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A tetrapeptide class of biased analgesics from an Australian fungus targets the μ-opioid receptor


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An Australian estuarine isolate of Penicillium sp. MST-MF667 yielded 3 tetrapeptides named the bilaids with an unusual alternating LDLD chirality. Given their resemblance to known short peptide opioid agonists, we elucidated that they were weak (K<sub>L</sub> low micromolar) μ-opioid agonists, which led to the design of bilorphin, a potent and selective μ-opioid receptor (MOPr) agonist (K<sub>i</sub> 1.1 nM). In sharp contrast to all-natural product opioid peptides that efficaciously recruit β-arrestin, bilorphin is G protein biased, weakly phosphorylating the MOPr and marginally recruiting β-arrestin, with no receptor internalization. Importantly, bilorphin exhibits a similar G protein bias to oliceridine, a small nonpeptide with improved overdose safety. Molecular dynamics simulations of bilorphin and the strongly arrestin-biased endomorphin-2 with the MOPr indicate distinct receptor interactions and receptor conformations that could underlie their large differences in bias. Whereas bilorphin is systemically inactive, a glycosylated analog, bilactorphin, is orally active with similar in vivo potency to morphine. Bilorphin is both a unique molecular tool that enhances understanding of MOPr biased signaling and a promising lead in the development of next generation analgesics.

Significance

Agonists of the μ-opioid receptor (MOPr) are currently the gold standard for pain treatment. However, their therapeutic usage is greatly limited by side effects including respiratory depression, constipation, tolerance, and dependence. Functionally selective MOPr agonists that mediate their effects preferentially through G proteins rather than β-arrestin signaling are believed to produce fewer side effects. Here, we present the discovery of 3 unusual tetrapeptides with a unique stereochemical arrangement of hydrophobic amino acids from an Australian estuarine isolate of Penicillium species. Building on these natural templates we developed bilorphin, a potent and selective highly G protein-biased agonist of the MOPr. Further, through the addition of a simple sugar moiety, we generated bilactorphin that is an effective analgesic in vivo.


Competing interest statement: Patent Application(s) corresponding to Australian Patent Application 2018091944 The University of Sydney and The University of Queensland has been filed concerning this peptide. Title: Analgesics and Methods of Use Thereof. This article is a PNAS Direct Submission. This open access article is distributed under Creative Commons Attribution-NonCommercial-NoDerivatives License 4.0 (CC BY-NC-ND).

Development of ligands that target G protein-coupled receptors (GPCRs) in multiple functional states has attracted great interest, particularly with increasing knowledge of GPCR structure (1–3). These novel ligands are expected to underpin the development of agonists with superior pharmacologically relevant properties, including biased receptor signaling (4, 5), whereby one downstream signaling pathway is favored over another. For example, biased agonists that signal by differentially recruiting G proteins over β-arrestin to the μ-opioid receptor (MOPr) could deliver better analgesics, based on the view that down-regulating β-arrestin recruitment diminishes adverse side effects (4, 6–8).

Exploiting this concept, the G protein-biased MOPr agonist oliceridine (TRV130) is a potent analgesic in rodents, with lower respiratory depression and gastrointestinal dysfunction compared to morphine (7). Indeed, human clinical trials of oliceridine show reduced respiratory impairment compared to morphine administered at equi-analgesic doses (9). Reduced respiratory depression delivers improved safety, potentially reducing the burden of opioid overdoses, now at epidemic proportions in many jurisdictions (10).

