial frontal cortex, and the amygdala (Bubser and Deutch, 1998; Otake and Nakamura, 1998; Su and Bentivoglio, 1990); they are involved in viscerolimbic functions and in the self-perception of viscerosensory stimuli (Cassel et al., 2013; Otake and Nakamura, 1998; Van der Werf et al., 2002).

Comparison of our findings with existing data on the defensive responses

The independent, multiple circuits that process the different types of defensive reaction involve the medial, but not the lateral, hypothalamus (Gross and Canteras, 2012; Jordan, 1990). In the classical work by Ranson's group, the defense responses was localized lateral to the fornix (Kabat et al., 1935), whereas the observation of Hess and his school (Akert, 1981; Fernandez De Molina and Hunsperger, 1962; Hess and Brügger, 1943; Hunsperger, 1956) implicated the perifornical and pericommissural (Gansser) regions, a finding that has since been confirmed in several studies (Nakao, 1958; Wasman and Flynn, 1962). The defensive response to the presence of a predator involves the dorsal premammillary nucleus (Canteras, 2002). Recent optogenetic work in mice has localized the center of an intermale attack to the ventrolateral subdivision of the ventromedial nucleus (Lin et al., 2011) and a stalking one to the lateral portion (Wasman and Flynn, 1962).

A participation of this most recondite portion of the lateral hypothalamus, which harbors the Foxb1-expressing neurons, in the stereotyped behavioral aspects of a defensive response has never been reported. Electrical stimulation of this zone in the cat (Hess, 1947, 1957; Karplus and Kreidl, 1910; Ranson and Magoun, 1933) leads to changes in cardiovascular, respiratory, and pupillary activity as well as to salivation, micturition, and defecation. The cardiovascular effects have been confirmed by the microinjection of L-glutamate, which activates neurons but not the axons of passage (Allen and Cechetto, 1992; Spencer et al., 1989). It may therefore be deduced that the parvafox nucleus is concerned not with the behavioral but with

the autonomic manifestations of the defense response. It is to be expected that electrode recording or chemo- and optogenetic techniques will help to establish the role played by this nucleus in various threatening, stressful, and other emotional situations.

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CONFLICT OF INTEREST STATEMENT

The authors declare that they have no conflicts of interest.

ROLE OF AUTHORS

All authors had full access to the data in the study and take responsibility for its integrity and the accuracy of the analysis. MRC designed and oversaw the project. GA-B supplied the Foxb1 mice; AB made the stereotactic injections, prepared the sections and performed immunohistochemistry. AB and MRC took pictures, evaluated the results, and together with GA-B wrote the manuscript.

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