

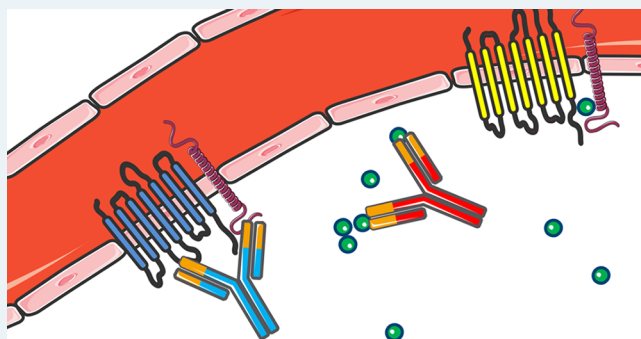
# CGRP-Based Migraine Therapeutics: How Might They Work, Why So Safe, and What Next?

Andrew F. Russo<sup>\*,†,‡,§,Ⓢ</sup>

<sup>†</sup>Departments of Molecular Physiology and Biophysics, <sup>‡</sup>Neurology, University of Iowa, Iowa City, Iowa 52242, United States

<sup>§</sup>Center for the Prevention and Treatment of Visual Loss, Iowa VA Health Care System, Iowa City, Iowa 52246, United States

**ABSTRACT:** Migraine is a debilitating neurological condition that involves the neuropeptide calcitonin gene-related peptide (CGRP). An exciting development is the recent FDA approval of the first in an emerging class of CGRP-targeted drugs designed to prevent migraine. Yet despite this efficacy, there are some fundamental unanswered questions, such as *where* and *how* CGRP works in migraine. Preclinical data suggest that CGRP acts via both peripheral and central mechanisms. The relevance of peripheral sites is highlighted by the clinical efficacy of CGRP-blocking antibodies, even though they do not appreciably cross the blood-brain barrier. The most likely sites of action are within the dura and trigeminal ganglia. Furthermore, it would be foolish to ignore perivascular actions in the dura since CGRP is the most potent vasodilatory peptide. Ultimately, the consequence of blocking CGRP or its receptor is reduced peripheral neural sensitization. Underlying their efficacy is the question of *why* the antibodies have such an excellent safety profile so far. This may be due to the presence of a second CGRP receptor and vesicular release of a large bolus of peptides. Finally, despite the promise of these drugs, there are unmet gaps because they do not work for all patients; so *what next*? We can expect advances on several fronts, including CGRP receptor structures that may help the development of centrally acting antagonists, combinatorial treatments that integrate other therapies, and development of drugs that target other neuropeptides. This is truly an exciting time for CGRP and the migraine field with many more discoveries on the horizon.



**KEYWORDS:** migraine, CGRP, therapeutic antibodies, neuropeptides, photophobia, pain

## INTRODUCTION

Migraine is more than just a bad headache. It is neurological disorder involving sensory abnormalities that occur preceding, during, and following the headache.<sup>1–3</sup> The prevalence and socio-economic impact of migraine cannot be overstated. It is estimated that 15% of all people suffer from migraine.<sup>4,5</sup> Migraine is the second leading cause of years lived with a disability, and the sixth most common disease globally (>1 billion sufferers).<sup>6</sup> The prevalence is 3–4-fold higher in women, such that over 40% of women experience migraine in their lifetime.<sup>7</sup> The annual cost of migraine in the US alone is ~\$36 billion.<sup>8</sup> Furthermore, headache is in the top five reasons for emergency room visits<sup>4</sup> and is associated with the overprescription of opioids.<sup>9</sup>

Clinical studies over the past 3 decades have demonstrated that migraine involves an increased sensitivity to the neuropeptide calcitonin gene-related peptide (CGRP). CGRP is a multifunctional peptide that has potent vasodilatory activity and has long been implicated in pain pathways.<sup>10</sup> Notably, CGRP is both necessary and sufficient to cause migraine for many people (Table 1). In particular, CGRP infusion is able to trigger delayed migraine-like headaches in most migraineurs. Interestingly, the majority of agents in these

types of provocation studies are vasodilators. However, this is not a property of all vasodilators since infusion of VIP and adrenomedullin do not trigger a migraine.<sup>11,12</sup> This suggests that the vasodilation activity of CGRP is not sufficient to trigger a migraine. Nonetheless, we cannot exclude the possibility that CGRP invokes both vasodilatory and non-vasodilatory actions in the perivascular,<sup>13</sup> or that there are important differences in vascular targets and pharmacokinetics.<sup>14,15</sup> On-going preclinical and clinical studies should resolve these possibilities.

