















RESEARCH ARTICLE

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[View Journal](#) | [View Issue](#)Cite this: *RSC Med. Chem.*, 2025, 16, 1172Novel sigma 1-antagonists with *cis*-(+)-normetazocine scaffold: synthesis, molecular modeling, and antinociceptive effect†Giuliana Costanzo, ^a Giuseppe Cosentino, ^a Margherita Grasso, ^b Vincenzo Patamia, ^a Sara Zuccalà, ^a Alessandro Coco, ^a Elisabetta Novello, ^c Mahmoud Al-Khrasani, ^d Raffaele Morrone, ^e Giovanni Mario Pitari, ^c Emanuele Amata, ^a Agostino Marrazzo, ^a Antonio Rescifina, ^a Lorella Pasquinucci ^{‡*a} and Carmela Parenti ^{‡a}

Inflammatory pain represents one of the unmet clinical needs for patients, as conventional therapies cause several side effects. Recently, new targets involved in inflammatory pain modulation have been identified, including the sigma-1 receptor ($\sigma 1R$). Selective $\sigma 1R$ antagonists have demonstrated analgesic efficacy in acute and chronic inflammatory pain models. Considering these findings, a series of novel *N*-normetazocine derivatives has been designed and synthesized to investigate the pivotal role of *N*-normetazocine stereochemistry in their pharmacological fingerprint. The affinity profile of new ligands *versus* sigma receptors and opioid receptors was evaluated *in vitro*, and compound **7** showed a relevant $\sigma 1R$ affinity, with $K_{\sigma 1} = 27.5 \pm 8.1$ nM, and selectivity over sigma-2 receptor ($\sigma 2R$) and opioid receptors. Furthermore, *in vivo*, compound **7** significantly reduced inflammatory pain in the second phase of the formalin test. Molecular modeling studies were also performed to analyze the binding mode and the key interactions between the new ligands and $\sigma 1R$.

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1. Introduction

Nowadays, sigma receptors (σR s) are considered promising therapeutic targets for the management of various debilitating disorders, including pain, depression, and neurodegenerative diseases.^{1,2} Initially, σR s were presented as subtypes of the opioid receptors; now, they are being reclassified as a unique receptor type.³ Two subtypes are identified and termed sigma-1 receptor ($\sigma 1R$) and sigma-2 receptor ($\sigma 2R$), different in structure, biological functions, and pharmacological profile. In particular, $\sigma 1R$ is a unique ligand-regulated chaperone membrane receptor, considered a regulatory subunit that modulates target protein functions,

thus playing a crucial role in various biological pathways.⁴ For instance, it can modulate opioid receptors,⁵ voltage-gated calcium channels,⁶ and the NMDA receptor activity.⁷ In particular, $\sigma 1R$ activation is considered a valid strategy to treat neurodegenerative disorders,⁸ while $\sigma 1R$ blockage is evaluated as an innovative approach in the oncology area for managing different cancers such as glioblastoma, pancreas, breast, and prostate tumors.⁹ Moreover, owing to its distribution in key regions of pain control, such as dorsal root ganglia, spinal dorsal horn, periaqueductal grey matter, locus coeruleus, and rostroventral medulla, $\sigma 1R$ antagonists have a central role in pain modulation.¹⁰ In fact, $\sigma 1R$ agonists counteract opioid receptor-mediated analgesia, acting as an endogenous anti-opioid system, whereas $\sigma 1R$ antagonists determined a marked morphine-induced antinociception enhancement, both at central and peripheral levels.¹¹ This observation is supported by further preclinical studies showing that $\sigma 1R$ antagonists produced antinociceptive effects in persistent pain conditions.¹² In this regard, it has been reported that the intrathecal blockade of $\sigma 1R$ prevents nerve injury-induced hyperalgesia, and the $\sigma 1R$ knockout (KO) mice and the treatment of wild-type animals with $\sigma 1R$ antagonists can counteract inflammatory pain behaviors evoked by carrageenan^{13,14} or complete Freund's adjuvant.¹⁵

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Furthermore, σ 1R antagonists showed antiallodynic and antihyperalgesic effects in sensitizing conditions without modifying the normal pain behavior.¹⁰

The role of σ 1R activation in inflammatory pain could be linked to its capability to enhance Ca^{2+} release, induced by bradykinin and nitric oxide (NO) signaling with a reinforcement of the inflammatory process.¹⁶ Pain sensitization after peripheral inflammation also involves plastic changes defined by increased spinal excitatory neurotransmission and activation of different kinases, like ERK1/2, modulated by σ 1R.¹⁷ Moreover, σ 1R plays a role in pain modulation at the spinal level by activating neurosteroids.¹⁸ In light of this, developing novel σ 1R antagonists is challenging in medicinal chemistry and is considered a good strategy to counteract persistent pain.

To develop new σ 1R ligands, we evaluated the profile of newly synthesized *N*-substituted normetazocine compounds **6–15** (Fig. 1). In particular, the (+)-*cis*-normetazocine scaffold was chosen due to its affinity for σ 1R, and it is known that (+)-SKF-10,047 was the first drug discovered to interact with σ 1R.¹⁹ Carrol *et al.* developed numerous analogs of (+)-SKF10,047, and the impact of altering its *N*-substituent with respect to the ligand's affinity and selectivity was evaluated. In particular, it has emerged that lipophilic *N*-substituents may improve σ 1R selectivity and potency, and the *N*-benzyl compound showed the best σ 1R affinity.²⁰ In a successive study,²¹ the *ortho*-, *meta*-, and *para*-substituted *N*-benzyl-analogues of (+)-*cis*-normetazocine were synthesized, and this study highlighted both the influence of substituent position and substituent volume. For instance, the analog with the F-atom in *para*-position retained the high σ 1R affinity profile of the lead compound. Thus, *para*-position and low steric hindrance analogs with *N*-benzyl-substituent were preferred. To obtain new insights into the structural requirements for σ 1R interaction, a series of 2-aminoethyl derivatives of *cis*-(+)-*N*-normetazocine was synthesized.²² Bearing a second basic nitrogen, these compounds displayed a good affinity for σ 1R while showing a reduced σ 2R/ σ 1R selectivity. Moreover, a compound with 2-(*N*-phenylcarbamoyl)ethyl substituent, (+)-LP1, was also synthesized, and it demonstrated a nanomolar σ 1R affinity and a potent analgesic effect in a mouse model of inflammatory pain. However, a low selectivity ratio to σ 2R was highlighted (σ 2R/ σ 1R = 10).²³ Subsequently, we synthesized the *cis*-(+)-*N*-normetazocine derivative of (+)-LP2 (Fig. 1), bearing an *N*-phenethylnormetazocine and a methoxy group at

the benzylic carbon. (+)-LP2 showed high affinity for σ 1R and selectivity *versus* σ 2R. Moreover, (+)-LP2 has shown a σ 1R antagonist profile with analgesic effects in both inflammatory and neuropathic animal pain models.^{24,25} (+)-LP2 was used as a diastereomeric mixture of (+)-(2'*R*)-LP2 and (+)-(2'*S*)-LP2.

Our new compounds **6–15** were designed as derivatives of (+)-LP2 and here we investigated the influence of various structural modifications. In particular, we examined the impact of altering the chain length of the *N*-substituent from 2 to 4 carbon atoms, assessed the importance of the phenyl ring by substituting it with a proton, and explored the effects of replacing the methoxy group with either an ester or carboxylic acid function. With respect to the methoxy group substitution of the (+)-LP2, we aimed to examine its impact on σ 1R affinity by introducing ester and acid functionalities that show different hydrogen bonding capabilities. A molecular modeling analysis was also performed to analyze the binding mode and the key interactions between the new ligands and σ 1R.

2. Results and discussion

2.1. Synthesis

The new compounds **6–15** were synthesized according to Scheme 1. Compounds **1–3** were obtained as previously described.²⁶ Briefly, the compound **1** was prepared by primary alcohol sulfonylation of ethyl tropate with methanesulfonyl chloride and triethylamine in CH_2Cl_2 . The ethyl 4-chloro-2-(*R/S*)-phenylbutanoate and the ethyl 5-chloro-2-(*R/S*)-phenylpentanoate were synthesized starting with a suspension of ethyl phenyl cyanoacetate in potassium *t*-butoxide in anhydrous DMF by adding 1-bromo-2-chloroethane and 1-chloro-3-iodopropane, respectively. The subsequent decarboxylation reaction on relative intermediate compounds in a saturated aqueous solution of K_2CO_3 and CH_3OH , followed by hydrolysis of their cyano group in a 1 : 1 mixture of HCl (12 N) and $\text{CH}_3\text{CH}_2\text{OH}$, allowed the compounds **2** and **3** to be obtained. Target esters **6–10** were prepared by alkylation of (+)-*cis*-normetazocine with the alkylation agents **1–5**, respectively. The acid derivatives **11–15** were synthesized by ester saponification. Compounds **6–15** possess a chiral stereocenter at the *N*-substituent of (+)-*cis*-normetazocine scaffold and exist as a mixture of diastereoisomers, which could not be separated using our purification procedures. An HPLC analysis on diastereomeric mixture of final compound **7** was performed with a Chiralcel

