A peptide toxin in ant venom mimics vertebrate EGF-like hormones to cause long-lasting hypersensitivity in mammals

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Venoms are excellent model systems for studying evolutionary processes associated with predator–prey interactions. Here, we present the discovery of a peptide toxin, MIITX\textsubscript{2}-Mg1a, which is a major component of the venom of the Australian giant red bull ant \textit{Myrmecia gulosa} and has evolved to mimic, both structurally and functionally, vertebrate epidermal growth factor (EGF) peptide hormones. We show that Mg1a is a potent agonist of the mammalian EGF receptor ErbB1, and that intransplantation into mice causes long-lasting hypersensitivity of the injected paw. These data reveal a previously undescribed venom mode of action, highlight a role for ErbB receptors in mammalian pain signaling, and provide an example of molecular mimicry driven by defensive selection pressure.

**Results**

MIITX\textsubscript{2}-Mg1a is a Major Component of \textit{M. gulosa} Venom. In a previous study (4), we showed that the venom of \textit{M. gulosa} was composed primarily of amphipathic peptides derived from the aculeatoxin gene family, but with one exception: a unique cysteine-rich peptide, MIITX\textsubscript{2}-Mg1a (hereafter Mg1a), which is unrelated to the aculeatoxins. The cysteine spacing and conservation of Gly39 and Arg41 residues (8) in Mg1a suggested that it might adopt an EGF-like fold (Fig. 1B). The EGF-like fold is characteristic of an ancient superfamily of peptide hormones that have diverged across Metazoa (9); in insects, 4 different EGF-like peptide hormones (Spitz, Vein, Gurken, and Keren) have been reported, while mammals have 11 EGF-like peptide hormones: EGF, amphiregulin, epgen, epiregulin, betacellulin, heparin-binding EGF-like growth factor (HBEGF), transforming growth factor-\(\alpha\) (TGF\(\alpha\)), and 4 neuregulins (10). While all EGF-like peptides retain the same overall fold, divergence of amino acid sequence and cysteine spacing has resulted in distinct pharmacological profiles. For example, the insect EGF-like peptide hormone Spitz is a ligand of the insect receptor (ErbB), while mammalian EGF-like peptide hormones are ligands of one or more mammalian receptors (ErbB1 to 4, also known as HER1 to 4, including heterodimers thereof) (10). A comparison of the amino acid sequence of Mg1a with that of several of the human EGF (hEGF)–like peptide hormones (Fig. 1B) illustrates the shared sequence features.

**Significance**

The targeting of mammalian ErbB receptor signaling by a venom toxin to cause hypersensitivity is a mode of action that has not previously been described. Natural selection of a defensive toxin to target ErbB signaling provides compelling independent evidence for a fundamental role of this receptor and its ligands in mammalian pain. The evolution of a toxin in ant venom to mimic a vertebrate nociceptive hormone serves as an example of both convergent evolution and molecular mimicry, illustrating how natural selection can shape the gene product of one organism to resemble that of another.

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The authors declare no competing interest.

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In order to assess Mg1α in the context of the whole venom, we collected “stung” venom from several individuals of *M. gulosa* (by inciting the ants to sting a thin layer of paraffin from which venom droplets were collected) and analyzed the venom by reversed-phase high-performance liquid chromatography (RP-HPLC) (Fig. 1C). In the HPLC chromatogram of venom collected from eight ants, one of the major peaks corresponded to Mg1α (Fig. 1C). Analysis of the expression of Mg1α in the venom-gland transcriptome of *M. gulosa* was also consistent with Mg1α being a major venom component: The transcript encoding Mg1α had an estimated expression level of 3,854 transcripts per million (TPMs), making it one of the most highly expressed transcripts (Fig. 1D). The Mg1α-encoding transcript is found in the highly expressed portion of the venom apparatus transcriptome with an estimated expression level like that of other venom peptides (highlighted in red). (https://doi.org/10.1073/pnas.2112630119 A peptide toxin in ant venom mimics vertebrate EGF-like hormones to cause long-lasting hypersensitivity in mammals)

**Mg1α Is an Analog of Mammalian EGF Peptide Hormones.** We used the mature peptide sequence of Mg1α as a query to search (using blastp) the National Center for Biotechnology Information nonredundant (NCBI nr) protein sequence database for related sequences. All of the top 100 alignments were vertebrate EGF-like peptide hormones, predominantly sequences annotated as HBEGF and betacellulin (79/100). When sorted by sequence identity, 5 of the top 10 hits (with sequence identity of 59% and identical intercysteine loop lengths) were to EGF-like peptide hormone sequences of marsupials (Mammalia: Marsupialia), for example the Tammar wallaby (*Notamacropus eugenii*), koala (*Phascolarctos cinereus*), and common wombat (*Vombatus ursinus*) (Fig. 2C). We repeated this analysis using the mature peptide sequence of Mg1α as a query to search (using tblastn) the NCBI Transcriptome Shotgun Assembly (TSA) Database. Again, the top 100 hits were vertebrate EGF-like peptide hormones, and of these the top 2 (with sequence identity of 59% and identical intercysteine loop lengths) were to EGF-like peptide hormone sequences of marsupials: the fat-tailed dunnart (*Sminthopsis crassicaudata*) and long-nosed bandicoot (*Perameles nasuta*). For comparison, the Spitz peptide hormones of *M. gulosa*, the Indian jumping ant (*Harpegnathos saltator*), western honey bee (*Apis mellifera*), and *Drosophila melanogaster* have sequence identities of 26 to 33% with Mg1α and a distinct intercysteine loop length (Fig. 2D). While these analyses were subject to limitations (e.g., the dynamic nature of the search databases and the relatively short length of Mg1α), it was clear that, despite its insect origin, the amino acid sequence and intercysteine loop length of the Mg1α mature peptide resembled vertebrate rather than invertebrate EGF-like peptide hormones, with a consistent signal emerging for similarity to those of marsupials.

