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## Advances in the neurobiological bases for food ‘liking’ versus ‘wanting’

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### HIGHLIGHTS

- We describe the anatomical localization of three “hedonic hotspots”.
- A description of how the hotspots functionally interact
- A proposed hedonic circuit that unites the hotspots

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### ABSTRACT

The neural basis of food sensory pleasure has become an increasingly studied topic in neuroscience and psychology. Progress has been aided by the discovery of localized brain subregions called hedonic hotspots in the early 2000s, which are able to causally amplify positive affective reactions to palatable tastes (liking) in response to particular neurochemical or neurobiological stimulations. Those hedonic mechanisms are at least partly distinct from larger mesocorticolimbic circuitry that generates the incentive motivation to eat (wanting). In this review, we aim to describe findings on these brain hedonic hotspots, especially in the nucleus accumbens and ventral pallidum, and discuss their role in generating food pleasure and appetite.

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### Q2 1. Introduction

Over the last 15 years, research has yielded several unexpected findings on how hedonic circuitry in the brain interacts with food to produce reward and appetite. Evidence now suggests that discrete, anatomically localized “hedonic hotspots” exist in limbic-related brain structures, and are able to magnify the hedonic impact of natural sensory rewards, such as sweet tastes. So far, these hotspots have been found in the forebrain nucleus accumbens (particularly in medial shell), ventral pallidum, and in the brainstem parabrachial nucleus. In this review, we will discuss where these hotspots were found, what neurochemical systems enhance hedonic impact in them, and how the hotspots may interact within hedonic circuitry and with a larger mesocorticolimbic circuitry that produces appetite or the motivation to eat.

#### 1.1. Nucleus accumbens hotspot

##### 1.1.1. The striatum

The nucleus accumbens (NAc), as well as the striatum as a whole, is well known to be involved in reward and motivation. However, it has also become increasingly clear that subregions within the nucleus accumbens and striatum can differently influence distinct aspects of behavior and motivation [6,39,84,134]. One potential contributing factor may be related to the anatomical make up of different zones within the striatum. For example, though there are general striatal neurobiological features shared by NAc and neostriatum (D1/Dynorphin and D2/Enkephalin descending projections, inputs from prefrontal cortex, amygdala, and hippocampal nuclei, etc.), there are also clear anatomical differences between ventral and dorsal striatum, between core and shell components within nucleus accumbens, and even between different subregions within the medial shell of the nucleus accumbens [52,61,73,117,133].

##### 1.1.2. Affective taste reactivity as a tool to measure hedonic function

The taste reactivity test can be used as an objective measure of hedonic impact or ‘liking’ reactions to taste palatability, based on

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quantifying discrete orofacial affective reactions to different tastes [112]. Originally applied to rats in behavioral neuroscience studies by Grill and Norgren for use in decerebrate and thalamic rats [48,49], this affective reactivity test was even earlier pioneered in human infants [111]. Converging evidence from animal and human comparisons showed that the orofacial reactions elicited by rats and humans (as well as several species of apes, monkeys, horses and mice), in response to palatable or unpalatable tastes, are strikingly homologous, with positive hedonic 'liking' reactions including tongue protrusions, lateral tongue protrusions and paw licks, and negative 'disgust' reactions including gapes, head shakes, and chin rubs [62,112]. 'Liking' and 'disgust' are placed in quotation marks to acknowledge that these are objective positive or negative hedonic reactions that are not necessarily accompanied by subjective feelings of pleasure or disgust (even if they often are) [94,125], and to distinguish them from the everyday use of the English term, liking. Similarly, 'wanting' in quotes refers specifically to the motivation process of incentive salience, which also can occur in brain and behavioral responses either with or without accompanying subjective feelings of ordinary wanting [94,94,125].

While at first it seemed possible that these taste-elicited reactions were merely sensory-specific reactions (e.g. sweet versus bitter), or merely brainstem reflexes rather than affective responses (taste reactions are emitted by decerebrates with only a brainstem to control behavior [47,49]), accumulating studies suggested that the orofacial reactions truly reflected hedonic impact for intact-brain individuals by the 1980s. For example, initially 'liked' tastes, such as sugars or saccharin, after being paired with injections of lithium chloride to produce a conditioned taste aversion (CTA), subsequently produced aversive gapes, which requires forebrain control [12,49,82,109,123]. Reciprocally, intraoral infusions of a normally disgusting hypertonic NaCl solution (e.g., 1.5 M) can produce hedonic reactions in a salt depleted state [19, 33,93,118]. Further, affective orofacial reaction patterns are not tied to particular sensory stimuli in any one-to-one fashion that would reflect sensory-specific coding; palatable sucrose, palatable NaCl at isotonic or hypotonic concentrations, and palatable fat emulsions can all evoke similar hedonic reactions [33,102,104]. Further, the affective taste reactivity pattern elicited by a particular taste can be altered by factors that also alter human palatability ratings, ranging from relevant appetite/satiety physiological states, to pharmacological opioid, endocannabinoid, etc. brain states of particular neuroanatomical structures, and types of neurobiological lesions [14,26,35,69,75,84,131]. Finally, specific brain microinjections, lesions, or optogenetic stimulations in forebrain structures can profoundly control taste-elicited 'liking' reactions as described below, which indicates a top-down or hierarchical control over brainstem circuitry that involves the entire brain. Altogether, these considerations indicate that the taste reactivity test reflects the affective (sensitive to homeostatic and learned cues), rather than merely a reflex or the sensory quality of a food reward.