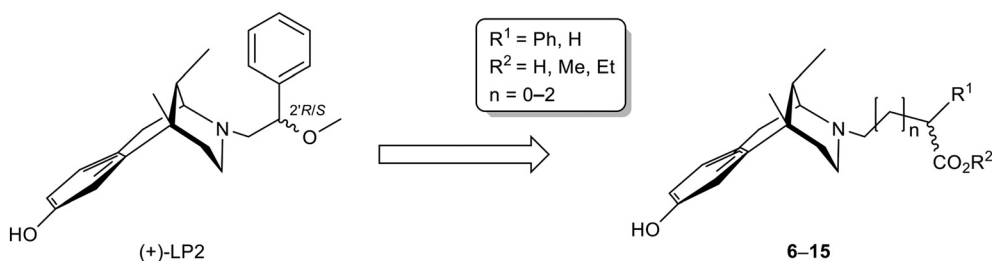
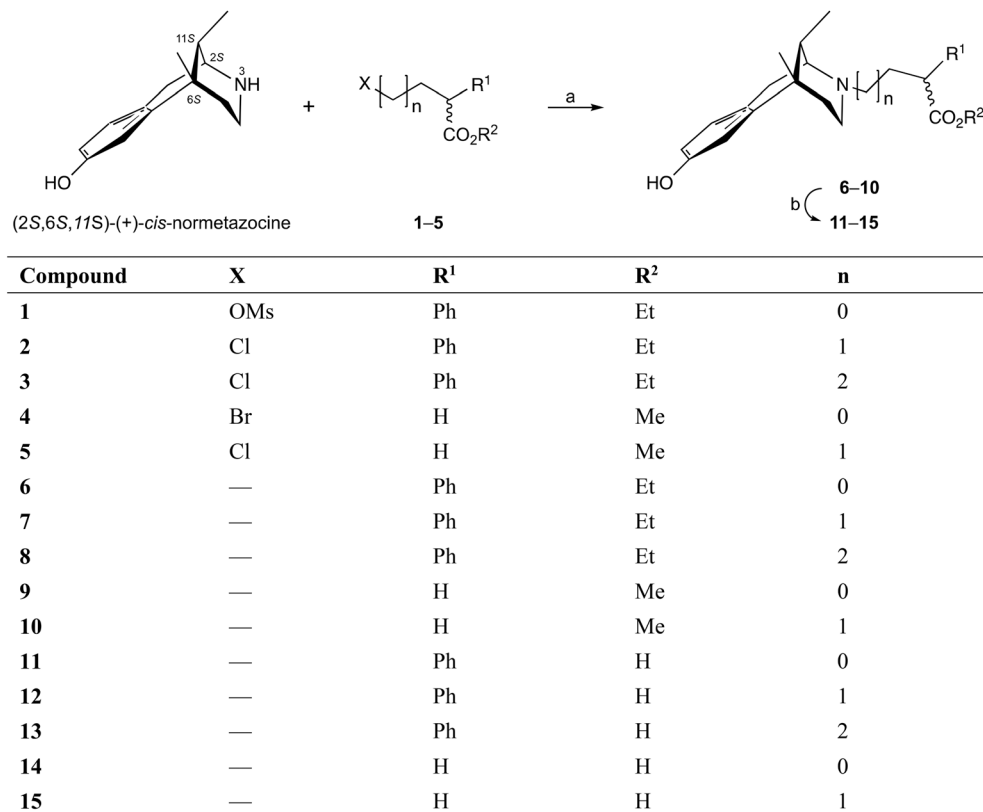


Fig. 1 Chemical structures of (+)-LP2 and the newly synthesized compounds.





Scheme 1 Synthesis of target compounds 6–15. Reagents and conditions: a) NaHCO₃, KI, DMF, 55 °C, 24 h; b) NaOH 1 N, 110 °C, 5 h.

OJ column to verify the diastereoisomers ratio. HPLC analysis highlighted the separation of the diastereoisomers of compound 7 with a peak at $t_R = 12.3$ min (55.6%) and a peak at $t_R = 14.6$ min (44.4%) (Fig. S1†). All compounds were characterized by ¹H NMR, ¹³C NMR, attached proton test (APT), and elemental analysis.

2.2. Binding properties on σ Rs and opioid receptors

Target compounds 6–15 were tested *versus* σ 1R and σ 2R with the [³H]-(+)-pentazocine and [³H]-1,3-di-*ortho*-tolylguanidine ([³H]-DTG) radio probes, respectively (Table 1). Competitive inhibition curves of compounds 7, 8, and 10–11 at 10^{−10}–10^{−5} M concentration range were reported in Fig. S2†. Inhibition constant (K_i) values were obtained using nonlinear regression analysis GraphPad Prism, version 7.0 (GraphPad Software Inc., San Diego, USA). Compared to (+)-LP2 ($K_{i\sigma 1} = 26.6 \pm 2.4$ nM), only compound 7 maintained high σ 1R affinity with a K_i value of 27.5 ± 28.1 nM. Shortening or increasing spacer length in compounds 6 and 8 significantly increased their K_i values *versus* σ 1R. Compound 10, with a similar distance between normetazocine scaffold and ester functionality but lacking the phenyl ring in the *N*-substituent, exhibited a moderate σ 1R affinity ($K_i = 159 \pm 23$ nM). Instead, compound 9, with a methylene deletion in the *N*-substituent spacer, showed negligible affinity for σ 1R. Replacing the ester group with the corresponding carboxylic acid group in compounds 11–15 was not tolerated, and all compounds showed

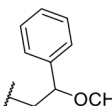
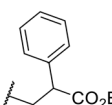
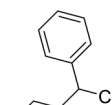
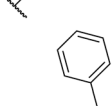
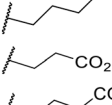
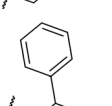
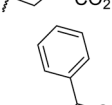
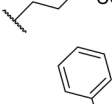
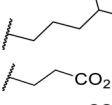
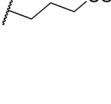
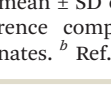
negligible σ 1R affinity. Regarding σ 2R, all new compounds have shown negligible binding affinity. Moreover, compounds 6–15 were tested *versus* mu-opioid receptor (MOR), delta-opioid receptor (DOR), and kappa-opioid receptor (KOR) with the [³H]-DAMGO, [³H]-Deltorphin and [³H]-U69,593 radio probes, respectively, with K_i values >5000.

2.3. Molecular modeling studies

To rationalize ligand-receptor interactions within the receptor site, *in silico* and molecular modeling studies have been performed, thanks to which we are better able to understand how the nature of the substituents of each ligand may influence the interaction with the various residues in the receptor pocket. *In silico* studies were carried out with AutoDock, integrated into the YASARA program, on compounds that demonstrated noteworthy inhibitory activity experimentally, *i.e.*, 7, 8, 10, and 11. At physiological pH (7.4), all the compounds had positively charged nitrogen, and for compound 11, the acidic function is deprotonated. As in our previous study, LP2 was used as a reference compound.²¹ Considering the protonated nitrogen atom as an asymmetric center, we first docked the four (+)-LP2 stereoisomers to identify the most suitable ones for further molecular modeling studies. The (2′S,3R)-(+)-LP2 and (2′R,3R)-(+)-LP2 stereoisomers had the computed K_i value closest to the observed one, with (2′S,3R)-(+)-LP2 being the more active of the two. Therefore, subsequent docking experiments focused



Table 1 σ 1R and σ 2R binding affinity of compounds 6–15

Compound	N-Substituent	K_i (nM) \pm SD ^a	
		σ 1R	σ 2R
(+)-LP2 ^b		26.6 \pm 2.4	2393 \pm 514
6		>10 000	>10 000
7		27.5 \pm 8.1	>1000
8		594.0 \pm 168	>10 000
9		>10 000	>10 000
10		159.0 \pm 23	>10 000
11		664.0 \pm 200	>10 000
12		>10 000	>10 000
13		>10 000	>10 000
14		>10 000	>10 000
15		>10 000	>10 000
(+)-Pentazocine DTG ^c		4.3 \pm 0.5 124.0 \pm 19	1465 \pm 224 18.0 \pm 1

^a Each value is the mean \pm SD of at least two experiments performed in duplicate. Reference compounds were tested with the same membrane homogenates. ^b Ref. 24. ^c 1,3-Di-*ortho*-tolylguanidine.

on the (2'*S*,3*R*)- and (2'*R*,3*R*)-stereoisomers, maintaining the nitrogen stereocenter as 3*R*. The stereoisomers with lower predicted K_i values are listed in Table 2. However, the inhibitory activity between the (2'*S*,3*R*)- and (2'*R*,3*R*)-stereoisomers did not differ significantly across most compounds. For σ 1R, a detailed analysis of the docking poses within the receptor site has been performed for compounds 7 and 10, which possess an ΔG of -10.21 and -9.05 kcal mol⁻¹, respectively.

Fig. 2 illustrates the 3D and 2D poses of the two most active compounds within the receptor site of σ 1R. Compound 7 (Fig. 2a) establishes a dense network of interactions within

the receptor cavity, similar to typical σ 1R inhibitors.^{27,28} It forms hydrogen bonds with the Asp126 residue and electrostatic interactions with Glu172. Additionally, the non-polar portion of compound 7 engages in hydrophobic interactions with various amino acid residues. In contrast, compound 10 (Fig. 2b) demonstrates lower inhibitory activity, lacking the typical interaction with Asp126. However, it forms a hydrogen bond and an electrostatic interaction with Glu172. Furthermore, compound 10 establishes an unconventional hydrogen bond and an electrostatic-type interaction with Tyr103, π - π interactions with Trp89 and Phe107, and several hydrophobic interactions with other residues.