Using principal-component analysis (PCA), we generated a quantitative map of protein sequence space for Mg1α and the vertebrate and invertebrate EGF-like peptide hormones (EGF, HBEGF, betacellulin, epiregulin, Spitz, and Vein) based on the biophysical properties of each residue in the EGF-like domain of each mature peptide sequence (Fig. 2B) (12). The map produced well-defined groupings for each of the EGF-like peptide hormone families. Consistent with the results of our blast analyses, Mg1α grouped near the vertebrate families (betacellulin, HBEGF, and epiregulin), in contrast to endogenous *M. gulosa* Spitz, which grouped with other Spitz sequences.
We then tested whether the observed relationships in primary structure were mirrored by the tertiary structure. For these experiments, we produced uniformly 15N, 13C–labeled Mg1a by recombinant expression in bacteria (SI Appendix, Fig. S2). The recombinant peptide was produced with an additional Gly residue at the N terminus, and we refer to it as rMg1a.

One-dimensional (1D) 1H and 2D [1H-15N]-HSQC (heteronuclear single quantum coherence) NMR spectra indicated that rMg1a adopted a single major conformation and was highly structured in solution (SI Appendix, Fig. S3). Structures were generated in CYANA using 671 nuclear Overhauser effect (NOE)–derived distance constraints and 75 dihedral-angle constraints. Structural constraints and statistics are summarized in SI Appendix, Table S1. Determination of the solution structure of rMg1a confirmed that it had an EGF-like fold (Fig. 2D) with disulfide connectivity of Cys I–III, II–IV, and V–VI and two small antiparallel β-sheets: The longer sheet (Tyr39–Tyr40 and His46–Ile47) is near the C terminus. Alignment of the closest-to-average structure of rMg1a with those of betacellulin (Protein Data Bank [PDB] ID code 1IP0) (Fig. 2D), HBEGF (PDB ID code 1XDT), epiregulin (PDB ID code 1K36), and Spitz (PDB ID code 3CA7) yielded backbone heavy-atom rmsd values of 1.37, 1.84, 1.96, and 2.87 Å, respectively. Thus, at the level of both primary (sequence identity, intercysteine loop lengths, biophysical properties) and tertiary structure, the venom peptide Mg1a more closely resembles vertebrate EGF-like peptide hormones than insect Spitz.

Together, these results led us to hypothesize that the natural target of Mg1a was not the endogenous insect ErbB receptor but a vertebrate ErbB receptor, and that its presence in the venom of M. gulosa reflects its evolution as a defensive weapon against vertebrate predators. The fact that several of the top alignments in the blast search were with EGF-like peptide hormones from Australian mammals implicated predation pressure by the insectivores among these as a potential driver of this evolutionary process. Several Australian mammals are known to feed opportunistically on solitary Myrmecia workers, but only one, the short-beaked echidna (Tachyglossus aculeatus), a monotreme, will attack colonies to feed on the brood (eggs, larvae, and pupae) (13). Because the latter likely represents a much stronger selection pressure on Myrmecia than occasional...
predation on solitary workers, we predicted that Mg1a may have evolved under selection pressure from—and therefore resemble an endogenous EGF-like peptide of—the echidna or other now-extinct mammals with similar foraging behavior. To further examine this potential evolutionary scenario, we searched the recently published genome of the echidna (14) using Mg1a as the query. Consistent with our hypothesis, we identified a peptide sequence in the echidna genome which, when aligned with Mg1a, was 60% identical (Fig. 2A). Thus, postdivergence of insect and vertebrate EGF-like peptide hormones, the ant venom peptide Mg1a appears to have convergently evolved to more closely resemble the EGF-like hormone(s) of its vertebrate predator(s).

**Mg1a Is a Potent Ligand of Mammalian ErbB1.** If Mg1a is the product of selection pressure to target a mammalian ErbB receptor, potent activity at this receptor would be expected. We determined the potency of rMg1a at human ErbB1 using a bio-luminescence resonance energy transfer (BRET) assay that quantifies the dimerization (required for activation) of the receptor, potent activity at this receptor would be expected. We product of selection pressure to target a mammalian ErbB receptor, potent activity at this receptor would be expected. We identified a peptide sequence in the echidna genome which, when aligned with Mg1a, was 60% identical (Fig. 2A). Thus, postdivergence of insect and vertebrate EGF-like peptide hormones, the ant venom peptide Mg1a appears to have convergently evolved to more closely resemble the EGF-like hormone(s) of its vertebrate predator(s).

However, it caused a significant decrease in paw-withdrawal latency (Fig. 4C) in the same assay. Thus, rMg1a was active at hErbB1 (Fig. 3A) with a half-maximal effective concentration (EC_{50}) of 6.3 nM (95% CI, 2.0 to 16.6 nM, n = 3) (Fig. 3B). For comparison, hEGF had an EC_{50} of 10.5 nM (95% CI, 6.4 to 16.6 nM, n = 3) in the same assay.