Bioactive peptides display great promise for their novel pharmacological properties (11). Since the discovery of the relatively nonselective mammalian opioid peptides, the enkephalins, other endogenous mammalian ligands, including the tetrapeptide endomorphins that target the MOPr with high selectivity over the related κ-opioid (KOPr) and δ-opioid (DOPr) subtypes, have been found (12, 13). Natural peptide agonists containing a D-Ala in the second position formed by a posttranslational modification isolated from frog skin, deroorphin and deltorphin II, selectively target MOPr and DOPr, respectively (14). Similar synthetic modifications have yielded enhanced biological stability and receptor selectivity. For example, introduction of ω-Ala stabilizes the enkephalins to proteolysis, and further substitutions yield highly stable, selective MOPr agonists such as DAMGO ([ω-Ala2, N-MePhe4, Gly5-ol]-enkephalin) (15). However, to our knowledge, all endogenous opioid peptides acting on MOPr robustly recruit arrestins and produce MOPr internalization (16–19). Here, we report the discovery of 3 tetrapeptides, bilaids A–C (Fig. 1, [1a–3a]), from an Australian estuarine isolate of Penicillium sp. MST-MF667.
[initially reported as *P. bilaii* (20); maximum likelihood tree presented in SI Appendix, Fig. S1]. Discovery of the bilaid, which resemble known opioid peptides but featuring an unusual alternating sequence of antipodal amino acids (LDLD), led to our design of bilorphin (3e), a MOPr agonist, with G protein signaling bias similar to oliceridine. Bilorphin adopts a distinct conformational shape and intermolecular interactions in molecular dynamics (MD) simulations of the bilorphin–MOPr complex, consistent with predicted G protein bias at related GPCRs. Together with its in vitro and in vivo activity, we demonstrate that bilorphin provides a scaffold for the development of stable, orally active opioid peptides that are biased toward G protein signaling (3g).

**Results**

To date, the major opioid peptide families have only been detected in vertebrates (21). Through a broader screening program, we now report the discovery of 3 closely related tetrapeptides (Fig. 1), bilaid A (1a, FvVf-OH), B (2a, FvVv-OH), and C (3a, YvVf-OH), from the Australian estuarine-derived *Penicillium sp.* MST-MF667. Solvent extracts of cultivations of MST-MF667 were subjected to solvent partition followed by reversed-phase high-performance liquid chromatography (HPLC) to yield 1a (0.15%), as well as traces of 2a (0.0018%) and 3a (0.0008%) (Fig. 1). The chemical structures for 1a–3a, which resemble opioid peptides albeit with a unique, alternating LDLD amino acid configuration, were identified by spectroscopic analyses, chemical derivatization and degradation, and Marley’s analysis and total synthesis. In addition to synthesizing 1a–3a, we prepared and screened a series of related analogs for inhibition of forskolin-induced cAMP formation in HEK cells expressing the human MOPr (hMOPr) (SI Appendix, Table S1). Five hits determined to be active at a concentration of 10 μM were tested for competitive binding against the hMOPr agonist [3H]DAMGO. Bilaid C (3a) bound with the highest affinity at the hMOPr (Kₐ = 210 nM, Fig. 2A and SI Appendix, Table S1), consistent with the requirement for a Tyr moiety in the majority of peptide ligands targeting opioid receptors (22). Bilaid A (1a) showed more modest affinity (Kᵢ = 3.1 μM) that was improved 4-fold through C-terminal amidation (Kᵢ = 0.75 μM, 1e), prompting us to apply this modification to further analogs. Analogos of bilaid A having a DLLDL (1b, 1f), LLDDL (1c, 1g), DDLL (1d), or LLLL (1h) configuration were generally inactive, with the exception of FVVFNH₂ (1b), which weakly inhibited cAMP formation at 10 μM. These findings are consistent with previous SAR studies, which revealed that having α-amino acids in position 2, other than those that are Nα-alkylated (e.g., L-Pro), reduce opioid agonist activity (23), as do α-amino acids in position 1 (24). Other bilaid analogs showed no MOPr activity at 10 μM.

C-terminial amidation of bilaid C (Fig. 2A, 3b and SI Appendix, Table S1) improved hMOPr affinity (Kᵢ = 93 nM), consistent with the increased binding of 1e over 1a. Dimethylation of the N-terminal tyrosine residue (3e) resulted in significantly increased potency (Kᵢ = 1.1 nM), as has been observed for other opioid peptides (25). This compound, which we name bilorphin, bound with nearly 200-fold selectivity for hMOPr over hDOPr (Kᵢ = 190 nM) and 700-fold selectivity over hKOPr (Kᵢ = 770 nM) (Fig. 2A and SI Appendix, Table S1).