Perhaps the most persuasive evidence of the importance of CGRP is efficacy of CGRP-based therapeutic antibodies and small molecule receptor antagonists.<sup>16</sup> Three monoclonal antibodies have now been approved by the Federal Drug Administration (FDA). Erenumab (Aimovig, Amgen/Novartis), blocks the CGRP receptor and fremanezumab (AJOVY, Teva Pharmaceuticals) and galcanezumab (Emgality, Eli Lilly) block the CGRP ligand. In addition, two receptor antagonists, ubrogepant (Allergan) and rimegepant (Biohaven Pharmaceuticals), and another ligand antibody (eptinezumab, Alder

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Table 1. Clinical Evidence of CGRP Involvement in Migraine

## 1. Elevation of CGRP in migraineurs

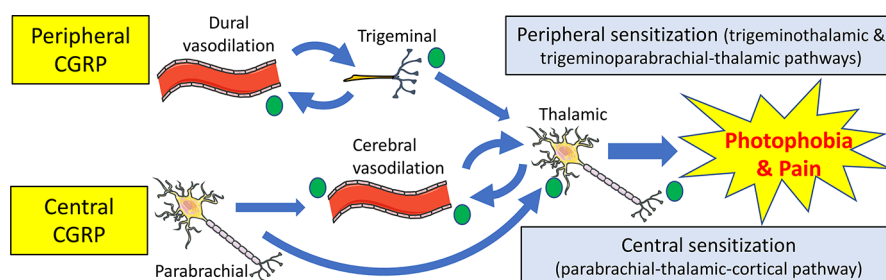
- CGRP levels elevated in plasma,<sup>80,81</sup> cerebrospinal fluid,<sup>82</sup> and saliva<sup>83</sup> during spontaneous migraine attacks.
  - Also elevated during nitroglycerin-induced migraine.<sup>84</sup>
  - Interictal levels elevated in episodic<sup>85</sup> and chronic migraineurs.<sup>86</sup>
  - Reduced by triptans.<sup>81,84</sup>
- However, elevation not seen in other episodic<sup>87</sup> and chronic<sup>88</sup> migraine studies.

## 2. Infusion of CGRP causes migraine

- In 66% of migraineurs, the infusion of CGRP is sufficient to induce a migraine-like headache.<sup>89,25,90,91</sup>
  - In contrast, healthy controls have only a mild headache,<sup>92</sup> suggesting that migraineurs are more sensitive to CGRP.
- However, CGRP infusion not effective in FHM1 patients<sup>93,94</sup> and apparently does not induce aura<sup>90</sup> or prodrome symptoms.<sup>91</sup>

## 3. Efficacy of CGRP-based drugs

- Small molecule CGRP receptor antagonists effective in clinical trials for abortive treatment of migraine.<sup>95–97</sup>
  - Lead antagonist dropped due to liver toxicity after repeated use,<sup>98</sup> although new compounds, rimegepant and ubrogepant, look promising as abortive and preventative drugs,<sup>73</sup> and are expected to be submitted for FDA approval soon.
- Antibodies that block CGRP or CGRP receptor are effective in clinical trials for prevention of both episodic and chronic migraine.<sup>18,19,73,99–101</sup>
  - FDA approvals of the receptor antibody erenumab (Aimovig) and two ligand antibodies, fremanezumab (AJOVY) and galcanezumab (Emgality), for migraine prevention, with another ligand antibody, eptinezumab, expected to be submitted soon.