Western blot analysis was used to investigate the ability of rMg1a to induce phosphorylation of extracellular signal-regulated kinases (ERKs) 1 and 2, downstream signaling molecules of ErbB receptors (15). Mg1a caused a concentration-dependent increase in phosphorylation of Erk1/2 (P < 0.01 for 2.5 nM and P < 0.0001 for 10 nM, Sidák’s multiple comparisons), equivalent to that caused by hEGF (Fig. 3 C and D).

These data indicate that, consistent with its vertebrate-like primary and tertiary structure, rMg1a is a potent ligand of mammalian ErbB1 with activity comparable to that of hEGF, an endogenous ligand of the receptor.

**Intraplantar Injection of Mg1a in Mice Causes Long-Lasting Hypersensitivity.** Myrmecia use their sting for both predation (to incapacitate other arthropods, which they feed to their larvae) and defense (to induce pain in, and thus deter, potential predators). Our results suggested that Mg1a has evolved as a defensive weapon against vertebrate predators. Accordingly, we observed no effect on crickets (Acheta domesticus), blowflies (Lucilia cuprina), or fruit flies (D. melanogaster) after intrabdominal injection of rMg1a up to a dose of 41 nmol/kg (SI Appendix, Fig. S5). Shallow intraplantar (i.pl.) injection of rMg1a (200 pmol) in mice did not cause spontaneous nocifensive behavior (sum of pain-behavior counts at 30 min for saline versus rMg1a, P = 0.58, t test, t = 0.5712, degrees of freedom [df] 10) (Fig. 4A and B). However, it caused a significant decrease in paw-withdrawal threshold to both mechanical (P < 0.01 to P < 0.0001, two-way ANOVA, Sidák’s multiple comparisons) and thermal (P < 0.05 to P < 0.0001, two-way ANOVA, Sidák’s multiple comparisons) stimuli (Fig. 4 C and D). This hypersensitivity was delayed in onset (beginning at 2 to 4 h after injection) and long-lasting (3 to 4 d after injection). Over the time course of this experiment, we observed no visual signs of inflammation (e.g., redness or swelling), and paw thickness measured at 3 d following injection was not significantly different between saline- and rMg1a-treated mice (P = 0.16, t test, t = 1.501, df 10), indicating that the hypersensitivity elicited by injection of rMg1a was independent of overt inflammation.

These data are consistent with our hypothesis that Mg1a has evolved specifically as a defensive weapon against mammalian predators and reveal that its likely function is to cause long-lasting hypersensitivity at the injection site.

**Origin of Weaponized EGF-like Peptides in Ant Venoms.** We next investigated how broadly EGF-like peptides were found in the venoms of ants and other taxa. We began by exploring other Myrmecia species (subfamily Myrmeciinae). To this end, we generated a venom-gland transcriptome of Myrmecia chrysogaster, the gold-tailed bull ant, which is also endemic to Eastern Australia. We identified a single, highly expressed (7,303 TPMs), ortholog of Mg1a that we named MHTX2-Mc1a. This ortholog shared the same precursor architecture as Mg1a and the mature peptide was also analogous to vertebrate EGF-like peptide hormones (SI Appendix, Fig. S6 A and B). As was the case with Mg1a, several EGF-like hormones of marsupials were among the top hits to Mc1a (the mature peptide) in a blastp search of the NCBI nr database.

We then explored ant venoms outside of the subfamily Myrmeciinae. In a venom-gland transcriptome of Rhytidoponera metallica (subfamily Ectatomminae), the greenhead ant, which is widely distributed in Australia, we identified a suite of precursors encoding EGF-like peptides with similar signal peptide sequences and the same precursor architecture as Mg1a (SI Appendix, Fig. S6A), indicative of homology and consistent with a role in venom. A search (using blastp) of the NCBI nr database suggested that the R. metallica peptides were most closely related to vertebrate betacellulin (SI Appendix, Fig. S6A), while PCA suggested an intermediate positioning between the invertebrate EGF-like domains Vein and Spitz and the vertebrate EGF-like domains (SI Appendix, Fig. S6C). Another EGF-like peptide,
Mr1a (UniProt accession no. A0A6G9JKM3; European Molecular Biology Laboratory accession no. QIQ51456.1), has been reported in the venom of the Eurasiatic ant *Manica rubida* (subfamily Myrmicinae) (6). Analysis of its precursor sequence and architecture revealed that it too belongs to the same gene family as Mg1a (SI Appendix, Fig. S6A). In the case of Mr1a, however, according to both blastp and PCA, the mature peptide sequence was clearly most similar to that of insect Spitz, as was the peptide’s intercyte spacing (SI Appendix, Fig. S6A and C). Thus, multiple species of ant have EGF-like peptide toxins in their venoms—in *Myrmecia* these venom peptides have evolved to resemble the EGF-like hormones of these ants’ vertebrate predator(s), while in *M. rubida* the venom peptide resembles the insect hormone Spitz and presumably reflects selection for efficacy in this ant’s insect predators/prey.