### 1.1.3. The nucleus accumbens hedonic hotspot

In an effort to uncover the neural mechanisms of hedonic processing, taste reactivity has been used in conjunction with brain manipulations, such as pharmacological microinjections in particular structures. Using this coupled paradigm, Susana Peciña in the Berridge lab was able to demonstrate that a unique hedonic function was localized to a subregion of the NAc medial shell; a 1<sup>3</sup> mm "hedonic hotspot" in the rostradorsal quadrant of the NAc medial shell [84]. Within the confines of the cubic-millimeter hotspot in shell, mu opioid receptor activation via microinjection of the mu agonist DAMGO [71] enhanced hedonic 'liking' reactions to a sweet sucrose solution, in addition to suppressing negative 'disgust' reactions to quinine [84,106]. Within the NAc hotspot, mu opioid stimulation was found double to triple the number of positive orofacial 'liking' reactions elicited by sweetness, in addition to dramatically stimulating intake of palatable food.

Outside the hotspot, the same opioid stimulation completely failed to increase 'liking' reactions, even though it increased intake just as

much. In fact, at posterior locations in the medial shell, opioid stimulation tended to oppositely suppress 'liking' reactions in a hedonic coldspot. However, at all sites in accumbens core and shell, DAMGO microinjections are equally effective at stimulating increases in food intake and in 'wanting' to obtain food, despite not enhancing 'liking' at most of those sites [7,84,85,134]. Indeed, food intake can be stimulated at a number of related sites outside the NAc, without enhancing 'liking' reactions, including the central nucleus of the amygdala [46,68] and even regions of the ventral and dorsal neostriatum [39,134]. Thus, opioid circuitry for 'wanting' to eat is more widely distributed throughout the NAc and related structures than opioid circuitry for 'liking'.

More recently, we have replicated the original mu opioid hotspot localization in the rostradorsal quadrant of the NAc shell for enhancements of sucrose 'liking' by DAMGO microinjections [30] (Fig. 1). Further, we have found evidence that the same anatomical site for the rostradorsal mu hotspot can mediate opioid hedonic enhancements for delta stimulation (DPDPE) by three-fold and even kappa stimulation (U50488H) by two-fold, whereas no hedonic enhancements are produced at other locations in the medial shell of NAc by either mu, delta or kappa stimulations [30] (Fig. 1). Oppositely instead, in a hedonic coldspot in the posterior half of the medial shell, all three forms of opioid stimulation suppress hedonic reactions to sucrose, apparently reducing 'liking'. However, each specific agonist had different anatomical patterns of effects on 'wanting' to eat in the sense of changing food intake despite their similar (rostral) enhancement hotspot versus (caudal) suppressive coldspot pattern of 'liking' effects [30] (Fig. 1). Mu stimulation increased eating at all sites throughout the medial shell, as previously reported, both in the caudal coldspot and the rostral hotspot. However, delta stimulation only increased eating in the rostral hotspot but not at other sites, and kappa stimulation never consistently increased food intake at any site in the medial shell. These differences speak again to the fundamental differences in mechanisms mediating 'liking' versus 'wanting', even within opioid systems contained in the medial shell of NAc. Related evidence has demonstrated endocannabinoid stimulation in the NAc hotspot and even GABAergic hyperpolarizations in the same hotspot can also enhance 'liking' reactions to sweet tastes [43,69,84,91,92].

All three types of opioid receptors couple to Gi subunits, subsequently leading to ERK activation and typically decreasing neuronal activity, which conceivably could be related to shared enhancement effects in the rostral hedonic hotspot and shared hedonic suppression effects in the caudal coldspot. However, though mu, delta and kappa pathways converge to activate ERK, they do so via different intracellular channels, which might possibly be relevant to how the three receptors have such different effects on motivated 'wanting' to eat reflected in food intake. However, the precise relation between intra-cellular mechanisms and 'liking'/'wanting' effects still remains to be clarified.

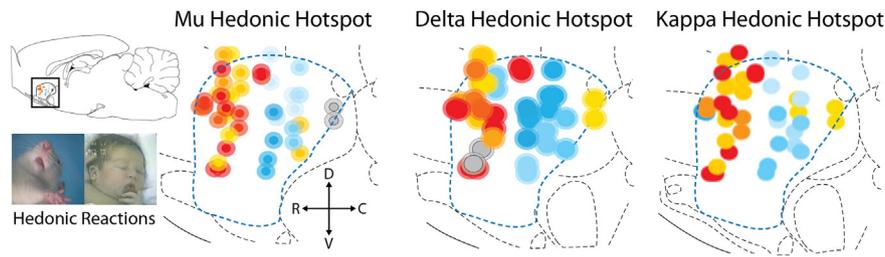
### 1.1.4. Dopamine fails to alter taste reactions

By contrast to hedonic neurochemical manipulations, NAc dopamine stimulation by amphetamine microinjections within or outside the shell hotspot [106,129], by genetic elevation of dopamine in the synapse (via knockdown of dopamine transporter in presynaptic dopamine neurons) [87], or by systemic amphetamine administration [119,121], all consistently fail to enhance positive hedonic reactions to sweet tastes. Conversely, reduction of NAc dopamine by 6-OHDA lesions [17,18], or by systemic dopamine blockade [86] all fail to reduce positive hedonic reactions. However, those same dopamine manipulations do potently alter motivated 'wanting' for the food rewards. Thus, unlike opioid or endocannabinoid neurotransmitters, dopamine in the NAc does not appear to be a mechanism for hedonic 'liking', but rather is restricted to motivation 'wanting' roles regarding food rewards.