Molecular modeling studies support that these *N*-substituents can establish interactions with key amino acid residues of the σ 1R receptor site. As shown in Fig. 2, which presents the poses of the two ligands within the receptor site, the presence of the phenyl group in compound 7 increases the number of hydrophobic interactions. Specifically, compound 7 can interact with residues Leu182 and Met93 through its phenyl group, interactions that do not occur with compound 10. Additionally, the presence of the aromatic group allows ligand 7 to adopt a conformation within the receptor site that enables it to form a crucial interaction with the residue Asp126, an interaction absent in compound 10.

2.4. Chemical stability of compound 7

Compound 7's chemical stability was evaluated *in vitro* at 37 °C in aqueous phosphate buffer (PBS) at pH 7.4 using HPLC with a diode array detector (DAD). Across all incubation times, the concentration of compound 7 remained constant (Fig. S3 and S4[†]), indicating optimal stability under physiological conditions. It has a half-life ($t_{1/2}$) of 300 min.

2.5. Plasma stability assay of compound 7

Stability studies were performed to determine the decay over time of compound 7 in mouse plasma. As indicated by correspondent chromatograms (Fig. S5[†]), compound 7 in mouse plasma was clearly identifiable as a sharp peak at t_R = 10.52 min (in red box). Compound concentrations remained stable up to 5 min, starting to decline at times ≥ 20 min. Half-life estimations (Fig. S6[†]) indicated that compound 7 was reduced to 50% of its initial concentrations at 31.07 min, supporting the temporal pharmacology of compound 7 in the inflammatory pain model examined in the present studies.

2.6. ADME profile of compound 7

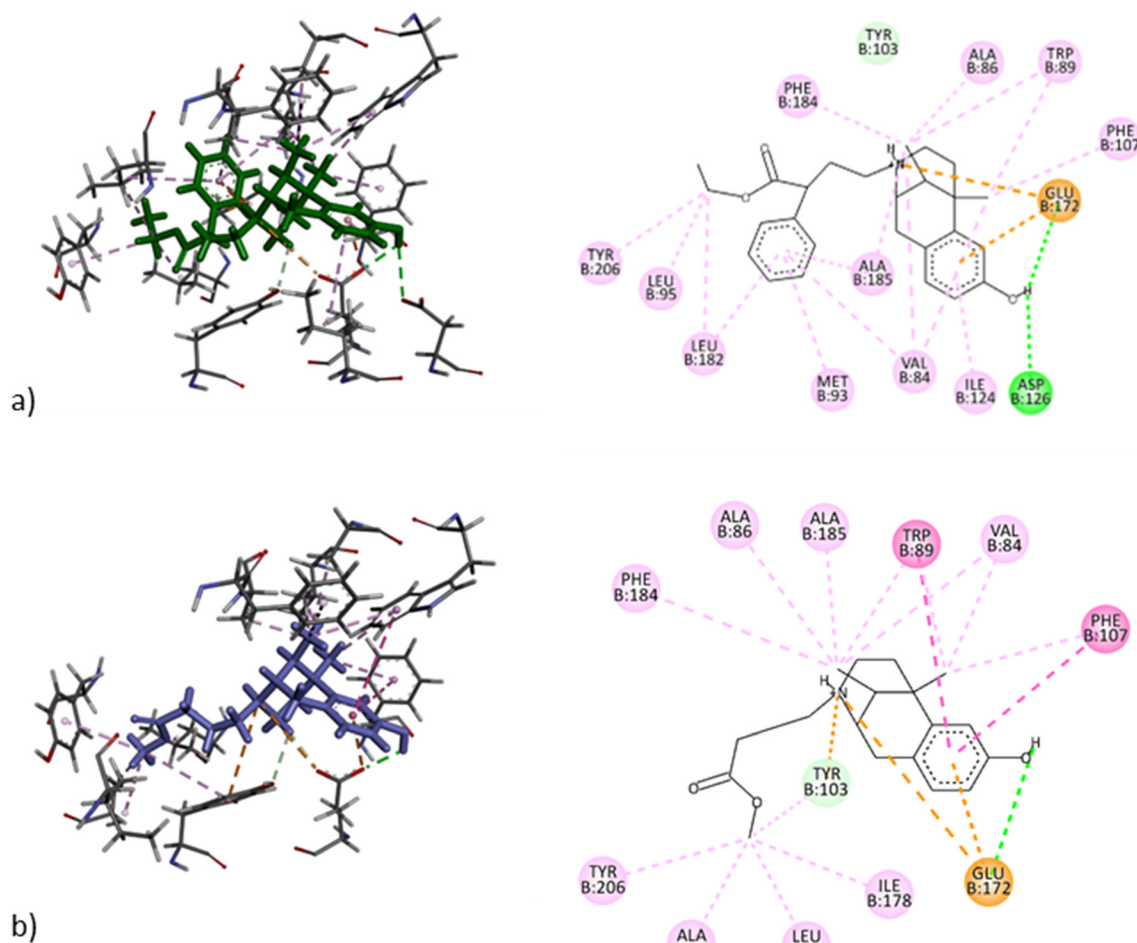
A comprehensive computational prediction of the pharmacokinetic profile and brain/plasma distribution of compound 7 was conducted using the Calculator Plugins of ChemAxon and the SwissADME web tool.²⁹ Notably, with high accuracy, the SwissADME platform incorporates advanced algorithms capable of identifying potential false positive results, a common issue in biochemical assays for small molecules. This provides valuable insights into



Table 2 Calculated free energies of binding, ΔG (kcal mol⁻¹), and constants of binding, K_i (nM), of synthesized compounds versus $\sigma 1R$ and $\sigma 2R$

Compound	Calcd ΔG $\sigma 1R$	Calcd K_i $\sigma 1R$	Exp K_i $\sigma 1R^a$	Calcd ΔG $\sigma 2R$	Calcd K_i $\sigma 2R$	Exp K_i $\sigma 2R^a$
(2'S,3R)-(+)-LP2	-9.70	77.0	26.6 \pm 2.4	-7.40	3741	2393 \pm 514
(2'R,3R)-(+)-LP2	-9.67	81.0		-7.38	3870	
(3'R,3R)-7	-10.21	32.5	27.5 \pm 8.1	-7.14	5803	>1000
(3'S,3R)-7	-10.14	36.6		-7.14	5803	
(4'S,3R)-8	-8.48	603.9	594.0 \pm 168	—	—	>10 000
(4'R,3R)-8	-8.46	624.7		—	—	
(3R)-10	-9.05	230.6	159.0 \pm 23	—	—	>10 000
(2'R,3R)-11	-8.38	715.0	664.0 \pm 200	—	—	>10 000
(2'S,3R)-11	-8.37	727.2		—	—	

^a The experimental values of K_i refer to the mixture of diastereoisomers.

**Fig. 2** 3D and 2D poses of compounds 7 (a) and 10 (b) within the $\sigma 1R$.

compound 7's reliability in terms of pharmacological profiling.

Several key descriptors essential for determining bioactivity parameters were utilized in these predictions, as summarized in Table 3.

Compound 7 was evaluated for drug-likeness according to Lipinski's rule of five, a widely accepted guideline for predicting oral bioavailability. The analysis confirmed compound 7 meets all criteria for orally active drugs without

any violations, reinforcing its suitability for development. It also adheres to Lipinski's³⁰ and Veber's³¹ rules, highlighting its drug-like properties. Veber's rule, in particular, emphasizes the importance of molecular flexibility and polar surface area in drug absorption, parameters in which compound 7 excelled. The compound's bioavailability score of 0.55 indicates moderate oral bioavailability, suggesting it may achieve sufficient systemic exposure when administered orally.



Table 3 Descriptors for compound 7

Descriptor	Value
MW	407.55
Log <i>P</i>	4.57
Log <i>D</i>	2.90
Topological polar surface area	49.77 Å ²
Number of hydrogen bond donors	1
Number of hydrogen bond acceptors	4
Number of nitrogen atom count	1
Oxygen atom count	3
Nitrogen-oxygen atoms count	4
Number of aromatic rings	2
Number of heavy atoms	30
MWHBN	0.20
p <i>K</i> _a	9.37
Solubility	5.23 × 10 ⁻⁵ mg mL ⁻¹ ; 1.28 × 10 ⁻⁷ mol L ⁻¹
Chemaxon BBB score	5.04
Chemaxon CNS MPO score	3.49

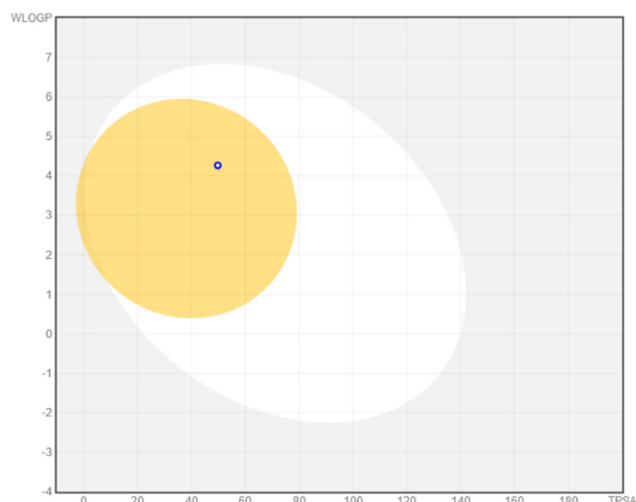
Crucially, compound 7 passed the PAINS (pan assay interference compounds) model,³² which screens for chemical structures likely to interfere with high-throughput biological assays and produce false positive results. The absence of PAINS alerts underscores its potential as a reliable lead compound for further development, free from problematic structures that often complicate drug discovery efforts. These assessments position compound 7 as a promising candidate for drug development, with minimal risks associated with common small-molecule pitfalls.