**Figure 1.** Model of peripheral and central perivascular CGRP actions. In the periphery, CGRP (green ovals) released from dural trigeminal afferents acts on perivascular cells to cause vasodilation and neurogenic inflammation, with positive feedback loops (blue arrows) leading to peripheral sensitization.<sup>33</sup> Nociceptive signals are relayed to the CNS to the thalamus directly and via the parabrachial nucleus. In the CNS, release of CGRP (green ovals) from parabrachial neurons into the posterior thalamic region modulates neural signaling. CGRP in the brain also causes vasodilation,<sup>49</sup> which could lead to neural activation by vascular-neural coupling<sup>51</sup> and further dilation of vessels in a positive feedback loop (blue arrows) that triggers further CGRP release. CGRP-mediated neuromodulation of glutamatergic synapses results in central sensitization<sup>33</sup> and signaling to the cortex that leads to migraine symptoms of photophobia and pain.

Biopharmaceuticals) are expected to be submitted for FDA approval in 2019. It is especially encouraging that the antibodies are effective for at least 15 months,<sup>17</sup> and have minimal adverse effects.<sup>18–21</sup> This is a stimulating time in the field because CGRP-based drugs are the first new class of migraine therapeutics in nearly 30 years.

## ■ HOW IS CGRP ACTING IN MIGRAINE?

The role of CGRP and the vasculature in migraine is central to many of the concepts in this article. Historically migraine has been viewed as a vascular disorder,<sup>22</sup> yet over the past two decades the vascular theory has been challenged by more neuro-centric theories. This shift was triggered by reports that vascular changes are neither necessary nor sufficient to trigger migraine, and by evidence that brain functions are altered during migraine.<sup>3,23</sup> Nonetheless, the debate on vascular contributions continues (see citations in ref 24).

In this Perspective, I suggest that CGRP actions at the vasculature should not be ignored. We have recently reviewed the vascular connections to migraine.<sup>24</sup> For example, local changes in vascular tone are difficult to rule out and recent studies support a role for meningeal vasodilation.<sup>25,26</sup> Furthermore, a meta-analysis of >1 million people concluded that migraine is associated with elevated risk of cardiovascular and cerebrovascular events,<sup>27</sup> and many genes that are associated with migraine are expressed in the vasculature.<sup>28</sup>

Finally, the high therapeutic efficacy of monoclonal antibodies that do not cross the blood-brain barrier<sup>16,29</sup> argues for a peripheral site of CGRP action. On the basis of these observations, a critical reevaluation of vascular contributions to migraine seems justified. Indeed, a neurovascular model of migraine involving peripheral sensitization in the trigemino-vasculature was articulated over 25 years ago, although it was limited to peripheral nerves.<sup>22</sup> The model I am proposing is similar, with the exceptions that the process can be triggered in both the meninges and CNS, and that it can go in both directions, that is, *neural to vascular* and *vascular to neural*.

I propose a model of perivascular CGRP actions that involves a cascade from the trigeminovascular to the thalamus and eventually the cortex (Figure 1). A key point is that at both sites of action the blood vessels can modulate neural activity. In this model, CGRP can act in both the periphery and CNS to cause migraine symptoms. This hypothesis is based on our unexpected finding from photophobia studies described below. Namely, when mice were sensitized to CGRP by overexpression of a receptor subunit (hRAMP1) in neural tissue, they showed enhanced light aversive behavior following central administration of CGRP, but not following peripheral administration.<sup>30</sup> Although this finding does not rule out involvement of the nervous system, it does indicate that the limiting site of peripheral CGRP action

is not neuronal. If neurons are not the limiting site in the periphery, then which cells are?