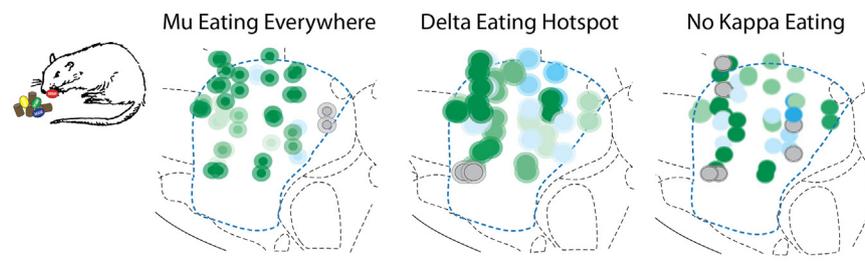
### 1.1.5. Anatomical basis for functional uniqueness of NAc hotspot

What anatomical basis might help explain the functional existence of an anatomically unique hotspot for opioid/endocannabinoid

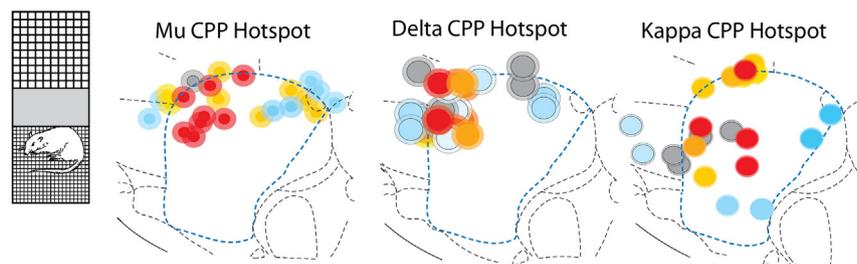
## Opioid Hedonic Hotspot in Nucleus Accumbens Enhances 'Liking'



## Opioid Stimulation Differently Enhances Food Intake



## Hotspot Opioid Stimulation Induces a CPP



**Fig. 1.** *Mu, delta or kappa* opioid hotspots for 'liking' enhancements in the nucleus accumbens. Top row: 'liking' reactions to sweetness. All three types of opioid signaling mechanisms share essentially the same hedonic hotspot in NAc medial shell. Activation of any of the three types of receptor (*mu*, *delta* or *kappa*) enhances hedonic 'liking' reactions to sucrose taste within the same rostral cubic millimeter hotspot. Conversely, all three opioid stimulations suppress hedonic reactions in a caudal coldspot in medial shell. Each circle represents a single microinjection site. Yellow to red colors indicate increases in positive 'liking' reactions caused by opioid stimulation, and gray to blue indicate suppression 'liking' to below normal control levels in the same individual rat. Middle row: different effects on 'wanting' to eat food. Stimulation of the three receptor subtypes have very different effects on food intake, highlighting that there are differences in the opioid neural mechanisms mediating 'liking' and 'wanting'. Gray to green symbols indicate increases in food intake, and gray to blue indicate a suppression of food intake. *Mu* stimulation enhanced food intake at all sites throughout the entire NAc. *Delta* stimulation enhanced eating only within the hotspot (similarly to 'liking' enhancement). *Kappa* stimulation never consistently enhanced eating at any anatomical site. Bottom row: confirmation of hotspot identity via place preference conditioning. Conditioned place preferences are an independent way of measuring reward, which turns out to confirm that the rostrrodorsal quadrant of medial shell is unique for opioid reward effects. Stimulation of either *mu* or *kappa* receptors within the hotspot also generated a conditioned place preference (and *delta* showed a similar trend), whereas no preference was induced at other sites in medial shell. Yellow to red symbols indicate a positive place preferences, and gray to blue symbols indicate induction of negative place avoidances. Modified by permission from [30]. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

200 amplification of sensory pleasure, and why is it uniquely able to enhance hedonic impact to tastes, compared to other regions of the NAc shell?

203 Recently, two independent groups of neuroanatomists have evaluated the anatomical connectivity patterns of the NAc rostrrodorsal quadrant of medial shell, and found that this hotspot region differs from other subregions of the medial shell (e.g., caudal shell). Thompson and Swanson [117] revealed, using a double injection of anterograde and retrograde tracers, that the rostrrodorsal quadrant appears to belong to a different striato-pallido-hypothalamo-thalamo-cortical closed circuit loop from other subregions of the medial shell. In other words, if one follows the projections from the rostrrodorsal quadrant of the medial shell along a point to point axis, one will end up back in the hotspot. This loop travels from the NAc hotspot to particular subregions of pallidum or hypothalamus, up to paraventricular nucleus of the thalamus, next passing through the infralimbic region of the prefrontal cortex, and finally projecting back again to the rostrrodorsal medial shell. The subregions of each of these structures are distinct from the subregions visited by other parallel loops that pass through more posterior regions of the

219 medial shell. Exactly how many parallel loops pass through the medial 220 shell of NAc remains to be elucidated, but it seems clear now that 221 there are at least two (visiting rostral vs caudal shell) and possibly additional loops that more finely dissect NAc shell into further subregions, each belonging to its own loop [117].