To assess functional activity of bilorphin, patch-clamp recordings of G protein-activated, inwardly rectifying potassium channel (GIRK) currents were made in rat locus coeruleus (LC) neurons, which natively express MOPr but not DOPr or KOPr (26). Bilorphin acted as an agonist with potency greater than morphine. Its actions were completely reversed by the highly MOPr selective antagonist CTAP (27) (n = 9, Fig. 2B and C and SI Appendix, Table S1), establishing that bilorphin does not act on the closely related nociceptin or somatostatin (SST) receptors expressed by LC neurons (28). The native peptide, bilaid C, was 14-fold weaker than morphine in the same assay system (29). Bilaid C analogs with an acetylated N terminus (3d), or containing all α-amino acids (3e and 3f), were inactive in LC neurons at 10–30 μM, confirming the importance of a free N terminus and the native LDLD motif for maintaining MOPr activity (SI Appendix, Table S1).

![Fig. 1. Structures of bilaid, bilorphin, and bilactorphin.](image-url)
The relative potency and intrinsic efficacy of bilorphin in signaling pathways was examined in AtT20 cells stably expressing FLAG-tagged mouse MOPr (mMOPr) to enable bias analysis (Fig. 2 C and E). To reliably determine the relative intrinsic efficacy of bilorphin to activate G proteins (G\textsubscript{GIRK}), we partially, irreversibly inactivated receptors to reduce the maximum response to the high efficacy natural agonist, Methionine5-enkephalin (Met-enk), to 80% of that produced by SST acting on native SST receptors in the same cells (Fig. 2 C and E). Maximal activation of SST receptors normally produces a G\textsubscript{GIRK} increase equivalent to a maximal activation of MOPr (30). Under these conditions, bilorphin, morphine, and endorphin-2 all displayed similar maximal responses. All agonists displayed similar potencies in the brain slice and cell assay, with the exception of Met-enk, the reduced potency of which in brain slices is known to result from its degradation by peptidases (31). As expected from brain slice experiments, all 3 opioids were moderately efficacious but less so than Met-enk (Fig. 2E and SI Appendix, Fig. S2A). In contrast, the G protein-biased, small molecule agonist oliceridine, activated G\textsubscript{GIRK} significantly less efficaciously than either morphine or bilorphin (Fig. 2E).

MOPr C-terminal phosphorylation, \(\beta\)-arrestin recruitment, and internalization are thought to contribute to on-target opioid analgesic side effects so that G protein-biased opioids that avoid \(\beta\)-arrestin signaling should show an improved side effect profile (4, 7, 8). We therefore assayed the activity of bilorphin for inducing C-terminal phosphorylation, \(\beta\)-arrestin recruitment, and MOPr internalization in the same AtT20 cell line. Agonist-induced phosphorylation of residue serine 375 (Ser375) drives \(\beta\)-arrestin recruitment and internalization (32). We determined bilorphin-induced phosphorysine Ser375 (pSer375) phosphorylation using a phosphosite-specific antibody (Fig. 3 A and D) (33). Surprisingly, and unlike other opioid peptides (19, 32, 33), bilorphin produced low levels of pSer375 (30 \(\mu\)M, Fig. 3 A and D and SI Appendix, Fig. S2B), which appeared less than that produced by morphine, but this difference was not statistically significant (SI Appendix, Fig. S2B). Using MOPr-luciferase and \(\beta\)-arrestin 2-YFP constructs, a bioluminescence resonance energy transfer (BRET) assay was performed to determine \(\beta\)-arrestin 2 recruitment (Fig. 3 B and D) (34). Similar to phosphorylation, saturating concentrations of bilorphin induced very low levels of BRET (33), significantly less than that produced by morphine (30 \(\mu\)M, Fig. 3B). MOPr internalization was assessed immunocytochemically after 30 min of agonist treatment (Fig. 3 C and D). Bilorphin produced almost no detectable internalization of MOPr, compared to low levels induced by morphine and robust internalization driven by both endomorphin-2 and Met-enk (Fig. 3D). In summary, when normalized to the maximum response to Met-enk in each pathway, bilorphin displayed similar maximal G protein efficacy to morphine with progressive reduction in relative efficacy across pathways from Ser375 phosphorylation, \(\beta\)-arrestin recruitment to internalization (Fig. 3D), suggesting that bilorphin is a G protein-biased opioid.