In the periphery, it seems likely that a combination of CGRP actions at multiple sites could alter the microenvironment of the trigeminovascular system (Figure 1). Peripheral targets most likely are (1) perivascular cells in the meninges, including mast cells, glia, vessels, primary afferents and (2) trigeminal ganglia cell bodies, including neurons and glia.<sup>31</sup> Notably, CGRP is the most potent peptide vasodilator, especially of cranial vessels,<sup>10</sup> and CGRP leads to neurogenic inflammation by activating mast cells, which release agents that can sensitize neurons and cause further vasodilation in the dura layer of the meninges.<sup>32</sup> This modulation of neural activity could then trigger positive feedback loops that lead to peripheral sensitization of nociceptors.<sup>31,33</sup> Dural vessels are fully peripheral, unlike vessels of the pial layer of the meninges. The pial afferents are exposed to cerebrospinal fluid and hence, unlike dural afferents, are within the blood brain barrier.<sup>34</sup> CGRP-induced dilation could trigger further dilation by autoregulation of local cerebral blood flow in the pia and parenchyma. In the periphery, CGRP is most likely acting by peripheral sensitization that set up vascular and neural actions in the CNS.

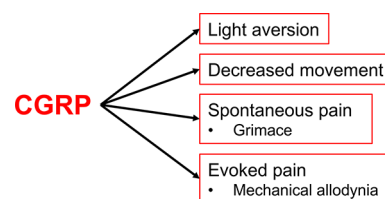
In the CNS, we expect that multiple neural pathways influence photophobia and pain, with the thalamus being a focal point (Figure 1). CGRP and its receptor subunits are widely distributed throughout the CNS, including the retina.<sup>35,36</sup> The trigeminal nerve carries nearly all the pain signals from the anterior part of the head<sup>31</sup> and is generally thought to be involved in most photophobia pathways.<sup>33</sup> The trigeminothalamic tract consists of second order trigeminovascular neurons that directly connect from the trigeminal nucleus to posterior thalamic nuclei (PoT). Note that we use the designation PoT to include all posterior thalamic nuclei, including the ventral posteromedial nucleus (VPM). A second path is via the parabrachial nucleus (PBN), which is known to relay sensory signals, including pain-producing signals, to the forebrain. The role of CGRP-expressing neurons in the PBN in a variety of aversive responses,<sup>37</sup> has recently been reviewed.<sup>38</sup> In addition, the PBN receives some direct monosynaptic input from the trigeminal nucleus, which contributes to affective pain.<sup>39</sup> A key study by Burstein and colleagues identified neurons in the rat PoT (primarily in the dorsal border) that might be involved; they are coordinately activated by dural stimulation of trigeminal afferents and melanopsin-containing retinal ganglion cells.<sup>40</sup> A role for CGRP in trigeminal and thalamic pathways is also supported by the presence of CGRP and its receptors in the nociceptive pathway from the meninges to the PoT,<sup>41,42</sup> and PoT activation during migraine.<sup>43</sup> The presence of CGRP in PoT neurons of the subparafascicular and intralaminar thalamic nuclei is especially intriguing because they receive somatosensory and nociceptive stimuli from ascending pathways.<sup>44</sup> In addition, the convergence of auditory and nociception inputs to the CGRP-positive subparafascicular thalamic nucleus are important for conditioned auditory and visual fear responses.<sup>45,46</sup>

An under-appreciated function of central CGRP is its ability to trigger vasodilation of cerebral blood vessels. While long documented *in vitro*,<sup>47,48</sup> central CGRP-induced vasodilation was only recently shown *in vivo*,<sup>49</sup> ironically, in the thalamus. Furthermore, CGRP could potentially act at vessels distant from its release site by diffusion through interstitial space (volume transmission).<sup>50</sup> This would create a second positive

feedback loop (Figure 1). In the CNS, this activity would be restricted to perivascular CGRP since it would not pass into the lumen of cerebral vessels. As an aside, CGRP detected in the blood is most likely spillover from peripheral nerves (e.g., from trigeminal afferents),<sup>16</sup> although there are also reports of nonneuronal sources of CGRP.<sup>10</sup> Dilation of cerebral vessels could then increase neural activity in the CNS by vascular-neural signals. This mechanism is supported by studies with *in vitro* cortical slices, and might occur either directly or via glial intermediates.<sup>51</sup> Coupled with vascular actions, CGRP released from neurons is known to act as a neuromodulator. This activity has been shown to increase glutamatergic transmission, and thus could cause central sensitization.<sup>52,53</sup>