224 Similarly, Zahm and colleagues [133] recently found a related pattern of distinct connectivity that distinguishes the rostral hotspot from more caudal subregions of NAc medial shell. Those authors suggest that the rostral hotspot projects to particular regions of the lateral preoptic area and lateral hypothalamus, and receives inputs from infralimbic (analogous to Brodmann's area 25) and other nearby regions of prefrontal cortex such as prelimbic and orbitofrontal cortex. They also suggest that the projection patterns of NAc rostral shell are similar to those of lateral septum, compared to the caudal shell, and that the rostral zone of medial shell is a unique transition region between NAc and lateral septum. In contrast, they suggest that the caudal zone is a different transition region blending features of NAc and extended amygdala. While the Zahm et al. and the Thompson and Swanson studies differ on some points, the overall anatomical scheme

presented by the two studies seems to agree that the circuitry belonging to the rostromedial hotspot quadrant of the NAc medial shell is fundamentally different compared to the connectivity patterns of the rest of the medial shell, and that these anatomical differences may in part contribute to the hotspot's unique abilities to amplify the hedonic impact of taste sensations.

In addition to differences in projection patterns, there may also be other local neurobiological features of neurons in the NAc medial shell that are relevant to hedonic contributions compared to other NAc components such as core. Meredith et al. [73] suggest that the local characteristics of neurons in the NAc medial shell are different from other regions of NAc and striatum. For example, the projecting medium spiny neurons (MSNs) within the medial shell are less spiny and smaller compared to NAc core or dorsal striatum. Furthermore, the distinction between different MSNs belonging to D1/dynorphin/direct pathway versus D2/enkephalin/indirect pathway, which is known from dorsal striatum, is somewhat diluted in the NAc medial shell, where at least 17% of MSNs harbor both D1 and D2 receptors [22,61]. Intriguingly, volume ratios of patch/matrix compartments in dorsal striatum (as delineated by mu opioid or calbindin binding) may also be flipped, or at the very least are not as cleanly split in the nucleus accumbens [63,72]. Although the roles of these neurobiological features is still unclear, some of these unique anatomical or cellular features of the NAc medial shell might be relevant to its ability to generate hedonic functions that are fundamentally different from other regions of striatum.

## 1.2. Ventral pallidum hotspot

### 1.2.1. Evidence for a ventral pallidum hotspot

The ventral pallidum (VP) receives the densest projections from NAc, compared to other target structures [77,78]. Similar to NAc, VP also has been shown to be important for rewards [35,70,74,104,110,114,115,130]. Also similar to NAc, the VP has been shown to contain a hedonic hotspot of its own [104].

In an initial microinjection mapping study of the VP hedonic hotspot, Kyle Smith in the Berridge lab made microinjections of DAMGO throughout the ventral pallidum and measured taste reactivity responses to sucrose and quinine, as well as changes in food intake [104]. Results showed that DAMGO microinjections in a roughly cubic-millimeter site of caudal VP enhanced hedonic reactions to sucrose, revealing a hedonic hotspot in the posterior half, as well as stimulating the motivation to eat more food. In behavioral and anatomical contrast to the posterior VP hotspot, microinjections into more rostral subregions of VP suppressed 'liking' reactions to sucrose and reduced food intake, indicating a VP opioid coldspot [104]. The caudal VP zone which enhanced hedonic reactions was slightly smaller (~0.8<sup>3</sup> mm) than the 1<sup>3</sup> mm NAc hotspot, although it is proportionally similar to the NAc hotspot when the relative size of the structures are taken into account. Thus, like NAc, the VP also appears to house a hedonic hotspot (but positioned caudally in VP, rather than rostrally as in NAc).

### 1.2.2. An orexin hotspot in VP

In addition to opioid signals, orexin signals in the posterior VP also can enhance the hedonic impact of sucrose [60]. This was found by performing microinjections of orexin-A directly into the VP hotspot or into the surrounding regions of lateral hypothalamus (lateral preoptic area) or into the extended amygdala. Chao-Yi Ho in the Berridge lab found that orexin microinjections enhanced 'liking' reactions when infused into the VP hotspot, but did not do so when infused into rostral ventral pallidum or into nearby structures such as the lateral hypothalamus or extended amygdala [60]. Whether or not orexin also acts in the NAc hotspot to enhance hedonic impact is still unknown, but preliminary observations in our lab suggest that orexin may also perform a similar role in this NAc region as well (Castro and Berridge, unpublished observations).