Moreover, compound 7 was predicted to have moderate water solubility, a favorable attribute for oral delivery, as sufficient solubility is essential for absorption through the gastrointestinal tract. The absorption and distribution characteristics were further analyzed using the extended BOILED-Egg model (Brain Or IntestinaL EstimatedD). This computational tool visually predicts a drug's capacity to

permeate biological barriers. As depicted in Fig. 3, compound 7 resides within the “yellow yolk” region, suggesting a high probability of passive diffusion across the blood–brain barrier (BBB). This indicates strong potential for targeted therapies for the central nervous system (CNS), where BBB penetration is a critical challenge. Simultaneously, the compound is predicted to exhibit high gastrointestinal absorption, further enhancing its feasibility for oral administration. Interestingly, despite its BBB permeability, a blue dot marker in the model indicates that compound 7 is likely a substrate for P-glycoprotein (P-gp), a transporter protein responsible for pumping drugs out of the brain. This suggests that while compound 7 can cross into the CNS, it may be subject to efflux from the brain *via* P-gp, an essential consideration for CNS drug development. This efflux mechanism could limit its retention within the brain, and strategies to modulate or circumvent this effect may be necessary for the future development of CNS-targeted therapies.

The Chemaxon BBB permeability score of 5.04 and CNS MPO score of 3.49 validate the predictions obtained using the SwissADME platform, confirming compound 7's potential for CNS drug development. A BBB score of 5.04 indicates a strong likelihood of the compound crossing the blood–brain barrier, which is critical for CNS-targeted therapies. This aligns with the results from SwissADME, which suggested high BBB permeability. The CNS MPO score of 3.49, while slightly below the ideal threshold of 4, still indicates reasonable suitability for CNS activity. Although this score suggests room for optimization in specific molecular properties, it confirms that compound 7 is a promising candidate for CNS applications, consistent with the SwissADME predictions of its drug-like behavior. Overall, the combination of these scores supports the viability of compound 7 for further development, particularly in therapies aimed at the brain, with potential adjustments to improve its CNS MPO score in future optimizations.

Metabolic stability and interaction with cytochrome P450 (CYP) enzymes were also explored *via in silico* predictions. Compound 7 was identified as an inhibitor of two key CYP isoforms: CYP2D6 and CYP3A4. These isoforms play significant roles in the metabolism of many drugs, so inhibition could lead to potential drug–drug interactions (DDIs) if compound 7 is co-administered with other medications metabolized by these enzymes. However, compound 7 showed no significant inhibition of CYP1A2, CYP2C19, or CYP2C9, indicating a more selective interaction with the CYP system. While the inhibition of CYP2D6 and CYP3A4 suggests that compound 7 could pose DDI risks, this can be managed through careful dosing strategies and further optimization. Understanding these inhibitory interactions is critical for anticipating side effects, managing metabolic liabilities, and ensuring the compound's safety profile in clinical applications.

**Fig. 3** BOILED-Egg plot for compound 7.

2.7. *In vivo* formalin test

Based on competition binding assays and molecular modeling studies, we evaluated the antinociceptive efficacy of compound 7 in the mouse formalin test. The intraplantar (i.pl.) injection of a dilute solution of formalin (5%) into the plantar surface of the mouse hind paw induces a biphasic response characterized by an early nociceptive phase (phase I) and a tonic, inflammatory pain phase (phase II). Phase I is induced by the direct activation of peripheral nerve terminals, while phase II is sustained by mechanisms of central sensitization in the dorsal horn of the spinal cord.

To assess the systemic antinociceptive effect of compound 7 in both phases of the formalin test, vehicle or compound 7 (3, 5, and 7 mg kg⁻¹) was administered intraperitoneally (i.p.) 20 minutes before formalin injection. As shown in Fig. 4, pretreatment with compound 7 at the dose of 3 mg kg⁻¹ i.p. did not significantly modify the nociceptive behavior (licking and flicking or shanking the injected paw) in either phase of the formalin test. On the other hand, pretreatment with 5 or 7 mg kg⁻¹ dose did not alter the pain behaviors of the first phase but significantly decreased the nociceptive behavior during the second phase. This reduction was evidenced by decreased flinches and the lifting/licking time compared to vehicle-treated mice, suggesting a σ 1R antagonist profile for compound 7. Compound 7 showed the same pharmacological profile as (+)-LP2, which evidenced an antinociceptive effect in the second phase of the formalin test.²⁴

3. Conclusions

In summary, we investigated the influence of the *N*-substituent on the affinity profile and selectivity profile of (+)-(2*S*,6*S*,11*S*)-normetazocine-based compounds, known for their capacity to bind σ 1R.³³ Among the newly synthesized ligands, compound 7 exhibited high σ 1R affinity, while compound 10 displayed moderate affinity. Both compounds showed negligible affinity for σ 2R and opioid receptors. These findings underscore the crucial role of normetazocine

stereochemistry in achieving σ 1R affinity. Regarding the *N*-substituent structure, compounds 7 and 10 possess an ester functional group positioned at an optimal distance from the nitrogen atom of the normetazocine scaffold despite the absence of a phenyl ring in compound 10. The higher σ 1R affinity of compound 7 compared to compound 10 aligns with the σ 1R pharmacophore model proposed by Glennon *et al.*,³⁴ which features two hydrophobic regions at an appropriate distance from the nitrogen atom.³⁵

Moreover, compound 7 produced a potent antinociceptive effect in a mouse model of inflammatory pain, suggesting that it acts as a σ 1R antagonist. These *in vivo* results, consistent with literature reports,³⁶ support the potential of compound 7 as a therapeutic agent for managing inflammatory pain through σ 1R antagonism. Notably, the i.p. administration of compound 7 reduces formalin-induced nociception associated with the second phase of the formalin test without affecting the first phase. The injection of formalin induces a biphasic pain response reflecting two different forms of pain. Phase I is a rapid pain response occurring immediately after formalin injection, lasting about 10 minutes, representing acute phasic pain due to direct activation of nociceptors. Phase II is a late pain response starting after 10–15 minutes and lasting up to 50–60 minutes, representing more persistent, tonic, inflammatory pain. This second phase is clinically significant as it reflects mechanisms of nociceptive sensitization, particularly in the dorsal horn of the spinal cord,³⁷ where σ 1Rs play a crucial role in pain transmission and modulation.³⁸ Given its ability to reduce pain during the second phase, compound 7 could represent a promising drug for treating persistent pain conditions, such as neuropathic pain, characterized by central sensitization. Previously reported σ 1R compounds with a similar profile in formalin-induced inflammatory pain have shown efficacy in counteracting neuropathic pain.³⁹ In particular, (+)-LP2, which exhibits a comparable profile to compound 7 during the second phase of the formalin test, demonstrated inhibition of mechanical allodynia and modulation of neuroinflammation in models of chronic neuropathic pain.²⁵

In silico ADME studies demonstrate compound 7 has a favorable balance of pharmacokinetic and pharmacodynamic properties, meeting critical criteria for oral bioavailability, solubility, absorption, and CNS penetration. While its interaction with CYP enzymes requires consideration, the compound holds significant potential for further development, particularly in CNS-targeted therapies. The absence of PAINS liabilities and compliance with major drug-likeness rules enhance its prospects as a lead candidate. Stability of compound 7 was also evaluated. While compound 7's chemical stability *in vitro* at 37 °C in aqueous phosphate buffer (PBS) at pH 7.4 was established and a half-life (*t*_{1/2}) of 300 min was measured, the plasma stability assay on compound 7 showed its progressive hydrolysis with a half-life (*t*_{1/2}) of 31.07 min. However, its high BBB permeability score measured and the *in vivo* evidence, as compound 7

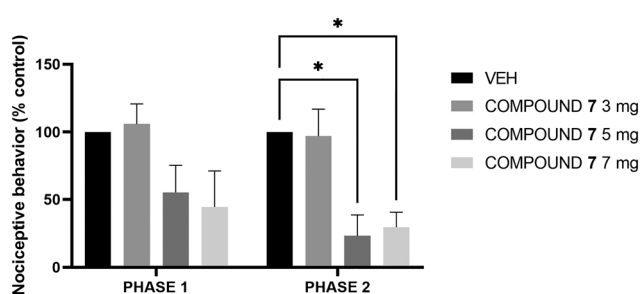


Fig. 4 Effects of compound 7 at the doses of 3.0, 5.0, and 7.0 mg kg⁻¹ i.p. in phase I (0–10) and phase II (10–60) of formalin test. Results are reported as MEAN \pm SEM (*n* = 4–11 per group) of the percentage of inhibition of pain responses of the respective vehicle group. **p* < 0.05, versus vehicle, two-way ANOVA followed by Bonferroni's *post hoc* test.



demonstrated pain modulation in the inflammatory pain model, support the pharmacological profile of compound 7. Therefore, future studies are required to improve stability of compound 7 with the aim to obtain its analogues useful in treating neuropathic pain.