Operational analysis, the de facto standard for quantifying biased signaling (35, 36), suggests that bilorphin is G protein biased relative to morphine (SI Appendix, Fig. S2) but this requires accurate determination of EC\textsubscript{50}, which was impractical for bilorphin due to its very low internalization efficacy (SI Appendix, Fig. S2) (35). Calculation of relative efficacy from maximum response in each pathway provides a complementary estimate of bias in signaling assays where all agonists are partial (36, 37). This approach (SI Appendix, Fig. S2 and Table S2) substantiated the G protein bias of bilorphin relative to morphine (19, 34). To further substantiate the bias of bilorphin compared with morphine and the established G protein biased agonist oliceridine, both of which produce very little internalization, we enhanced internalization by overexpressing GRK2-YFP and \(\beta\)-arrestin 2-HA (38). In GRK2-positive cells, morphine, oliceridine, and bilorphin all produced clear internalization signals (SI Appendix, Fig. S3A). Quantification shows that even under these amplified conditions, bilorphin induces similar MOPr internalization to oliceridine but significantly less than morphine at saturating concentrations (SI Appendix, Fig. S3 B and C). Calculation of bias relative to morphine with the \(\Delta E_{\text{max}}\) method suggests that bilorphin exhibits significant G protein bias that is similar to oliceridine, further establishing it to be a G protein biased opioid (SI Appendix, Fig. S3C).

The widely accepted mechanism for ligand bias is the stabilization of distinct GPCR conformations that favor coupling to different intracellular proteins (39). However, how such biased agonists interact with the receptor binding pocket and the nature of these biased receptor conformations, particularly for the MOPr, is poorly understood. To this end, we performed molecular docking and MD simulations (40) to compare possible biased receptor conformations of the G protein-biased bilorphin versus \(\beta\)-arrestin–biased endorphin-2. First, we analyzed the predicted binding poses of bilorphin and endorphin-2 after 1 \(\mu\)s of MD simulation to determine if interactions with the MOPr binding pocket could differentiate these oppositely biased compounds. The binding pose of bilorphin is shown in Fig. 4A and SI Appendix, Fig. S4A and that of endorphin-2 in Fig. 4B and SI Appendix, Fig. S4B. There were similarities in how bilorphin and endorphin-2 interacted with the MOPr; both peptides bound in the orthosteric pocket with the phenol groups of the Dmt (bilorphin) or Tyr1 (endorphin-2) orientated toward the intracellular side of MOPr, and maintained interactions with Asp147, Trp293, and His297 (Fig. 4C). For the latter
residue, bilorphin switched between a direct interaction and hydrogen bonding via a bridging water molecule (Fig. 4A. Inset). However, there were also important differences in how these peptides interact with the MOPr binding pocket. For bilorphin, the rest of the tetrapeptide chain extended out toward the extracellular side of MOPr, making contacts with residues at the top of TMs 1, 2, and 7, resulting in bilorphin, but not endomorphin-2, interacting with Tyr75. Conversely, endomorphin-2 extended toward ECL1 and ECL2 and the top of TM2. Endomorphin-2 therefore interacted with the extracellular loops, contacting W133ECL2 in ECL1 for the entire simulation time, and transiently interacting with Cys217ECL2, Thr218ECL2, and Leu219ECL2 of ECL2. Whereas, these interactions were absent for bilorphin. Moreover, endomorphin-2 made a greater number of contacts with residues in TM3 and TM5 than bilorphin.

These predicted binding poses were validated by root mean square deviation analysis showing stability of each ligand in the binding pocket (SI Appendix, Fig. S4 C and D), and our docking and MD protocol successfully recapitulated the binding pose of DAMGO when compared to the cryo-electron microscopy structure of the DAMGO-bound MOPr-Gi complex (SI Appendix, Fig. S5) (41).