### 1.2.3. Necessity of the VP hotspot

During the 1960s and 70s, it was reported that lesions to LH would produce intense aphagia [23,80,99,116]. In particular, Teitelbaum and Epstein [116] reported that LH lesions, in addition to disrupting eating and drinking behavior, also disrupted hedonic/appetitive reactions to sweet solutions and replaced them with aversive or 'disgust' reactions, which suggests a role for LH in affective processing and behavior. However, with the benefit of hindsight, it can be noted that those hypothalamic lesions were very large by modern standards, and the damage actually extended well outside the lateral hypothalamus. Additional structures were damaged, ranging from caudal ventral pallidum in a direction anterior to LH, and as far back as premammillary nucleus in a caudal direction. Subsequently Schallert and Wishaw [98] identified the anterior direction as most important, showing that electrolytic lesions only in anterior LH produced intense 'disgust' reactions to sucrose in addition to producing aphagia, whereas posterior LH lesions produced merely aphagia without any aversion. To more thoroughly localize the site of 'disgust' release, Cromwell and Berridge [35] made discrete excitotoxic lesions in VP (anterolateral to LH) or in nearby regions such as lateral hypothalamus and the preoptic area. They confirmed that lesions to all LH and VP sites produced aphagia, but found that only lesions that damaged VP produced the flip in affective responses to sucrose from 'liking' to 'disgust'. Even anterior LH lesions did not release 'disgust' if VP was spared. Temporary inhibitions by muscimol microinjections into VP also have been reported to increase aversive reactions to sucrose [101]. More recently, a PhD dissertation study by Chao-Yi Ho, which mapped the increase of aversive reactions to sucrose, demonstrated that it was the VP hotspot in caudal VP that appears responsible for both lesion-induced 'disgust' and muscimol-induced 'disgust': sites for either in the posterior VP hotspot produced intense 'disgust' reactions to sucrose, whereas other sites in anterior VP as well as in anterior LH did not (as long as the posterior VP hotspot remained untouched) [59]. Such findings suggest that the VP hotspot in particular is especially important for generating normal hedonic impact, as well as for amplifying intense hedonic impact, since it is the only region in the brain known so far in which lesions not only suppress hedonic reactions, but also replace them with aversive reactions to sweetness.

### 1.2.4. Anatomical basis for the VP hotspot

The larger anatomical zone in which VP is located was traditionally called the substantia innominata (SI), or unnamed substance. This was due to its lack of distinguishing features (as far as was then known), and the confusing nature of what constituted its borders, however the term substantia innominata was later criticized as too vague [57]. The VP boundaries reveal themselves when tissue is stained for enkephalin or substance P; VP produces more enkephalin and substance P than other nearby SI regions, and has distinct afferent and efferent patterns from that of the dorsally positioned globus pallidus [50,54], marking it as a relatively distinct structure within SI.

Like the NAc hotspot, the VP hotspot in its posterior region has several unique characteristics that differ from other VP subregions that may contribute to its hedonic function. For example, Kupchik and Kalivas [66] showed that the electrophysiological signature of the neurons in VP change, depending on where they recorded along a rostrocaudal axis. Neurons in anterior VP included a mix of "Type I" and "Type II" neurons, whereas posterior VP was characterized solely by Type I neurons. Type I neurons are tonically active and easily excited, while Type II neurons have low basal firing rates, and require more stimulation to elicit an action potential. In addition to this, Type II neurons morphologically resemble the accumbens medium spiny neurons, whereas Type I neurons that predominate in posterior VP are relatively aspiny and are somewhat larger than Type II. Although it is still unclear how Type I and II neurons differ functionally, it is interesting to note that the change in neuron type follows the rostrocaudal functional difference between caudal VP hotspot and rostral VP coldspot sites.

366 1.3. *Parabrachial nucleus hotspot*367 1.3.1. *Brainstem mechanisms of reward*

368 In addition to the two forebrain hotspots of NAc and VP, there is also  
369 some evidence for a brainstem hedonic hotspot within the parabrachial  
370 nucleus (PBN) of the pons [108]. Although best known as a visceral/  
371 taste sensory relay [38,79], the PBN has additional functions, including  
372 food intake [28,40,124,127], establishing a conditioned taste aversion  
373 [28,36,130], and REM sleep [90,120].

374 As noted above, Grill and Norgren [47,49] pioneered the taste re-  
375 activity paradigm in order to compare normal and decerebrate (and thal-  
376 amic or detelencephalic) rats. Mesencephalic decerebrate rats receive  
377 transections above the superior colliculus at the level of the midbrain,  
378 removing inputs from the hypothalamus, thalamus and all telencephalic  
379 forebrain structures, and display no voluntary eating. However, despite  
380 the complete lack of spontaneous eating behavior, decerebrates show  
381 normal taste reactivity patterns to palatable sucrose or aversive quinine  
382 [47,49]. Although decerebrate taste reactions are reflexive in nature,  
383 another potential implication of that finding is that even at the level of  
384 the brainstem, the beginnings of some elementary hedonic processing  
385 may be occurring [15].

386 To more directly assess brainstem hedonic function, Berridge [13]  
387 made systemic injections of chlordiazepoxide, a benzodiazepine drug  
388 that enhances hedonic reactions in normal rats as well as enhancing  
389 food intake [34,121], into decerebrate rats and found that this benzodi-  
390 azepine stimulation of the functional midbrain and hindbrain was still  
391 sufficient to enhance sucrose 'liking' reactions. Pecina and Berridge  
392 [83] then went on to show that fourth ventricular microinjections of di-  
393 azepam into the brainstem fourth ventricle of intact rats also enhanced  
394 hedonic 'liking' reactions even at low doses that were ineffective in the  
395 forebrain lateral ventricles, again indicating that indeed there was  
396 a brainstem site capable of amplifying hedonic impact for normal  
397 animals.