4. Experiment section

4.1. Chemistry

4.1.1. General remarks. Reagent-grade chemicals were purchased from Merck (Darmstadt, Germany) and were used without further purification. (+)-*cis*-Normetazocine was obtained from Fabbria Italiana Sintetici. The methyl-3-bromopropanoate (4) and methyl 4-chlorobutanoate (5) were purchased from Merck (Darmstadt, Germany). Flash column chromatography was carried out on Merck silica gel 60 (230–400 mesh). Reactions were monitored by thin-layer chromatography (TLC) performed on 250 μ m silica gel Merck 60 F254 coated aluminum plates; the spots were visualized by UV light or iodine chamber. Melting points were determined in open capillary tubes with a Büchi 530 apparatus and are uncorrected. ^1H , ^{13}C spectra were recorded at 200 and 500 MHz on Varian Inova spectrometers in CDCl_3 or $\text{DMSO}-d_6$. ^{13}C spectra were ^1H -decoupled, and the attached proton test (APT) pulse sequence determined the multiplicities. Chemical shifts δ are expressed in parts per million (ppm). The following abbreviations are used to designate the multiplicities: s = singlet, d = doublet, t = triplet, m = multiplet, br = broad. Elemental analyses (C, H, N) were performed on a Carlo Erba 1106 analyzer (Milan, Italy), and the analysis results were within 0.4% of the theoretical values.

The intermediates 1–3 were obtained as previously reported.²⁶

4.1.2. Ethyl 3-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-2-(*R*/*S*)-phenylpropanoate (6). (+)-*cis*-Normetazocine (1.17 mmol, 1 eq.) was dissolved in DMF (5 mL) and ethyl 3-[(methylsulfonyl)oxy]-2-(*R*/*S*)-phenylpropanoate (1) (1.17 mmol, 1 eq.), NaHCO_3 (1.76 mmol, 1.5 eq.) and KI (catalytic quantity) were added. The mixture was stirred overnight at 55 °C. At the reaction mixture, H_2O (4 mL) was added, and the aqueous phase was extracted with EtOAc; the organic phase was dried over Na_2SO_4 and concentrated *in vacuo*. The crude was purified by flash chromatography ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 95:5) to give a white solid. Yield: 88%; m.p.: 183–186 °C. TLC $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (95:5 v/v) R_f = 0.49. ^1H NMR (500 MHz, CDCl_3): δ = 7.83–7.79 (m, 1H, *CH*-aryl), 7.32–7.26 (m, 4H, *CH*-aryl), 6.86 (d, 1H, J = 8.3 Hz, *CH*-aryl benzomorphan), 6.70 (d, 1H, J = 3.4 Hz, *CH*-aryl benzomorphan), 6.58 (dd, 1H, J = 8.3 and 3.4 Hz, *CH*-aryl benzomorphan), 3.82–3.79 (m, 1H, *CH*-CO), 3.68 (s, 3H, OCH_2CH_3), 2.99–2.97 (m, 2H, CH_2 -benzomorphan), 2.91–2.88 (m, 2H, CH_2), 2.89 (d, 2H, J = 15.0 Hz, CH_2 - CH_3), 2.56–2.67 (m, 4H, CH_2 -benzomorphan), 1.79–1.70 (m, 2H, CH_2 -benzomorphan), 1.32 (s, 3H, CH_3 -benzomorphan), 0.79 (d, 3H, J = 7.0, CH_3 -benzomorphan). The –OH signal of

benzomorphan is not observed (Fig. S7†). APT NMR (126 MHz, CDCl_3) δ = 174.11 (C=O), 154.62 (*C*-aryl-OH benzomorphan), 137.83 (*C*-aryl benzomorphan), 137.57 (*C*-aryl), 128.74 (*CH*-aryl), 128.25 (*CH*-aryl benzomorphan), 128.32 (*CH*-aryl), 128.16 (*CH*-aryl), 128.10 (*C*-aryl benzomorphan), 112.99 (*CH*-aryl benzomorphan), 112.39 (*CH*-aryl benzomorphan), 59.95 ($\text{O}-\text{CH}_2\text{CH}_3$), 58.28 (*CH* piperidine benzomorphan), 57.34 ($\text{N}-\text{CH}_2$), 46.40 (CH_2 piperidine benzomorphan), 42.36 (CH_2 piperidine benzomorphan), 41.72 (*CH* piperidine benzomorphan), 41.62 (CH_2 -*CH*-aryl/ $\text{COOCH}_2\text{CH}_3$), 36.56 (*C*-piperidine benzomorphan), 25.71 (CH_3 benzomorphan), 24.95 (CH_2 aliphatic benzomorphan), 14.33 ($\text{O}-\text{CH}_2\text{CH}_3$), 13.71 (CH_3 benzomorphan) (Fig. S17†). Anal. ($\text{C}_{25}\text{H}_{31}\text{NO}_3$) C, H, N (Table S1†).

4.1.3. Ethyl 4-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-2-(*R*/*S*)-phenylbutanoate (7). (+)-*cis*-Normetazocine (3.09 mmol, 1 eq.) was dissolved in DMF (6 mL), and ethyl 4-chloro-2-(*R*/*S*)-phenylbutanoate (2) (3.09 mmol, 1 eq.), NaHCO_3 (5.85 mmol, 1.5 eq.) and KI (catalytic quantity) were added. The mixture was stirred at 65 °C for 72 h. After, the mixture was transferred to a separatory funnel and partitioned (EtOAc/ H_2O). The organic phase was washed with brine, dried on Na_2SO_4 , and concentrated under vacuum. The reaction crude was purified by flash chromatography ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 95:5) to give an orange solid. Yield: 25%; m.p.: 158–161 °C; TLC $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (95:5 v/v) R_f = 0.36. Chiral HPLC: 55.6%, t_R = 12.3 min; 44.4%, and t_R = 14.6 min (Fig. S1†). ^1H NMR (200 MHz, CDCl_3): δ = 7.40–7.32 (m, 5H, *CH*-aryl), 6.87 (d, 1H, J = 8.0 Hz, *CH*-aryl benzomorphan), 6.95 (d, 1H, J = 2.2 Hz, *CH*-aryl benzomorphan), 6.71 (dd, 1H, J = 8.0 and 2.2 Hz, *CH*-aryl benzomorphan), 4.21 (q, 2H, J = 7.2 Hz, $\text{O}-\text{CH}_2-\text{CH}_3$), 3.30–3.08 (m, 1H, *CH*-CO), 3.01–2.93 (m, 7H, *CH*-benzomorphan, CH_2 -benzomorphan, CH_2 , CH_2), 2.66–2.46 (m, 4H, CH_2 -benzomorphan), 1.46–1.44 (m, 1H, *CH*-benzomorphan), 1.20 (s, 3H, CH_3 -benzomorphan), 1.12 (t, 3H, J = 7.2 Hz, $\text{O}-\text{CH}_2-\text{CH}_3$), 0.89 (d, 3H, J = 7.0 Hz, CH_3 -benzomorphan). The –OH signal of benzomorphan was not observed (Fig. S8†). ^{13}C NMR (126 MHz, CDCl_3) δ = 167.36 (C=O), 155.66 (*C*-aryl-OH benzomorphan), 133.45 (*C*-aryl benzomorphan), 133.06 (*C*-aryl), 129.25 (*CH*-aryl), 129.20 (*CH*-aryl benzomorphan), 129.11 (*CH*-aryl), 129.08 (*C*-aryl benzomorphan), 128.28 (*CH*-aryl), 114.19 (*CH*-aryl benzomorphan), 112.43 (*CH*-aryl benzomorphan), 54.04 ($\text{O}-\text{CH}_2\text{CH}_3$), 54.01 (*CH* piperidine benzomorphan), 54.00 ($\text{N}-\text{CH}_2-\text{CH}_2$), 53.37 (CH_2 piperidine benzomorphan), 53.22 (CH_2 piperidine benzomorphan), 35.24 (*CH* piperidine benzomorphan), 34.84 (CH_2 -*CH*-(aryl/ $\text{COOCH}_2\text{CH}_3$), 34.85 (*C*-piperidine benzomorphan), 34.55 ($\text{N}-\text{CH}_2-\text{CH}_2$), 29.48 (CH_3 benzomorphan), 23.84 (CH_2 aliphatic benzomorphan), 13.53 ($\text{O}-\text{CH}_2\text{CH}_3$), 13.50 (CH_3 benzomorphan) (Fig. S18†). Anal. ($\text{C}_{26}\text{H}_{33}\text{NO}_3$) C, H, N (Table S1†).