## ■ WHAT ARE THE MIGRAINE-LIKE ACTIONS OF CGRP?

Much of what we know about CGRP actions comes from preclinical studies with mice (Figure 2). Admittedly, we will



**Figure 2.** Summary of CGRP-induced migraine-like symptoms. Injection of CGRP into mice can cause the indicated symptoms of light aversion, decreased movement, and spontaneous and evoked pain.

never be sure if a mouse has a migraine; however, we can examine behaviors that are surrogates for migraine symptoms, such as photophobia and pain. Light aversion can be used as a photophobia assay.<sup>30,54,55</sup> Grimace and touch sensitivity can be used as a measure of CGRP-induced spontaneous and evoked pain.<sup>56,57</sup>

**Photophobia and Decreased Movement.** Light aversive behavior has been established as a surrogate for photophobia.<sup>30,54,55</sup> Photophobia reflects an allodynic response, that is, nonpainful light becomes noxious, and is one of the diagnostic criteria and most common symptoms of migraine.<sup>58</sup> Bright light can trigger a migraine, and even dim light causes discomfort and pain during a migraine. Administration of CGRP to mice drove them into the dark zone.<sup>30,54,55</sup> While we cannot be certain that mice experience photophobia, we have ruled out that increased anxiety alone drives the behavior. Thus, we presume that aversion to light must reflect an unpleasant sensation that can overcome their strong, innate exploratory drive. Coupled with this behavior, we noticed that the mice also moved less after being given CGRP. Interestingly, the mice only moved less in the dark zone, which may be similar to how migraineurs prefer to rest when they go into the dark.

One important feature of the assay is that either dim or bright light triggers a response. In dim light (55–70 lx), CGRP-sensitized mice, that is, mice in which hRAMP1, the rate-limiting subunit of the CGRP receptor, is overexpressed in the nervous system (nestin/hRAMP1), but not wildtype or littermate control mice, spent less time in the light following intracerebroventricular (icv) injection of CGRP.<sup>54</sup> As mentioned above, this sensitized phenotype was not seen in nestin/hRAMP1 mice after peripheral (intraperitoneal, ip) CGRP



delivery compared to wildtype mice.<sup>30</sup> The finding that nestin/hRAMP1 mice are sensitized to central CGRP but not to peripheral CGRP suggests that neurons are not the rate-limiting step for peripheral CGRP actions. Given that people who do not get migraine are bothered by bright light, it seemed that if the light intensity is great enough, then wildtype mice might also show CGRP-induced light aversion. This proved to be the case for both icv<sup>55</sup> and ip<sup>30</sup> CGRP delivery to wildtype mice when the light intensity was increased to the equivalent of that on a sunny day (25–27K lux). Antimigraine drugs of the triptan family blocked the effect of bright and dim light in the contexts of both icv and ip CGRP,<sup>30,55</sup> consistent with their ability to inhibit CGRP-induced migraine in humans.<sup>25</sup>

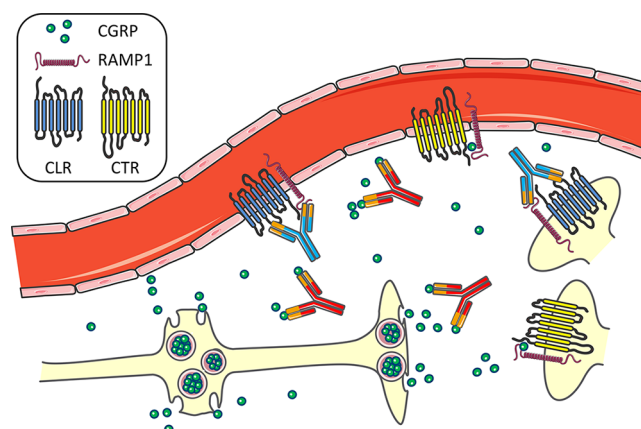
**Spontaneous and Evoked Pain.** Pain is admittedly a difficult parameter to objectively measure in mice as well as in people. An indicator of spontaneous pain is the mouse grimace scale developed by Mogil.<sup>59</sup> Injection of CGRP induces a grimace response that lasts for over an hour.<sup>56</sup> The grimace is partially attenuated by sumatriptan, consistent with the pain being at least partially migraine-like. Importantly, given the role of CGRP in light aversive behavior, the grimace is independent of light, being observed both in the dark and light.<sup>56</sup> A strength of this assay is that it is translatable to humans, who also grimace when in pain.<sup>59</sup>