398 Providing further localization of brainstem benzodiazepine mecha-  
399 nisms of food motivation, Higgs and Cooper [58] demonstrated that mi-  
400 croinjections of a related benzodiazepine, midazolam, into the pontine  
401 parabrachial nucleus (PBN), but not nearby regions of brainstem,  
402 could significantly enhance food intake in non-deprived rats. Building  
403 on these findings, Soderpalm and Berridge [107] found that similar mi-  
404 croinjections of midazolam into the lateral parabrachial nucleus of nor-  
405 mal rats enhanced positive 'liking' taste reactivity patterns to sucrose  
406 taste, in addition to its hyperphagic effects, whereas microinjections  
407 into the hindbrain nucleus of the solitary tract or into midbrain ventral  
408 tegmental area did not.

409 Taken together, these studies implicate PBN benzodiazepine mecha-  
410 nisms in hedonic processing, extending the hedonic hotspot circuit to  
411 include the brainstem, as well as forebrain, sites of action.

412 Recent work on the parabrachial nucleus has supported its role in  
413 food intake. For example, work by Simansky and colleagues showed  
414 that opioid and endocannabinoid stimulation within the parabrachial  
415 nucleus also robustly increases consumption of palatable food [40,  
416 124]. Further, endogenous opioid function within PBN appears to be re-  
417 quired for food motivation, as infusions of naloxonazine completely  
418 prevented DAMGO induced hyperphagia [32].

419 More recently, Palmiter and colleagues have shown that the  
420 PBN interacts with hypothalamic mechanisms to control appetite  
421 [28,126–128]. Wu et al. [127] showed that PBN neurons are nor-  
422 mally inhibited by GABAergic projections from agouti-related pro-  
423 tein (AgRP) neurons in the hypothalamic arcuate nucleus, and that  
424 destruction of AGRP neurons abolished eating. They then went on  
425 to show that the starvation effects they observed through AgRP  
426 neuron ablation were not due to increased melanocortin signaling  
427 [126], but rather to over-excitation of PBN from glutamate projections  
428 originating in the hindbrain nucleus of the solitary tract or serotonin  
429 neurons [128]. Oppositely, Carter et al. [28] showed that direct  
430 optogenetic inhibition of lateral PBN calcitonin neurons produced

excessive eating. Such mechanisms are potential candidates for future  
studies of PBN roles in hedonic impact.

433 1.4. *A functional circuit for hedonic processing*

434 The existence of multiple hedonic hotspots allows for the possibility  
435 that the hotspots interact and work together within a coordinated he-  
436 donic circuit. A functional circuit would not necessarily imply that the  
437 hotspots are all directly connected anatomically, since intermediary  
438 stops could be equally effective in creating a functional circuit. To deter-  
439 mine whether at least a functional interaction existed, Smith and  
440 Berridge unbalanced the circuit by infusing DAMGO into one hotspot  
441 (e.g. NAc), while simultaneously infusing naloxone, an opioid antago-  
442 nist, into another hotspot (e.g. VP) [105]. The guiding hypothesis was  
443 that if the simultaneous opioid neurotransmission is required in both  
444 hotspots, essentially creating unanimous opioid votes for enhancement  
445 in both sites, to increase 'liking' reactions to a palatable sweet solution,  
446 then blocking endogenous opioid signals in one hotspot should prevent  
447 exogenous opioid stimulation by DAMGO microinjection in the other  
448 from causing any hedonic enhancement. The results supported this hy-  
449 pothesis: opioid blockade in either the VP or NAc hotspot prevented  
450 DAMGO enhancement of positive 'liking' reactions in the other hotspot.  
451 Further supporting the functional relationship between the NAc and VP  
452 hotspots, it was also found that DAMGO activation in one hotspot en-  
453 hanced Fos activity both locally and in the other hotspot, and in both di-  
454 rections, demonstrating that their functional interactions could be  
455 detected via neural markers of genomic transcription. It should be  
456 noted that although naloxone in VP prevented DAMGO-enhanced  
457 'liking' in the NAc hotspot, enhancements of eating by NAc DAMGO  
458 were still robustly generated, suggesting again independent controls  
459 for hedonic 'liking' versus motivated 'wanting' of the same food reward.