To further investigate the origin of EGF-like peptide toxins in ant venoms, we used tblastn to search (using the Mg1a, Mc1a, and Mr1a precursor protein sequences as queries) the NCBI TSA Database and the Hymenoptera Genome Database (16), which included genomic datasets from 58 species of Hymenoptera including 24 ant species. We identified additional homologous precursor sequences from genomes or whole-body transcriptomes of five other ant species (*Myrmica salicinus, Myrmica rubra, Formica aquilonia, Pogonomyrmex californicus, and Pogonomyrmex barbatus*). No homologous ant venom EGF-like toxin sequences were detected in hymenopterans outside of the formicoid clade of the Formicidae, suggesting that within Hymenoptera, the EGF-like toxin gene family is a unique innovation of the formicoid ant lineage. Some of the detected sequences encoded mature peptides which, like Mr1a, were insect-like, while other sequences encoded mature peptides which, like Mg1a and Mc1a, were vertebrate-like (according to blastp searches of the NCBI nr database). However, of the predicted vertebrate-like mature peptides, none exhibited the similarity to marsupial monotreme EGF-like hormones that we observed for the *Myrmecia* venom peptides.

The similarity of the venom peptide Mr1a to the insect hormone Spitz was suggestive of an evolutionary link between the ant EGF-like venom peptide toxins and the endogenous insect hormone; that is, the ant venom EGF-like gene family appears to be derived from the spitz gene. We were unable to resolve the phylogeny of the ant venom EGF-like toxins, presumably due to saturation effects (SI Appendix, Fig. S7). However, endogenous Spitz hormones from diverse hymenopterans grouped closely together on short branches indicative of minimal sequence divergence. By contrast, the ant venom EGF-like toxins, which were derived from a much narrower range of species, exhibited high levels of sequence divergence—a signature of many gene families that have been co-opted for use in venoms (17, 18), and suggestive of functional divergence. The high level of sequence divergence in the ant venom EGF-like peptides may have been driven by selection for efficacy at the receptor(s) of different predators or prey. For example, the similarity of Mr1a from *M. rubida* to insect Spitz suggests that this peptide has retained its ancestral activity, to target the insect ErbB receptor of this species’ insect predators/prey, either as a defensive or predatory weapon. On the other hand, the structural and functional similarity of Mg1a and Mc1a from *Myrmecia* spp. to vertebrate EGF-like hormones suggests that they evolved as a defensive weapon to target the ErbB receptor(s) of the ant’s vertebrate predator(s).

Outside Formicidae, there have been reports of EGF-like peptides in the venoms of sea anemones of the genus *Stichodactyla* (19–21). These venom peptides share a similar “refined” precursor architecture to the ant venom EGF-like peptides. Analysis of the mature peptide sequences revealed highest similarity to the arthropod EGF-like peptide Vein (SI Appendix, Fig. S6B), suggesting that the receptor for these venom peptides is the arthropod ErbB homolog and that their function is for predation or defense against arthropods. This represents an independent convergent incorporation of EGF-like peptides into venom, the origins of which are unclear given the reported absence of an endogenous EGF signaling system in Cnidaria (9).

**Discussion**

Here, we present the discovery and characterization of an EGF-like peptide from venom of the ant *M. gulosus*, which is a potent ligand of the mammalian ErbB1 receptor and causes long-lasting hypersensitivity in mammals. Most algesic venom toxins characterized to date target ion channels or receptors in the nervous or cardiovascular system, or target cell membranes. The exploitation of ErbB signaling by a toxin in order to cause pain extends the known molecular mechanisms of envenomation beyond conventional neurotoxins, cardiotoxins, and membrane...
perturbants. Moreover, while most described venom toxins have an immediate and direct effect, the delayed and long-lasting hypersensitivity caused by Mg1a is, to our knowledge, unprecedented.

In the context of defense, the advantage of causing immediate nociception is easily understood. Indeed, stings of *Myrmecia* do cause immediate intense pain, which results from the action of one or more membrane-targeting peptides (4). However, the advantages of a delayed and indirect, albeit long-lasting, effect are less obvious. One explanation for this effect is that an initial dose of Mg1a, as might occur in an attack by a predator on a *Myrmecia* nest, would cause long-lasting hypersensitivity and perhaps reduce the duration of the attack, discourage future attacks, or enhance the algesic actions of subsequent exposure to other pain-causing venom peptides. Further direct observations of *Myrmecia*-predator interactions are necessary to test this hypothesis.

One important outcome of this study is the implication of ErbB signaling in mammalian pain—more specifically long-lasting hypersensitivity. The fact that the exploitation of this signaling pathway for defensive purposes in *Myrmecia* has been naturally selected for implies that it may play a fundamental role in nociception in mammals, one that until recently has gone largely unnoticed. Mammalian EGF-like peptide hormones have mostly been studied in the context of development, although they also play important physiological roles in adults. For example, some are major components of wound fluid (22) and are involved in epithelialization in skin wound healing (23). ErbB receptor inhibitors are used for the treatment of a range of cancers, and several case reports have indicated that ErbB receptor inhibition may relieve certain cancer pain (for a review, see ref. 24). Recent independent studies have reported roles for epiregulin-, EGF-, and HBEGF-ErbB signaling in wound (25–27) and, as in the present study, each reported a specific role in hypersensitivity.

From these studies, we can begin to formulate a hypothesis for the role played by ErbB signaling in pain. Following an injury, high concentrations of EGF-like peptide hormones may be released at the wound site where they serve a critical role in wound healing and, independent of inflammation, sensory neurons innervating the site of injury may become sensitized via activation of ErbB receptors expressed on nerve terminals or cells interacting with these. While long-term pain is often viewed as pathological, hypersensitivity of a wound site may be adaptive (i.e., for protection of a wound during healing). Mg1a in the venom of *M. gulosa* may have evolved to exploit this pain pathway in the absence of a genuine injury. Thus, activation of ErbB by certain EGF-like peptide hormones may represent an important noninflammatory mediator of wound hypersensitivity, and inhibition of these pathways may be an effective treatment for certain pain pathologies.