4.1.4. Ethyl 5-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-2-(*R*/*S*)-phenylpentanoate (8). (+)-*cis*-Normetazocine (3.50 mmol, 1



eq.) was dissolved in DMF (6 mL), and ethyl 5-chloro-2-(*R/S*)-phenylpentanoate (**3**) (4.15 mmol, 1 eq.), NaHCO₃ (5.25 mmol, 1.5 eq.) and KI (catalytic quantity) were added. The reaction mixture was stirred at 65 °C for 72 h. After, the mixture was transferred to a separatory funnel and partitioned (EtOAc/H₂O). The organic phase was washed with brine, dried on Na₂SO₄, and concentrated in vacuum. The crude was purified by flash chromatography (CH₂Cl₂/MeOH 95:5) to give an orange solid. Yield: 40%; m.p.: 160–163 °C; TLC CH₂Cl₂/MeOH (95:5 v/v) *R*_f = 0.47. ¹H NMR (500 MHz, CDCl₃): δ = 7.48–7.31 (m, 4H, *CH*-aryl), 7.28–7.27 (m, 1H, *CH*-aryl), 7.01 (d, 1H, *J* = 10.0 Hz, *CH*-benzomorphan), 6.76 (d, 1H, *J* = 10 Hz, *CH*-aryl benzomorphan), 6.69 (d, 1H, *J* = 10.0 Hz, *CH*-aryl benzomorphan), 3.47–3.24 (m, 2H, CH₂-CH₃), 3.17–3.06 (m, 3H, CH₂-benzomorphan, *CH*-CO), 2.14–1.97 (m, 6H, CH₂, CH₂, CH₂), 1.66–1.65 (m, 3H, CH, CH₂-benzomorphan), 1.55–1.40 (m, 2H, CH₂-benzomorphan), 1.30 (s, 3H, CH₃-benzomorphan), 1.27 (s, 3H, CH₂-CH₃), 1.21–1.24 (m, 1H, *CH*-benzomorphan), 0.93 (d, 3H, *J* = 7.2 Hz, CH₃-benzomorphan). The –OH signal of benzomorphan is not observed (Fig. S9†). ¹³C NMR (126 MHz, CDCl₃) δ = 167.34 (C=O), 155.65 (*C*-aryl-OH benzomorphan), 133.44 (*C*-aryl benzomorphan), 133.05 (*C*-aryl), 129.23 (*CH*-aryl), 129.19 (*CH*-aryl benzomorphan), 129.10 (*CH*-aryl), 129.07 (*C*-aryl benzomorphan), 128.27 (*CH*-aryl), 114.17 (*CH*-aryl benzomorphan), 112.42 (*CH*-aryl benzomorphan), 63.30 (O-CH₂CH₃), 54.03 (*CH* piperidine benzomorphan), 54.02 (N-CH₂-CH₂-CH₂), 54.00 (CH₂ piperidine benzomorphan), 53.36 (CH₂ piperidine benzomorphan), 53.35 (*CH* piperidine benzomorphan), 53.21 (N-CH₂-CH₂-CH₂), 35.23 (CH₂-CH-aryl/COOCH₂CH₃), 34.84 (*C*-piperidine benzomorphan), 34.53 (N-CH₂-CH₂-CH₂), 29.47 (CH₃ benzomorphan), 24.01 (CH₂ aliphatic benzomorphan), 13.65 (O-CH₂CH₃), 13.49 (CH₃ benzomorphan) (Fig. S19†). Anal. (C₂₇H₃₅NO₃) C, H, N (Table S1†).

4.1.5. Methyl 3-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[d]azocin-3(2*H*)]-propanoate (9**).** (+)-*cis*-Normetazocine (1.84 mmol, 1 eq.) was dissolved in DMF (5 mL), and methyl-3-bromopropanoate (**4**) (1.84 mmol, 1 eq.), NaHCO₃ (2.76 mmol, 1.5 eq.) and KI (catalytic quantity) were added. The reaction mixture was stirred overnight at 55 °C. At the reaction mixture, H₂O (50 mL) was added, and the aqueous phase was extracted with EtOAc; the organic phase was dried over Na₂SO₄ and concentrated *in vacuo*. The crude was purified by flash chromatography (CH₂Cl₂/MeOH 93:7) to give a white solid. Yield: 78%; m.p.: 139–142 °C. TLC CH₂Cl₂/MeOH (93:7 v/v) *R*_f = 0.49. ¹H NMR (200 MHz, CDCl₃): δ = 6.83 (d, 1H, *J* = 8.0 Hz, *CH*-aryl benzomorphan), 6.63 (m, 1H, *CH*-aryl benzomorphan), 6.55 (d, 1H, *J* = 8.0 Hz, *CH*-aryl benzomorphan), 3.58 (s, 3H, O-CH₃), 2.98–2.91 (m, 4H, CH₂-benzomorphan, CH₂-CH₂-CO), 2.82–2.57 (m, 4H, CH₂-benzomorphan, CH₂-CH₂-CO), 2.21–2.09 (m, 1H, *CH*-benzomorphan), 1.90–1.76 (m, 2H, CH₂-benzomorphan), 1.29–1.44 (m, 1H, *CH*-benzomorphan), 1.21 (s, 3H, CH₃-benzomorphan), 0.75 (d, 3H, 8.0 Hz, CH₃-benzomorphan). The –OH signal of benzomorphan was not

observed (Fig. S10†). APT NMR (126 MHz, DMSO-*d*₆) δ = 174.63 (C=O), 155.67 (*C*-aryl-OH benzomorphan), 141.26 (*C*-aryl benzomorphan), 127.62 (*CH*-aryl benzomorphan), 124.96 (*C*-aryl benzomorphan), 113.21 (*CH*-aryl benzomorphan), 111.72 (*CH*-aryl benzomorphan), 57.11 (O-CH₃), 54.06 (N-CH₂-CH₂), 44.26 (CH₂ piperidine benzomorphan), 40.81 (CH₂ piperidine benzomorphan), 40.80 (*CH* piperidine benzomorphan), 40.79 (*CH* piperidine benzomorphan), 35.88 (N-CH₂-CH₂), 34.54 (*C*-aliphatic benzomorphan), 25.30 (CH₃ benzomorphan), 23.43 (CH₂ aliphatic benzomorphan), 13.94 (CH₃ benzomorphan) (Fig. S20†). Anal. (C₁₈H₂₅NO₃) C, H, N (Table S1†).

4.1.6. Methyl 4-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[d]azocin-3(2*H*)]-butanoate (10**).** (+)-*cis*-Normetazocine (2.30 mmol, 1 eq.) was dissolved in DMF (5 mL), and methyl 4-chlorobutanoate (**5**) (2.30 mmol, 1 eq.), NaHCO₃ (3.43 mmol, 1.5 eq.) and KI (catalytic quantity) were added. The reaction mixture was stirred overnight at 55 °C. At the reaction mixture, H₂O (50 mL) was added, and the aqueous phase was extracted with EtOAc; the organic phase was dried over Na₂SO₄ and concentrated *in vacuo*. The crude was purified by flash chromatography (CH₂Cl₂/MeOH 93:7) to give a white solid. Yield: 75%; m.p.: 143–146 °C. TLC CH₂Cl₂/MeOH (93:7 v/v) *R*_f = 0.50. ¹H NMR (200 MHz, CDCl₃): δ = 6.61 (d, 1H, *J* = 10.0 Hz, *CH*-aryl benzomorphan), 6.38 (m, 1H, *CH*-aryl benzomorphan), 6.34 (d, 1H, *J* = 10.0 Hz, *CH*-aryl benzomorphan), 3.38 (s, 3H, O-CH₃), 2.66–2.57 (m, 4H, CH₂-benzomorphan, CH₂-CH₂-CH₂-CO), 2.04–1.96 (m, 1H, *CH*-benzomorphan), 1.96–1.90 (m, 2H, CH₂-benzomorphan), 1.90–1.85 (m, 2H, CH₂-CH₂-CH₂-CO), 1.85–1.62 (m, 4H, CH₂-benzomorphan, CH₂-CH₂-CH₂-CO), 1.02–0.95 (m, 1H, *CH*-benzomorphan), 0.91 (s, 3H, CH₃-benzomorphan), 0.54 (d, 3H, 10.0 Hz, CH₃-benzomorphan). The –OH signal of benzomorphan was not observed (Fig. S11†). APT NMR (126 MHz, DMSO-*d*₆) δ = 174.65 (C=O), 155.68 (*C*-aryl-OH benzomorphan), 141.27 (*C*-aryl benzomorphan), 127.64 (*CH*-aryl benzomorphan), 124.97 (*C*-aryl benzomorphan), 113.22 (*CH*-aryl benzomorphan), 111.73 (*CH*-aryl benzomorphan), 57.12 (O-CH₃), 54.07 (N-CH₂-CH₂-CH₂), 44.27 (CH₂ piperidine benzomorphan), 40.83 (CH₂ piperidine benzomorphan), 40.80 (*CH* piperidine benzomorphan), 40.79 (*CH* piperidine benzomorphan), 38.02 (N-CH₂-CH₂-CH₂), 35.90 (N-CH₂-CH₂-CH₂), 34.40 (*C*-aliphatic benzomorphan), 25.31 (CH₃ benzomorphan), 23.44 (CH₂ aliphatic benzomorphan), 13.95 (CH₃ benzomorphan) (Fig. S21†). Anal. (C₁₉H₂₇NO₃) C, H, N (Table S1†).