460 In a further electrophysiological demonstration of NAc–VP hotspot  
461 interactions, Smith et al. [106] recorded taste reactivity responses and  
462 extracellular neuronal firing patterns in the VP hotspot during an  
463 intraoral infusion of sucrose. They found that neurons in the VP hotspot  
464 appeared to encode the impact of sucrose in neuronal firing, correlating  
465 with behavioral 'liking' reactions. This hedonic pattern manifested itself  
466 by steadily increasing the neural firing rate in a slow-onset but  
467 sustained the burst of action potentials, becoming evident during the  
468 first 1.5 s after the sweet taste was introduced, and sustaining this ele-  
469 vation in firing for the duration of the 10-s sucrose infusion. DAMGO mi-  
470 croinjection into the NAc hotspot enhanced both behavioral hedonic  
471 taste reactivity to sucrose and the hedonic pattern of neural firing in  
472 VP elicited by the sweet taste. In behavioral contrast, amphetamine  
473 microinjections that potentiated dopamine transmission in the NAc  
474 hotspot only increased food intake and a more transient VP neural sig-  
475 nal burst that encoded cue-triggered 'wanting', and correlated with  
476 amount of food eaten, but had no effect on behavioral taste reactivity  
477 'liking' patterns or on the hedonic-encoding VP neural response to  
478 sucrose. Altogether, these results show that the VP and NAc hotspots  
479 interact to form a larger functional circuit that mediates the hedonic  
480 reaction to a palatable taste.

481 1.4.1. *Anatomically unconnected hotspots?*

482 Although the evidence presented so far clearly indicates a functional  
483 relationship between the hotspots, it may be surprising to note that the  
484 NAc, VP and PBN hotspots do not have any known direct reciprocal an-  
485 atomical connections between them. For example, although the NAc  
486 hotspot sends robust projections to the ventral pallidum, they are  
487 primarily directed toward rostromedial VP, and not to the posterior  
488 hotspot [50,117,122,133]. Instead, the caudolateral core sends projec-  
489 tions to the caudolateral VP region that contains the hotspot [50].  
490 Beyond the NAc–VP projection, a NAc–PBN projection also exists. How-  
491 ever, these NAc projections originate from the ventral half of medial  
492 shell, and not the dorsal half that primarily houses the rostral hotspot,

493 leaving it unclear if the NAc hotspot and PBN hotspot are directly  
 494 connected [122] (Fig. 2).

495 An analysis of VP connections shows that it sends topographic effer-  
 496 ents to NAc, so that anterior NAc connects with anterior VP, whereas  
 497 posterior NAc connects with posterior VP [51]. This suggests that  
 498 these two hotspots do not anatomically connect directly to each other  
 499 (despite their clear functional relationship). Unlike NAc, VP does not  
 500 project to PBN at all, although VP does reach other brainstem areas  
 501 such as locus coeruleus and the raphe nuclei [51]. Similarly, PBN effer-  
 502 ents do not appear to innervate NAc as far as is known [3], though  
 503 they still might possibly interact, such as via lateral PBN efferents to  
 504 the VP hotspot [96]. However, no study to our knowledge has systemat-  
 505 ically mapped PBN projections to caudal VP, leaving this connection  
 506 somewhat unresolved [53,76].

507 Altogether, an anatomical analysis of what is known of the current  
 508 hotspot boundaries suggests that although the hotspots must work to-  
 509 gether, it cannot be via direct connections. If this is true, then hotspot

activity is likely monitored and mediated by an as yet unidentified  
 brain region that shares reciprocal connections with the hotspots. 511

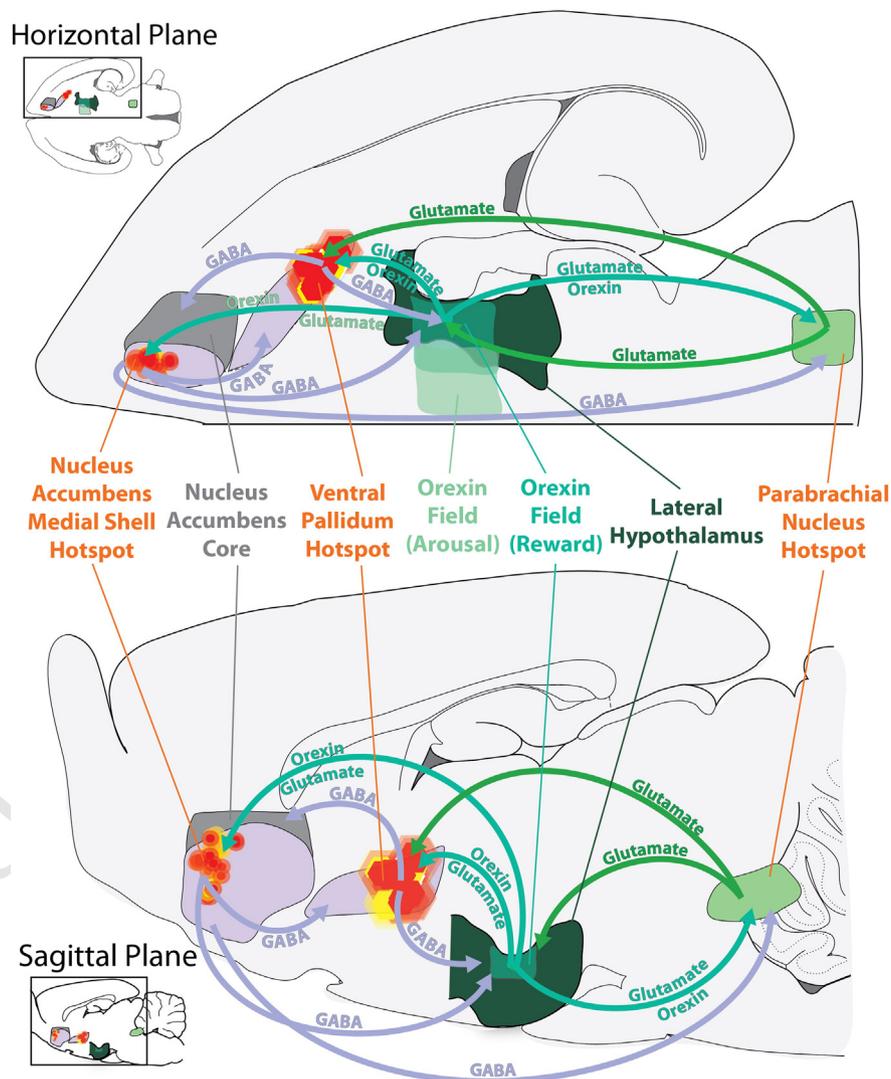
#### 1.4.2. A role for orexin in hedonic processing 512