Our results allow us to build a hypothesis of the evolution of EGF-like peptide toxins in the venoms of ants. We can deduce that early in the formicoid lineage, a duplication of a gene, possibly that encoding the peptide hormone Spitz, allowed its modification for direct secretion (i.e., loss of domains other than the signal and mature peptide or recombination of the mature peptide with another signal peptide) and recruitment to the venom. Supporting this interpretation, we found that the venom of *M. rubida* contains a peptide that closely resembles endogenous Spitz, suggesting retention of its ancestral function targeting the insect ErbB receptor but which is now directed exogenously for predation and/or defense against other insects. In *Myrmecia*, and possibly other lineages of ants, selection pressure for defense against one or more vertebrate predators resulted in the evolution of the venom peptide to resemble, both structurally and functionally, a nociceptive peptide hormone of the predator(s).

This scenario draws strong parallels with the recent discovery of an insulin-like toxin in the venom of the fish-hunting cone snail *Conus geographus*, that has convergently evolved to mimic, structurally and functionally, the endogenous insulin hormone of its fish prey (28). When hunting, the snail releases the venom insulin into the water, where it acts on the fish’s endocrine system to cause sedation, thus facilitating prey capture.

Both of these toxins are examples of convergent evolution and, more specifically, of biological mimicry—where one organism (the mimick) gains a selective advantage by mimicking an aspect (or multiple aspects) of another organism (the model) in order to deceive it or another (the dupe). The scenario involving the cone snail and the fish is a form of aggressive molecular mimicry, known as Batesian–Wallacian mimicry (a bipolar aggressive system in which the model belongs to the dupe’s species and the dupe is deceived by the model). The scenario of the bull ant and its predator(s) is different. It is a rare form of mimicry known as Gilbertian mimicry, a bipolar protective system in which the model belongs to the dupe’s species and the dupe is deceived by the model (29). Gilbertian mimicry has been described in certain species of the plant genus *Passiflora*, where parts of the plants have evolved to visually mimic *Heliconius* butterfly eggs that are near to hatching (30). To avoid intraspecific competition, *Heliconius* females avoid ovipositing on plants already bearing such eggs (or egg mimics) and thus the plant is afforded protection (31). In this study, the scenario presented is analogous to the Gilbertian mimicry described in *Passiflora* and *Heliconius*, yet it occurs not at the organismic level but at the molecular level. Here, a molecule (Mg1a) produced by the ant *M. gulosa* mimics an endogenous nociceptive molecule of its vertebrate predator(s), and its injection causes long-lasting hypersensitivity and presumably a reduction in predation, thus protecting the ant. While Gilbertian mimicry appears to be a rare phenomenon (until now it had only been described in the *Passiflora–Heliconius* interaction), we predict that more examples of this type of mimicry may have evolved, particularly in cases where nociceptive molecules are concerned.

This study illustrates how natural selection can mold the gene product of one organism to resemble that of another (in this case, the gene product of the prey resembles that of the predator). It is an example of both convergent evolution and an unusual form of molecular mimicry, one that brings to mind the ancient proverb “To know your enemy, you must become your enemy.”

**Materials and Methods**

**Collection of Venom and Purification of Mg1a.** Eight adult female *M. gulosa* were collected from a single site near Brisbane, QLD, Australia. Venom was acquired by inciting ants to sting a thin layer of paraffin, which was rinsed with purified water.

Approximately 500 μg of venom was separated on a Phenomenex Gemini NX-C18 column (250 × 4.6 mm; particle size 3 μm; pore size 110 Å) using a gradient of 5 to 80% solvent B (90% acetonitrile [ACN], 0.05% trifluoroacetic acid) over 60 min at a flow rate of 1 mL min⁻¹. Fractions were collected on the basis of absorbance at 214 nm and matrix-assisted laser desorption ionization mass spectrometry was used to confirm the identity of the peptide(s) present in each fraction. The fraction corresponding to Mg1a was quantified via absorbance at 280 nm (NanoDrop 2000C spectrophotometer; Thermo Fisher Scientific), divided into aliquots, and dried by vacuum centrifugation.

The complete primary structure of Mg1a was confirmed by liquid chromatography-mass spectrometry analysis of native, reduced, and alkylated peptide. A 2-μg aliquot of Mg1a was analyzed on an AB Sciex TripleTOF 5600 mass spectrometer equipped with a TurboV source heated to 550 °C. A second 2-μg aliquot was first reduced and alkylated (incubated in 50 mM ammonium carbonate, pH 11, 48.25% ACN, 1% 2-iodoethanol, 0.25% triethylphosphine for 1 h at 37 °C) and analyzed on the same instrument. Experimental MS/MS spectra of native, reduced, and alkylated Mg1a were compared against theoretical peak lists for each peptide generated using MS-Product in ProteinProspector v5.22.1 (https://prospector.ucsf.edu/prospector/).