4.1.7. 3-[(2*S*,6*S*,11*S*)-8-Hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[d]azocin-3(2*H*)]-2-(*R/S*)-phenylpropanoic acid (11**).** 1 N NaOH solution (3.06 mmol, 9.27 eq.) was added to ethyl 3-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[d]azocin-3(2*H*)]-2-(*R/S*)-phenylpropanoate (**6**) (0.82 mmol, 1 eq.). The suspension was strongly stirred and refluxed at 110 °C for 5 h. After cooling, the mixture was partitioned (CHCl₃/H₂O). A 1 N solution of HCl was added to the aqueous phase to give a



pH of 5–6. The resulting yellow precipitate was crystallized by EtOH/diethyl ether. Yield: 80%; m.p.: 181–184 °C; TLC CH₂Cl₂/MeOH (95:5 v/v); *R_f* = 0.32. ¹H NMR (500 MHz, DMSO-*d*₆): δ = 9.08 (br, 1H, OH), 7.28–7.24 (m, 5H, CH-aryl benzomorphan), 6.83–6.82 (m, 1H, CH-aryl benzomorphan), 6.81–6.80 (m, 1H, CH-aryl benzomorphan), 6.56–6.50 (m, 1H, CH-aryl benzomorphan), 4.07–4.02 (m, 1H, CH-CO), 3.12–3.08 (m, 4H, CH₂-benzomorphan, CH₂), 2.80–2.75 (m, 3H, CH, CH₂-benzomorphan), 2.41–2.42 (m, 1H, CH₂-benzomorphan), 1.99–1.76 (m, 2H, CH-benzomorphan), 1.19 (s, 3H, CH₃-benzomorphan), 0.68 (d, 3H, CH₃-benzomorphan). The –OH signal of benzomorphan was not observed (Fig. S12†). APT NMR (126 MHz, DMSO-*d*₆) δ = 174.44 (C=O), 156.95 (C-aryl-OH benzomorphan), 141.65 (C-aryl benzomorphan), 138.81 (C-aryl), 129.54 (CH-aryl), 129.48 (CH-aryl benzomorphan), 129.16 (CH-aryl), 129.02 (CH-aryl), 128.94 (C-aryl benzomorphan), 114.50 (CH-aryl benzomorphan), 112.83 (CH-aryl benzomorphan), 60.52 (CH piperidine benzomorphan), 56.26 (N-CH₂), 52.00 (CH piperidine benzomorphan), 47.74 (CH₂ piperidine benzomorphan), 46.08 (CH₂-CH-(aryl/COOH), 41.00 (CH₂ piperidine benzomorphan), 36.17 (C-piperidine benzomorphan), 27.58 (CH₃ benzomorphan), 26.39 (CH₂ aliphatic benzomorphan), 13.93 (CH₃ benzomorphan) (Fig. S22†). Anal. (C₂₃H₂₇NO₃) C, H, N (Table S1†).

4.1.8. 4-[(2*S*,6*S*,11*S*)-8-Hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-2-(*R*/*S*)-phenylbutanoic acid (12). 1 N NaOH solution (3.06 mmol, 11.3 eq.) was added to ethyl 4-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-2-(*R*/*S*)-phenylbutanoate (7) (0.27 mmol, 1 eq.). The obtained suspension was vigorously stirred and refluxed at 110 °C for 5 h. After, the mixture was cooled at rt, transferred to a separatory funnel, and partitioned (CHCl₃/H₂O). A 1 N solution of HCl was added to the aqueous phase to pH 5–6. A yellow precipitate was obtained that was separated from the aqueous phase by vacuum filtration. Yield: 53%, m.p.: 190 °C; TLC CH₂Cl₂/MeOH (95:5 v/v) *R_f* = 0.30. ¹H NMR (500 MHz, DMSO-*d*₆): δ = 8.43 (br, 1H, OH), 7.22–7.20 (m, 2H, CH-aryl), 7.13–7.10 (m, 2H, CH-aryl), 7.03–7.01 (m, 1H, CH-aryl), 6.74 (d, 1H, *J* = 10.0 Hz, CH-aryl benzomorphan), 6.52 (d, 1H, *J* = 10.0 Hz, CH-aryl benzomorphan), 6.43–6.41 (dd, 1H, *J* = 10.0 and 3.4 Hz, CH-aryl benzomorphan), 3.13–3.10 (m, 2H, CH-CO, CH-benzomorphan), 2.68–2.62 (m, 2H, CH₂-benzomorphan), 2.38–2.21 (m, 4H, CH₂, CH₂), 1.80–1.77 (m, 2H, CH₂-benzomorphan), 1.80–1.77 (m, 2H, CH₂-benzomorphan), 1.64–1.60 (m, 1H, CH-benzomorphan), 1.16 (s, 3H, CH₃-benzomorphan), 0.66 (d, 3H, *J* = 7.2 Hz, CH₃-benzomorphan). The –OH signal of benzomorphan was not observed (Fig. S13†). APT NMR (126 MHz, DMSO-*d*₆) δ = 176.93 (C=O), 155.91 (C-aryl-OH benzomorphan), 144.63 (C-aryl benzomorphan), 142.17 (C-aryl), 127.89 (CH-aryl), 127.64 (CH-aryl benzomorphan), 127.51 (CH-aryl), 126.59 (C-aryl benzomorphan), 125.43 (CH-aryl), 113.12 (CH-aryl benzomorphan), 111.92 (CH-aryl benzomorphan), 56.43 (CH piperidine benzomorphan), 52.75 (N-CH₂-CH₂), 51.75 (CH

piperidine benzomorphan), 44.80 (CH₂ piperidine benzomorphan), 41.78 (CH₂ piperidine benzomorphan), 36.21 (CH₂-CH-aryl/COOH), 32.91 (C-piperidine benzomorphan), 28.89 (N-CH₂-CH₂), 25.28 (CH₃ benzomorphan), 22.43 (CH₂ aliphatic benzomorphan), 13.44 (CH₃ benzomorphan) (Fig. S23†). Anal. (C₂₄H₂₉NO₃) C, H, N (Table S1†).

4.1.9. 5-[(2*S*,6*S*,11*S*)-8-Hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-2-(*R*/*S*)-phenylpentanoic acid (13). 1 N NaOH solution (3.06 mmol, 11.3 eq.) was added to ethyl 5-[(2*S*,6*S*,11*S*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-2-(*R*/*S*)-phenylpentanoate (8) (0.95 mmol, 1 eq.). The obtained suspension was vigorously stirred and refluxed at 110 °C for 5 h. After, the mixture was cooled at r.t., transferred to a separatory funnel, and partitioned (CHCl₃/H₂O). A 1 N solution of HCl was added to the aqueous phase to pH 5–6. A yellow precipitate was obtained that was separated from the aqueous phase by vacuum filtration. Yield: 53%, m.p.: 196 °C; TLC CH₂Cl₂/MeOH (95:5 v/v) *R_f* = 0.28. ¹H NMR (500 MHz, DMSO-*d*₆): δ = 7.48–7.31 (m, 5H, CH-aryl), 7.28–7.27 (m, 1H, CH-aryl), 7.01 (d, 1H, *J* = 10.0 Hz, CH-aryl benzomorphan), 6.76 (d, 1H, *J* = 10 Hz, CH-aryl benzomorphan), 6.69 (dd, 1H, *J* = 10.0 and 3.4 Hz, CH-aryl benzomorphan), 3.47–3.24 (m, 2H, CH-CO, CH-benzomorphan), 3.17–3.06 (m, 2H, CH₂-benzomorphan), 2.14–1.97 (m, 4H, CH₂, CH₂), 1.66–1.65 (m, 4H, CH₂-benzomorphan), 1.23 (s, 3H, CH₃-benzomorphan), 1.21–1.22 (m, 1H, CH-benzomorphan), 0.93 (d, 3H, *J* = 7.2 Hz, CH₃-benzomorphan). The –OH signals of benzomorphan and carboxylic acid moiety were not observed (Fig. S14†). APT NMR (126 MHz, DMSO-*d*₆) δ = 174.50 (C=O), 156.10 (C-aryl-OH benzomorphan), 143.39 (C-aryl benzomorphan), 139.79 (C-aryl), 128.68 (CH-aryl), 128.39 (CH-aryl benzomorphan), 128.05 (CH-aryl), 127.88 (CH-aryl), 127.14 (C-aryl benzomorphan), 113.76 (CH-aryl benzomorphan), 112.50 (CH-aryl benzomorphan), 57.76 (CH piperidine benzomorphan), 51.24 (CH piperidine benzomorphan), 50.79 (N-CH₂-CH₂-CH₂), 39.62 (CH₂ piperidine benzomorphan), 35.10 (CH₂ piperidine benzomorphan), 33.57 (C-piperidine benzomorphan), 31.22 (N-CH₂-CH₂-CH₂), 29.09 (CH₂-CH-(aryl/COOH), 25.25 (N-CH₂-CH₂-CH₂), 24.86 (CH₂ aliphatic benzomorphan), 19.73 (CH₃ benzomorphan), 13.74 (CH₃ benzomorphan) (Fig. S24†). Anal. (C₂₅H₃₁NO₃) C, H, N (Table S1†).