Next, principal component analysis was employed to examine the overall conformational changes in the receptor transmembrane helices in the presence of bilorphin or endomorphin-2. The receptor conformations at each time point were projected onto principal components (PC) 1 and 2 and plotted in Fig. 4D. PC1 and PC2 accounted for 28.9% and 10.9% of the variance, respectively. Both peptide–MOPr complexes sampled conformations across PC2, but clustered differently based on PC1, suggesting that the MOPr adopted distinct conformations when bound to each biased agonist.

As depicted in SI Appendix, Fig. S4E, PC1 primarily described alternative conformations in the extracellular region of the receptor close to the orthosteric binding site and, to a lesser extent, differences in the intracellular portions of the helices. With endomorphin-2 bound, there was an overall contraction of the orthosteric site, due to inward movements of TMs 2, 3, 4, and 7. Whereas, with bilorphin bound, there was a bulging of the middle portion of TM1 and a shift outwards from the helix bundle, relative to the endomorphin-2–bound receptor. These alternative conformations of the helices were also reflected in the volume of the orthosteric site, as with bilorphin bound the binding pocket volume was on average 1.6 times greater than with endomorphin-2 bound (42) (SI Appendix, Fig. S4F). On the intracellular side of MOPr, PC1 described inward movements of TMs 5, 6, and 7 with endomorphin-2 bound, compared to the bilorphin-bound MOPr (SI Appendix, Fig. S4E). The result was a more occluded intracellular cavity in the endomorphin-2–bound MOPr, compared to that with bilorphin bound.

Analysis of the MD data therefore suggests that the different ligand–residue interactions for these oppositely biased peptides (with respect to the extracellular loops and TM1) may lead to the alternative receptor conformations described by the principal component analysis and, hence, the opposing bias profiles of bilorphin and endomorphin-2.

We then evaluated bilorphin in vivo. Bilorphin failed to inhibit noiception in the hotplate test in mice when administered subcutaneously (s.c.) (100 mg/kg, n = 4) or intravenously (i.v.) (50 mg/kg n = 4). By contrast, bilorphin was antinociceptive after intrathecal injection (5 nmol per mouse, peak effect 41 ± 9% MPE n = 4, versus 0 ± 1.5% for vehicle, n = 4), suggesting the lack of systemic activity is due to poor penetration of the blood–brain barrier (BBB). We therefore developed several bilorphin analogs with substitutions thought to enhance BBB permeability, including glycosylation near the C terminus (43). The diglycosylated analog, bilorphin (3g), was a potent analgesic after systemic administration (s.c.: ED50 of 34 μmol/kg, 95% confidence interval [CI] = 28–40 μmol/kg, SI Appendix, Fig. S6 A and B), was nearly equipotent with morphine (ED50 of 27 μmol/kg, 95% CI = 24–30 μmol/kg SI Appendix, Fig. S6B) and was antagonized by coadministration of the opioid antagonist, naltrexone (SI Appendix, Fig. S6C). Bilorphin was also active after i.v. (peak effect of 88.9 ± 11.8 versus 14.4 ± 1.8% MPE for vehicle, n = 3–4) or oral administration (SI Appendix, Fig. S7). In contrast, the monoglycosylated analog 3h was systemically inactive, consistent with previous modified opioid peptides (43). These findings establish that the natural LDLD opioid peptide backbone we have
discovered is a viable framework for development of G protein-biased opioid analgesics. Like bilorphin, bilactorphin was a potential partial opioid agonist of G proteins in AtT20 cells but exhibited a small loss of potency compared with bilorphin (SI Appendix, Fig. S6 C and D). Bilactorphin did, however, display increased internalization and β-arrestin recruitment compared to bilorphin (SI Appendix, Fig. S6 E and F) and thus might not be suitable for directly testing the role of bias in opioid side-effect profile. Alternative modifications that do not disrupt G protein bias of the parent bilorphin might be better suited to enhance BBB permeability.