An indicator of evoked pain is tactile mechanical allodynia of the periorbital facial region and hindpaw using von Frey filaments. This tactile sensitivity is considered to be a painful response to ordinarily nonpainful touch. Sensitivity to light touch, predominantly in the cephalic region, is reported by over half of migraineurs.<sup>60</sup> We and others have reported that CGRP enhances sensitivity to von Frey filaments used as mechanical stimuli to the hindpaw.<sup>57</sup> Therefore, grimace and tactile allodynia assays provide independent assessments of pain in mice following CGRP treatment.

### ■ WHY ARE THE CGRP AND CGRP RECEPTOR BLOCKING ANTIBODIES SO SAFE THUS FAR?

To date, the antibodies have a remarkably clean safety profile. This is despite reasonable concerns about cardiovascular complications of blocking CGRP activity,<sup>61</sup> especially given its cardioprotective functions.<sup>62</sup> Toward this end, it is encouraging that the CGRP receptor antibody did not affect patients with stable angina on a treadmill test.<sup>21</sup> While we must remain cautious, why might the antibodies have been so safe thus far? The answer could lie in part because the antibodies *knockdown*, but do not *knockout* CGRP signaling (Figure 3).

In the case of the CGRP receptor antibody, it blocks only the canonical CGRP receptor (Figure 3). This receptor is an unusual G protein coupled calcitonin-like receptor (CLR) that requires the RAMP1 accessory protein for both trafficking to the cell surface and binding of CGRP.<sup>63,64</sup> Our data indicate that RAMP1 is the rate-limiting subunit.<sup>65</sup> Interaction of CLR with related RAMPs generates receptors for adrenomedullin, which is involved in nociception<sup>66</sup> but not migraine.<sup>11</sup> The hRAMP1 subunit is responsible for the species selectivity of the CGRP receptor antagonists used in clinical studies.<sup>67</sup> Importantly, RAMP1 can bind other G protein coupled receptors. Most notably, RAMP1 converts the calcitonin receptor (CTR) into an amylin-1 receptor, which can be a second CGRP receptor.<sup>68,69</sup> This receptor is also sensitized in the transgenic hRAMP1 mice.<sup>70,71</sup> The amylin-1 (CTR/RAMP1) receptor is reportedly not blocked, based on



**Figure 3.** Model of CGRP and CGRP receptor antibody actions in the trigeminovascular system. CGRP is released in a large bolus from dense core vesicles at varicosities and free nerve endings of trigeminal afferents in the meninges. The ligand antibodies are proposed to dampen CGRP signaling by binding some, but not all, of the released CGRP. The receptor antibody blocks the CLR/RAMP1 CGRP receptor, but apparently not the CTR/RAMP1 amylin-1 receptor, which would allow CGRP signaling to still occur at CTR/RAMP1 on vessels and trigeminal nerves.

pharmacological data, by the CGRP receptor antibody that was raised against CLR/RAMP1.<sup>72</sup> Since both the amylin-1 and CGRP receptors are expressed in vessels, as well as the trigeminal ganglia and trigeminal nucleus,<sup>68</sup> this suggests that there could be at least partial compensation when the CGRP (CLR/RAMP1) receptor is blocked. Thus, in the presence of the CGRP receptor antibody, CGRP could still potentially signal at the amylin-1 receptor.