A peptide toxin in ant venom mimics vertebrate EGF-like hormones to cause long-lasting hypersensitivity in mammals.
Principal-Component Analysis. Sequences of EGF, HBEGF, betacellulin, epiregulin, and Mesothelin were downloaded from UniProt, and were clustered to the EGF-like domain of each. Misannotated and fragmented sequences were removed and identical sequences were clustered using CD-hit v4.6.3 (2). The remaining 313 unique sequences were aligned using local pairwise alignment (L-INS-i) with the regional alignment tool (v0.2) in MAFFT v7.304b (33). After adjusting the alignment of structurally conserved cysteine residues, we then clustered and projected the biophysical properties of each sequence in a multidimensional sequence space as described previously (12). Briefly, the alignment was converted into a numerical matrix and analyzed using PCA to summarize the main covarying sets of properties using sequence space (R) codes obtained from GitHub, https://github.com/T5404-SeeSpace. The variables used were molecular mass (Da), net charge (coulombs), hydrophobicity (Doolittle index), disorder propensity (TOP-IDP), disulfide potential (binary descriptor), and occupancy (binary descriptor).

Recombinant Expression of Mg1a. A synthetic gene encoding Mg1a, with codons optimized for high-level expression in Escherichia coli, was produced by GeneArt (Life Technologies). The synthetic gene also incorporated a tobacco etch virus (TEV) protease recognition site immediately N-terminal to the Mg1a gene to minimize the presence of additional nonnative residues after protease cleavage. The synthetic gene was inserted into a pLICC expression plasmid (pMCSG9) (34), which enables expression of Mg1a as a His6-tagged maltose-binding protein (MBP) fusion protein, with the incorporated TEV protease recognition site between the MBP- and Mg1a-coding regions. E. coli (BL21(DE3)) were transfected with 10 ng/mL plasmid, incubated on ice, and then heat-shocked. Cells were plated on Luria–Bertani (LB) agar containing 100 μg/mL ampicillin (Astral Scientific) and grown for 18 h. Suitable single colonies were used to spike LB broth containing 100 μg/mL ampicillin, and propagated overnight at 37°C before being induced for expression induction. A modified autoinduction medium was used based on ZYP5052 (35), where lactose was replaced by a low concentration of isopropyl β-D-thiogalactopyranoside (IPTG) (36), resulting in increased expression yields. The medium, referred to as ZYP5050 (due to the presence of 0.5% [weight/volume; wt/vol] glycerol, 0.05% [wt/vol] glucose, 0.0% lactose, and the presence of IPTG), contained 10 g/L tryptone, 5 g/L yeast extract, 54.3 mM glycerol, 2.8 mM MgSO4, 0.1% FeCl3, 1 μM MnCl2, 1 μM ZnSO4, 0.2 μM CoCl2, 0.2 μM CuCl2, 0.2 μM NaMoO4, 0.2 μM NaSeO3, 0.2 μM H3BO3, 40 μg/mL IPTG, 100 μL/l Antifoam SE-15 (Sigma), and 100 μg/mL ampicillin. The culture was incubated at 30°C, with shaking at 300 rpm, until the optical density at 600 nm (OD600) reached 0.8 to 1.2, as determined via spectrophotometry (SmartSpec 3000; Bio-Rad). The temperature was then lowered to 17°C and cells were cultured for a further 22 h with shaking at 300 rpm. After incubation, cells were harvested and lysed, and Mg1a was purified as outlined above. The extent of 12C and 13N incorporation was determined via electrospray ionization MS/MS.

NMR Spectroscopy and Structure Determination. NMR samples containing 15N-labeled rMg1a were prepared by dissolving lyophilized peptide in 300 μL of NMR buffer (20 mM sodium phosphate, pH 6.0, 5% 2H2O) at a concentration of 350 μM. For resonance assignments, spectra were acquired at 293 K on a Bruker Avance III 600-MHz spectrometer equipped with a cryogenically cooled probe. Two-dimensional [15N-1H]-HSQC, 3D HNCO, 3D HNCA, 3D CBCA(CO)NH, 3D HBHA(CO)NH, and 3D HNCACB spectra were used to generate backbone-resonance assignments while 3D HCCH-correlation spectroscopy and 2D [13C-1H]-HSQC spectra were used for side-chain carbon and proton assignments. Nonuniform sampling data were reconstructed with compressed sensing using qMDD, uniform sampling data were processed in TOPSPIN (v3.2), and chemical shift analysis was performed using CcpNmr Analysis (v2.1.3), with the pyMOSFET (v2.1.3–01) software package to model the 1H–1H NOE and determine the NOE evolution rate. All other processing was performed with Mnce Eos. Synchronizing the NOE evolution rate with the NOESY cross-peak intensities allowed the sensitivity of the NOESY spectrum to be increased by 10X.