Hunger modulates the hedonic impact of food through the phenom-  
 enon known as alliesthesia [24,25]. One candidate mechanism to help  
 mediate interactions between regulatory-hedonic circuitry is the hypo-  
 thalamic orexin/hypocretin neurons, which both project to and receive  
 direct inputs from all of the hotspots [4,8,50,51,56,89,132]. 517

Orexin neurons relevant to reward appear localized within a small  
 portion of perifornical and lateral hypothalamus [4,9,29,55,56,88]. In  
 other hypothalamic regions, such as in dorsomedial hypothalamus  
 orexin/hypocretin neurons are mostly implicated in attention, arousal  
 and sleep/wake cycles [1,11,27,42,45]. 522

Reward-related orexin neurons in lateral hypothalamus are located  
 just medial to the internal capsule and lateral to the perifornical area,  
 heavily concentrated in the dorsal and magnocellular portions of LH. 525

## Anatomical Circuitry of the Hotspots



**Fig. 2.** Anatomical connections of the hotspots. Horizontal and sagittal maps show anatomical connections of nucleus accumbens and ventral pallidum (purple), and locations of hotspots within each (red/yellow), and interactions with brainstem parabrachial nucleus (light green) and lateral hypothalamus (dark green). As indicated, the three hotspots do not share any known reciprocal connections. Potentially relevant to the homeostatic regulation of the hedonic circuit, lateral hypothalamic orexin/glutamate neurons (shown in blue/green) have reciprocal connections with all the hotspots. Based on anatomical studies cited in text. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

While a few orexin neurons can be found as far dorsal as zona incerta, most are located more ventrally (although many MCH-containing neurons are located in zona incerta), though still more dorsal than medial tuberal nucleus. The anterior–posterior extent of orexin neurons more or less coincides with the medial tuberal nucleus, which appears just after and ends just before the orexin field boundaries [8,113].

Orexin is implicated in hunger alliesthesia [5,20,21,41,44,67,81,97]. As mentioned above, a recent work by Chao-Yi Ho in our lab found that direct orexin microinjections into the VP hotspot can selectively enhance sucrose ‘liking’ reactions [60], supporting the idea that activation of hypothalamic orexin projections to VP might enhance the hedonic impact of food.

We have recently conducted pilot studies of the role of LH-to-VP projections using optogenetic techniques to activate neurons [31]. Optogenetics has the special advantage of allowing stimulation of specific point-to-point projections [2,10]. This potentially includes LH to VP projections (by putting virus in one location such as LH to infect neuron cell bodies but putting the stimulating optic fiber in a different location such as VP that receives axon terminals).

We recently infused an excitatory channelrhodopsin-2 virus into the reward-related orexin field of lateral hypothalamus, and implanted an optic fiber in the VP hotspot, which contains orexin/glutamate terminals from that field. We found that VP illumination of the orexin terminals from LH enhanced the hedonic ‘liking’ reactions to sucrose and also enhanced the motivation to consume food (measured by intake of palatable M&M candies) [31]. In contrast, direct illumination of LH neurons, by placing both optic fiber and virus in LH, increased only food intake, but did not increase sucrose ‘liking’, consistent with similar effects previously found from electrical stimulation of the LH [16,35]. Finally, direct stimulation of VP neurons, by placing both illuminating optic fiber and virus microinjection in posterior VP, specifically enhanced hedonic reactions to sucrose, without increasing food intake, supporting VP hotspot involvement in amplifying hedonic impact [60,104]. Taken together, these results indicate that neurons in the VP hotspot, and LH projections to the VP hotspot, are capable of amplifying sweetness hedonic impact.

## 2. Conclusion

Since the identification of localized hedonic hotspots in NAc, VP and the brainstem, it has become increasingly clear that these hotspots are specialized generators of hedonic impact in food reward, and that they work together to form a larger functional hedonic circuit. Future work will extend this understanding, as well as the search for additional hotspots in the brain. Some potential targets for future searches include regions of the limbic prefrontal cortex, such as the orbitofrontal cortex and insula, which are known to encode food hedonic impact in human neuroimaging studies [37,64,65,95,100,103].

In conclusion, exciting advances have been made since the initial discovery of the hotspots, and future studies can be expected to further elucidate how the brain takes a simple sensory stimulus, such as the taste of sweet food, and applies a hedonic gloss to make that sensation become positively ‘liked’.

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