In the case of the ligand antibodies, the blocking antibodies likely only dampen, not completely block, the relatively slow and prolonged actions of neuropeptides over a relatively large area (Figure 3). The vesicular storage of neuropeptides means that when a neuron is activated, it rapidly releases peptides in a huge bolus that could potentially overwhelm the antibodies. In essence there would be a “binding race” driven by the kinetics of CGRP binding its receptor and being bound by an antibody. To appreciate the odds on this race, a relevant consideration is the number of peptides released from a single vesicle. Primarily on the basis of peptide and capacitance measurements of hypothalamic neurons, it can be estimated there are ~10 000 peptides per dense core vesicle, with ~10<sup>3</sup> vesicles released per sec.<sup>50</sup> Thus, millions of neuropeptides can be released in a short burst from just a single neuron. As a result, it seems likely that the blocking antibodies will be able to blunt, but are unlikely to completely block, neuropeptide signaling. Rather, I suggest that the amplitude and duration of volumetric transmission (dispersion in extracellular fluid) of the peptides would be decreased, but not abolished. For a more detailed discussion of the relevance of peptide release and transmission in migraine see ref 50. Hence, the inability to completely block CGRP upon release from neurons and the existence of a second CGRP receptor may contribute to the excellent safety profile that has been reported to date for the CGRP- and CGRP receptor-blocking antibodies in clinical trials.

### ■ WHAT NEXT?

The development of CGRP-based drugs is clearly a great beginning;<sup>73</sup> however, there are unmet gaps for treatments. For

example, a clinically relevant end point of 50% reduction of headache days is achieved in only about half of the subjects in trials.<sup>74</sup> To improve existing treatments and develop new drugs, we need to know more about where and how CGRP acts. Along this line, centrally acting CGRP antagonists should be considered. Importantly, given the actions of CGRP in central sensitization of pain,<sup>75</sup> it seems likely that CGRP targeted drugs will also be effective for other chronic pain syndromes. However, this potential benefit of a centrally acting antagonist will need to be balanced against the likely risk of greater side effects on CNS function. Likewise, a better understanding of the structures of both CGRP receptors will help direct drug design for both peripheral and central target engagement. In this regard, the recently solved cryo-EM structure of the active canonical CGRP receptor (CGRP:CLR:RAMP1:Gs heterotrimer)<sup>76</sup> is likely to provide new perspectives for drug development. In this vein, antibodies and drugs that target the second CGRP receptor should be considered. In addition, combinatorial treatments, in which CGRP targeted therapies may be integrated into other treatment regimens, such as NSAIDs, may prove effective.

Finally, where to after CGRP? New drug targets should include other neuropeptides. Indeed, at least two pharmaceutical companies are moving forward with antibodies that block either PACAP-38 or the PAC1 PACAP receptor. Given the shared activities of CGRP and PACAP,<sup>77,78</sup> this is likely to be a fruitful endeavor. But these two peptides are just the very tip of the iceberg. There are hundreds of neuropeptides that can act within the brain as neuromodulators and within the periphery as signaling molecules, which makes them well poised to alter sensory perception in migraine.<sup>50</sup> In the periphery, an example of neuropeptide actions is at the cerebrovasculature, which is heavily innervated by sensory, parasympathetic, and sympathetic nerves.<sup>79</sup> These nerves release several neuropeptides, including CGRP, PACAP, and NPY, that can either increase or decrease vascular tone. Whether in the perivasculature or deep in the CNS, members of the diverse families of neuropeptides could add to the pathogenesis and complexity of migraine.<sup>50</sup> Clearly CGRP will not be the only neuropeptide involved in migraine and given the emerging evidence for other peptides, it seems reasonable that altered neuropeptide actions may be a good target for reducing the heightened sensory state of migraine.

In summary, the advent of new CGRP-based migraine drugs established the significance of CGRP in migraine, yet also highlighted how little we know about the underlying mechanisms. It seems likely that we are heading for a very exciting time in the field.

## AUTHOR INFORMATION

### Corresponding Author

\*E-mail: [andrew-russo@uiowa.edu](mailto:andrew-russo@uiowa.edu). Address: Departments of Molecular Physiology and Biophysics, 51 Newton Road, University of Iowa, Iowa City, IA 52242.

### ORCID

Andrew F. Russo: 0000-0002-8156-5649

### Notes

The author declares no competing financial interest.

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