Bioluminescence Resonance Energy Transfer Assay. The plasmids pcdNA3-ErbB1/Rluc8 and pcdNA3-Grb2/Venus have been described previously (43) and were gifts of Kevin Pfleger, Harry Perkins Institute of Medical Research, University of Western Australia, Perth, WA, Australia. HEK293T cells were seeded at 25,000 cells per well in poly-o-lysine (0.1 mg/mL; Sigma-Aldrich; P6407)−coated white 96-well plates 24 h before transfection. FuGene (Promega; E2311) was used, as per the manufacturer’s instructions, to transfect 5 μg of plasmid (0.5% HEPES, 50 mM KCl, 50 mM NaPO4, 1 mM MgSO4, 5 μM FeCl3, 2 μM CaCl2, 1 μM MnCl2, 1 μM ZnSO4, 0.2 μM CoCl2, 0.2 μM CuCl2, 0.2 μM NaMoO4, 0.2 μM NaSeO3, 0.2 μM H3BO3, 40 μg/mL IPTG, 100 μL/l Antifoam SE-15 (Sigma), and 100 μg/mL ampicillin. The culture was incubated at 30°C, with shaking at 300 rpm, until the optical density at 600 nm (OD600) reached 0.8 to 1.2, as determined via spectrophotometry (SmartSpec 3000; Bio-Rad). The temperature was then lowered to 17°C and the culture was grown until the OD600 was >5 (~21 h later). The cells were then harvested via centrifugation (6,200 × g, 20 min, 4°C), the supernatant was discarded, and the pellet was dissolved in Tris HCl NaCl (TN) buffer (20 mM Tris, 200 mM NaCl, pH 8.0).

The cell pellet was homogenized and lysed using a constant-pressure cell disruptor (25 to 32 kpsi; TS Series; Constant Systems; 6 to 14°C). The cell lysate was then centrifuged (39,000 × g, 30 min, 4°C) and the soluble lysate was collected. The cell pellet was then incubated at 37°C for 3 h with 100 μL of NMR buffer (20 mM sodium phosphate, pH 6.0, 5% 2H2O) at a concentration of 350 μM. For resonance assignments, spectra were acquired at 37°C and cells were cultured via centrifugation, the supernatant was discarded, and pellets were resuspended in 250 mL phosphate buffer (5 mM Na2HPO4, 50 mM KH2PO4, 60 mM NaH2PO4) to remove any residual rich media. Cells were pelleted again via centrifugation, the supernatant was discarded, and pellets were resuspended in 250 mL autoinduction medium. The minimal autoinduction medium, which we called P-2050I (due to the presence of 0.2% [wt/vol] glycerol, 0.05% [wt/vol] glucose, 0.0% lactose, and the presence of IPTG), contained 1 mM MgSO4, 10 μM FeCl3, 4 μM CaCl2, 21.7 mM [13C]Glucose, 2.8 mM [13C]Glucose, 9.2 mM [15N4]NH4Cl, 5 mM Na2SO4, 50 mM KH2PO4, 60 mM NaH2PO4, 1 mM MgSO4, 5 μM FeCl3, 2 μM CaCl2, 1 μM MnCl2, 1 μM ZnSO4, 0.2 μM CoCl2, 0.2 μM CuCl2, 0.2 μM NaMoO4, 0.2 μM NaSeO3, 0.2 μM H3BO3, 40 μg/mL IPTG, 100 μL/l Antifoam SE-15 (Sigma), and 100 μg/mL ampicillin. The culture was grown at 30°C for 2.5 h before the temperature was switched to 17°C and cells were cultured for a further 22 h with shaking at 300 rpm. After incubation, cells were harvested and lysed, and Mg1a was purified as outlined above. The extent of 12C and 13N incorporation was determined via electrospray ionization MS/MS.

EGF Receptor Transfection and Western Blot. HEK293 cells were seeded on 12-well plates and grown at 37°C, 5% CO2 until they reached ~80% confluency. Transient transfection was achieved through incubation with 15 ng of plasmid containing human ErbB1 and Lipofectamine LTX reagent diluted in Opti-MEM. At 5 h posttransfection, the medium was aspirated off and replaced with Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 1-glutamine and 10% fetal bovine serum. Cells were grown for a further 24 h, and then the medium was aspirated off and replaced with serum-free DMEM to remove growth factors present in the medium. After 12 h, cells were stimulated for 5 min with either hEGF (R&D Systems; 236-EG), or rMg1a, or
PBS (negative control). Following stimulation, cells were placed on ice, washed twice with cold PBS, and lysed using RIPA lysis buffer (50 mM Tris HCl, pH 7.5, 100 mM NaCl, 2 mM ethylenediaminetetraacetate [EDTA], 50 mM NaF, 0.1% [wt/vol] sodium dodecyl sulfate, 0.5% [wt/vol] sodium deoxycholate, 1% [vol/vol] Triton X-100, 20 mM sodium pyrophosphate, 10 mM sodium orthovanadate) as well as complete EDTA-free protease inhibitor mixture (Roche) and a PhosSTOP phosphatase inhibitor tablet (Roche). The cell lysates were gently rocked for 1 h at 4 °C and centrifuged (15,000 × g for 15 min), and the supernatant was collected and frozen until analysis.

The supernatants were thawed, electrophoresed on a sodium dodecyl sulfate–polyacrylamide gel (12.5% resolving gel), and then transferred to a polyvinylidene difluoride membrane via electrophoretic transfer. Membranes were washed with PBS containing 0.1% Tween 20 (PBS/Tween) and then blocked with 1:1 PBS/Tween and intercost (PBS) blocking buffer (LI-COR Biosciences, 927-700001) to prevent nonspecific binding. Membranes were then incubated overnight at 4 °C with primary antibodies against glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and phosphorylated Erk1 and Erk2. The following morning, membranes were washed three times and then incubated with secondary anti-mouse and anti-goat fluorescent antibodies for 2 h before once again washing three times and visualizing using an Odyssey CLx near-infrared imaging system.