**Discussion**

Nature has inspired many of the most well-known and widely used analgesics, from natural salicin in willow (Salix) bark to synthetic aspirin, from opioid poppy alkaloids such as morphine and codeine to synthetic hydrocodone (Vicodin), oxycodone (OxyContin), and buprenorphine (Subutex). Notwithstanding their use in alleviating pain, serious adverse side effects, combined with the challenge of addiction, abuse, and acquired tolerance, render these analgesics (particularly opioids) far from ideal. There is an urgent need to discover and develop new, safer and more efficacious analogues, with mechanisms of actions that mitigate against risk.

We therefore investigated the analgesic potential of a class of tetrapeptides, the bilaidas (1a–3a), isolated from a Penicillium fungus. Taking advantage of an unprecedented natural scaffold comprising alternating LDLD configuration amino acids, which imparts inherent biostability, we designed a peptide-based G protein-biased MOPr agonist, bilorphin (3e). Furthermore, we assembled proof-of-concept data that this pharmacophore can be optimized to yield an orally active MOPr agonist analgesic, bilactorphin (3g).

G protein-biased opioid agonists have been proposed as a route to improving therapeutic profile (4, 7, 8). Among known peptide opioid agonists, which typically are biased toward β-arrestin signaling relative to morphine (19, 34), the pharmacological profile of bilorphin is most unusual, although a synthetic opioid cyclopeptide with G protein bias was recently reported (44). Bilorphin enjoys an opioid signaling bias comparable to oliceridine, a G protein-biased drug candidate in phase III clinical trials. G protein-coupled receptor depletion with a cAMP assay system (37). Similar low efficacy results have recently been reported for another opioid, PZM21, also claimed to be safer than morphine (4, 51). Furthermore, it is difficult to evaluate maximal β-arrestin efficacy of novel biased agonists in other studies because assays were insensitive to the relatively low G protein efficacy of morphine (7, 8). Very low G protein efficacy may indeed be a confounding factor in the preclinical and clinical studies of side effect profile, given that other agonists with very low G protein efficacy such as buprenorphine are not strongly G protein biased (19) but are well characterized to produce less respiratory depression, and overdose death than highly efficacious agonists such as morphine and methadone (52). Because bilorphin is strongly G protein biased and has nearly equivalent maximal G protein efficacy to morphine, further development of BBP penetrating analogs that can release the parent molecule will facilitate direct test of the influence of bias without being confounded by differing G protein efficacy.

Finally, we have elucidated an analgesic pharmacophore based on the discovery of the bilaidas and related bilorphin and bilactorphin motifs. Unusually, they derive from a microbial source, an untapped resource for analgesics and worthy of further investigation. This observation suggests that microbes may be an untapped resource for analogues, deserving of further investigation.

**Experimental Procedures.** Full details on the materials and methods used are available in SI Appendix. Solvent extracts solid phase cultivations of MST-MF667 were subjected to solvent partition followed by reversed-phase HPLC, and chemical structures were identified on the basis of detailed spectroscopic analysis, chemical derivatization and degradation, Marley's
analysis and total synthesis. All peptides were assembled manually by stepwise solid-phase peptide synthesis. MOPr activity was initially screened using competition opioid radioligand binding to membranes from cultured cells expressing hMOPr, hDOPr, or hKOPr receptors, then agonist activity screened at hMOPr using inhibition of forskolin-stimulated CAMP formation. Agonist activation of MOPr-coupled G protein channels was then quantified using superfusion onto LC neurons in rat brain slices using whole-cell patch clamp recording. Signaling pathway analysis was quantified in AT20 cells stably expressing mMOPr using perforated patch recording for G protein channel activation (G protein signal), fluorescence immunohistochemistry for Ser375 phosphorylation and MOPr internalization and arrestin recruitment with a BRET-based approach. Molecular docking was performed using the Bristol University Docking Engine (53). The selected peptide–MOPr complexes were then embedded in a lipid and cholesterol bilayer and used in all-atom MD simulations. Multiple simulations (8 × 125 ns), with different initial velocities, were performed under the Amber ff14SB and Lipid14 forcefields, to yield a total of 1 μs of trajectory data for each peptide. Behavioral assays of analgesia were performed using the hotplate latency assay in mice. Statistical analyses were performed as described in SI Appendix. All values are expressed as means ± SEM, except where noted otherwise (SI Appendix, Fig. S2).

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