Venom-Gland Transcriptome of M. chrysogaster. A single M. chrysogaster worker was collected near Logan, QLD, Australia. The venom duct, venom reservoir, and venom glands (together referred to as the venom apparatus) were dissected out and placed in RNAlater. Total RNA was extracted from the venom apparatus using TRIZol reagent (Thermo Fisher Scientific), according to the manufacturer’s instructions. Complementary DNA library preparation and sequencing were performed by the Institute for Molecular Bioscience Sequencing Facility at The University of Queensland. A dual-indexed library was constructed using the TruSeq-3 Stranded mRNA Sample Prep Kit (Illumina) with oligo(dT) selection and an average insert size of 180 bp; 150-cycle paired-end sequencing was performed on an Illumina NextSeq 500 instrument. Adapter trimming of demultiplexed raw reads was performed using fastqtrim (44), followed by quality trimming and filtering using PRINSEQ-lite (45) and error correction using BBnorm tdapole, a part of the BBTools package. Trimmed and error-corrected reads were assembled using Trinity v2.4.0 (46) with a k-mer value of 1e 29 and a UniRef90 database. Estimates of transcript abundance were made using the RSEM (48) plugin of Trinity (align_and_estimate_abundance).

Insect Assays. House crickets (A. domesticus; average mass 50 mg; Pisces Live Food) were injected intraabdominally with 2 μL water or rMg1a (doses ranging from 1.4 to 41.5 nmol/g). Following injection, crickets were placed on their back and the time taken to right themselves was recorded.

Sheep blowflies (L. cuprina; average mass 25.2 mg) were cooled on dry ice for 3 min and then injected into the lateral thorax with 2 μL water or rMg1a (41.5 nmol/g) using a Terumo insulin syringe (29-gauge needle) attached to an Arnold hand microapplicator (Burkard Manufacturing). Survival of injected flies was monitored at 10 min, 30 min, 1 h, and 24 h postinjection, with paralysis (twitching, but inability to walk or fly) and lethality recorded.

Plant-paralyzing experiments. Male 5-wk-old C57BL/6j mice used for behavioral experiments were purchased from the Animal Resources Centre (WA, Australia). Mice were housed in groups of up to four per cage, maintained on a 12-h light-dark cycle, and fed standard rodent chow and water ad libitum. RmG1a (200 pmol) diluted in saline containing 0.1% bovine serum albumin (BSA) was administered in a volume of 20 μL to the hind paw by shallow i.pl. injection. Negative-control animals were injected with saline containing 0.1% BSA. Following injection, spontaneous pain-behavior events were counted from video recordings by a researcher blinded to the treatment.

Prior to mechanical and thermal threshold testing, animals were acclimated (30 min) to the MouseMet test enclosures containing a bar bottom to permit access to the plantar hind paw. Measurements were performed at 1 h, 2 h, 4 h, 1 d, 2 d, 3 d, 4 d, and 7 d after i.pl. injection of rMg1a (dose) or saline containing 0.1% BSA. Mechanical paw-withdrawal thresholds were measured using an automatic von Frey apparatus (MouseMet; Topcat Metrology) as previously described (49). Thermal threshold measurements were performed using an automated MouseMet thermal apparatus (Topcat Metrology) as previously described (50). All measurements were performed by a researcher blinded to the treatment. Baseline paw-withdrawal thresholds were determined 3 d prior to i.pl. injection.

At each time point, animals were visually inspected for inflammation of the injected paw (e.g., swelling and reddening) and, at 3 d after injection, the thickness of both hind paws was measured using electronic calipers.

Animal experiments were approved by The University of Queensland Animal Ethics Committee (Approval no. 2021/AE00448).

Phylogenetic Analysis. Sequences of ant EGF-like toxin precursors and Spitz (truncated to signal and mature peptides only) were aligned using the LINS-i algorithm of MAFFT v7.309 (33). We selected the most appropriate evolutionary model (JTTDCMut+G4) using ModelFinder (51) before we used IQ-TREE v2.0.6 (52) to construct the molecular phylogeny by maximum likelihood, estimating branch support values by ultrafast bootstrap using 10,000 replicates (53). Trees were visualized in Archaeopteryx v0.9921 (54).

Statistics. All statistical analyses were done using GraphPad Prism (v9.0.0). For concentration–response curves of the ligand-induced BRET ratio, data were fitted to a nonlinear regression with variable slope. For quantitative Western blot data, an ordinary one-way ANOVA with Sidák’s multiple-comparisons test was used to test significance between EGF- or rMg1a-treated and negative-control cells. For the insect paralysis assay, an ordinary one-way ANOVA with Sidák’s multiple-comparisons test was used to test significance between rMg1a-treated and negative-control animals. For the insecticidal assays, data were plotted using survival curves. For analysis of spontaneous pain, a two-way ANOVA with Sidák’s multiple-comparisons test was used to test significance over the time course of the experiment and an unpaired t test was used to test significance of the sum of pain-behavior counts at 30 min between treated and negative-control animals. For measurements of both mechanical and thermal hypersensitivity, a two-way ANOVA with Sidák’s multiple-comparisons test was used to test for significance between rMg1a-treated and negative-control animals. For paw-thickness measurements, an unpaired t test was used to test for significance between treated and negative-control animals.

Data Availability. The precursor sequence of Mg1a reported in this article has been deposited in UniProt (accession no. PD5L4A). The solution structure and chemical shifts of Mg1a have been deposited in the PDB (ID code7R6P) and Biological Magnetic Resonance Data Bank (ID code30928), respectively. All study data are included in the article and/or SI Appendix.

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