4.1.10. 3-[(2*S*,6*S*,11*S*)-8-Hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-propanoic acid (14). 1 N NaOH solution (7.70 mmol, 11.3 eq.) was added to methyl 3-[(2*S*,6*S*,11*R*)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[*d*]azocin-3(2*H*)-yl]-propanoate (9) (0.68 mmol, 1 eq.). The obtained suspension was vigorously stirred and refluxed at 110 °C for 5 h. After, the mixture was cooled at r.t., transferred to a separatory funnel, and partitioned (CHCl₃/H₂O). A 1 N solution of HCl was added to the aqueous phase to pH 5–6. A white precipitate it was obtained that was separated from the aqueous phase by



vacuum filtration. Yield: 63%, m.p.: 198 °C; TLC $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (95 : 5 v/v) R_f = 0.28. ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ = 6.82 (d, 1H, J = 10.0 Hz, CH-aryl benzomorphan), 6.60 (d, 1H, J = 5.0 Hz, CH-aryl benzomorphan), 6.50 (dd, 1H, J = 10.0 and 5.0 Hz, CH-aryl benzomorphan), 2.76–2.81 (m, 2H, CH_2 -benzomorphan), 2.65–2.60 (m, 4H, CH_2 -benzomorphan, $\text{CH}_2\text{-CH}_2\text{-CO}$), 2.42–2.39 (m, 2H, m, CH_2 -benzomorphan), 2.07–2.05 (m, 2H, $\text{CH}_2\text{-CH}_2\text{-CO}$), 1.92–1.70 (m, 1H, CH -benzomorphan), 1.22 (s, 3H, CH_3 -benzomorphan), 1.18–1.15 (m, 1H, CH -benzomorphan), 0.72 (d, 3H, J = 10.0 Hz, CH_3 -benzomorphan). The $-\text{OH}$ signals of benzomorphan and carboxylic acid moiety were not observed (Fig. S15[†]). APT NMR (126 MHz, $\text{DMSO}-d_6$) δ = 171.47 (C=O), 156.01 (C-aryl-OH benzomorphan), 139.84 (C-aryl benzomorphan), 128.20 (CH-aryl benzomorphan), 122.98 (C-aryl benzomorphan), 113.74 (CH-aryl benzomorphan), 111.62 (CH-aryl benzomorphan), 57.90 ($\text{N-CH}_2\text{-CH}_2$), 48.70 (CH_2 piperidine benzomorphan), 44.98 (CH_2 piperidine benzomorphan), 40.80 (CH piperidine benzomorphan), 35.72 (CH_2 aliphatic benzomorphan), 34.58 ($\text{N-CH}_2\text{-CH}_2$), 33.46 (CH piperidine benzomorphan), 29.54 (C-aliphatic benzomorphan), 24.08 (CH_3 benzomorphan), 12.84 (CH_3 benzomorphan) (Fig. S25[†]). Anal. ($\text{C}_{17}\text{H}_{23}\text{NO}_3$) C, H, N (Table S1[†]).

4.1.11. 4-[(2S,6S,11S)-8-Hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[d]azocin-3(2H)yl]-butanoic acid (15). 1 N NaOH solution (4.40 mmol, 11.3 eq.) was added to methyl 4-[(2S,6S,11S)-8-hydroxy-6,11-dimethyl-1,4,5,6-tetrahydro-2,6-methanobenzo[d]azocin-3(2H)yl]-butanoate (**10**) (0.39 mmol, 1 eq.). The obtained suspension was vigorously stirred and refluxed at 110 °C for 5 h. After, the mixture was cooled at r.t., transferred to a separatory funnel, and partitioned ($\text{CHCl}_3/\text{H}_2\text{O}$). A 1 N solution of HCl was added to the aqueous phase to pH 5–6. A white precipitate was obtained that was separated from the aqueous phase by vacuum filtration. Yield: 60%, m.p.: 197 °C; TLC $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (95 : 5 v/v) R_f = 0.30. ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ = 7.09 (d, 1H, J = 4.0 Hz, CH-aryl benzomorphan), 6.85 (d, 1H, J = 2.0 Hz, CH-aryl benzomorphan), 6.77 (dd, 1H, J = 4.0 and 2.0 Hz, CH-aryl benzomorphan), 3.38–3.34 (m, 1H, CH -benzomorphan), 2.95–2.88 (m, 4H, CH_2 -benzomorphan, $\text{CH}_2\text{-CH}_2\text{-CO}$), 2.36–2.33 (m, 2H, CH_2 -benzomorphan), 2.16–2.15 (m, 2H, $\text{CH}_2\text{-CH}_2\text{-CH}_2\text{-CO}$), 1.97–1.92 (m, 4H, CH_2 -benzomorphan, $\text{CH}_2\text{-CH}_2\text{-CH}_2\text{-CO}$), 1.59–1.50 (m, 1H, CH -benzomorphan), 1.46 (s, 3H, CH_3 -benzomorphan), 0.96 (d, 3H, J = 4.0 Hz, CH_3 -benzomorphan). The $-\text{OH}$ signals of benzomorphan and carboxylic acid moiety were not observed (Fig. S16[†]). APT NMR (126 MHz, $\text{DMSO}-d_6$) δ = 173.44 (C=O), 154.48 (C-aryl-OH benzomorphan), 140.07 (C-aryl benzomorphan), 126.43 (CH-aryl benzomorphan), 123.77 (C-aryl benzomorphan), 112.02 (CH-aryl benzomorphan), 110.52 (CH-aryl benzomorphan), 52.87 ($\text{N-CH}_2\text{-CH}_2\text{-CH}_2$), 43.07 (CH_2 piperidine benzomorphan), 39.62 (CH_2 piperidine benzomorphan), 39.50 (CH piperidine benzomorphan), 39.48 (CH piperidine benzomorphan), 34.89 ($\text{N-CH}_2\text{-CH}_2\text{-CH}_2$), 33.35 (C-aliphatic benzomorphan), 24.11 (CH_3 benzomorphan), 22.24 ($\text{N-CH}_2\text{-CH}_2\text{-CH}_2$), 20.27 (CH_2 aliphatic benzomorphan), 12.75 (CH_3

benzomorphan) (Fig. S26[†]). Anal. ($\text{C}_{18}\text{H}_{25}\text{NO}_3$) C, H, N (Table S1[†]).

4.2. Radioligand binding assays

4.2.1. Radioligand binding assays for $\sigma 1\text{R}$ and $\sigma 2\text{R}$. The radioligand binding assays and the data analysis were performed as previously reported.⁴⁰

4.2.1.1. Materials. [^3H](+)-Pentazocine (26.9 Ci mmol^{-1}) and [^3H]1,3-di-*o*-tolylguanidine ([^3H]DTG, 35.5 Ci mmol^{-1}) were purchased from PerkinElmer (Zaventem, Belgium). Ultima Gold MV Scintillation cocktail was from PerkinElmer (Milan, Italy). All the other materials were obtained from Merck Life Science S.r.l. (Milan, Italy). The test compound solutions were prepared by dissolving approximately 10 μmol of the test compound in DMSO to obtain a 10^{-2} M stock solution. The required test concentrations for the assay (from 10^{-5} to 10^{-10} M) have been prepared by diluting the DMSO stock solution with the respective assay buffer. All experiments were performed using ultrapure water obtained with a Millipore Milli-Q Reference Ultrapure Water Purification System. All the laboratory glassware was washed with 6 M HCl water solution and then rinsed with ultrapure water.

4.2.1.2. Preparation of membrane homogenates from rat liver. Rat livers (~21 g, male Sprague Dawley rats – ENVIGO RMS S.R.L., Udine, Italy) were cut into small pieces with a scalpel and homogenized in two portions with 6 volumes of cold 0.32 M sucrose with a Potter–Elvehjem glass homogenizer. The suspension was centrifuged at $1030 \times g$ for 10 min at 4 °C. The supernatant was separated and centrifuged at $31\,100 \times g$ for 20 min at 4 °C. The pellet was resuspended with 6 volumes of ice-cold Tris buffer (50 mM, pH 8) and incubated at rt for 30 min. Then, the suspension was centrifuged at $31\,100 \times g$ for 20 min at 4 °C. The final pellet was resuspended with 6 volumes of ice-cold Tris buffer and stored at -80 °C in ~1 mL portions containing about 6 mg protein mL^{-1} .^{41,42}

4.2.1.3. Protein determination. The protein concentration was determined using Bradford's method. The Bradford solution was prepared by dissolving 10 mg of Coomassie Brilliant Blue G 250 in 5 mL of 95% ethanol. To this solution, 10 mL of 85% phosphoric acid were added, and the mixture was stirred and filled to a total volume of 100 mL with ultrapure water. The calibration was carried out with bovine serum albumin as a standard at different concentrations. In a 96-well plate, 30 μL of the calibration solution or 30 μL of the membrane receptor preparation were mixed with 240 μL of the Bradford solution, respectively. After 5 min of incubation at rt, the UV absorbance was measured at λ = 595 nm using a microplate spectrophotometer reader (Synergy HT, BioTek).

4.2.1.4. $\sigma 1\text{R}$ ligand binding assays. *In vitro* $\sigma 1\text{R}$ ligand binding assays were carried out in Tris buffer (50 mM, pH 8.0) for 150 min at 37 °C. The thawed membrane preparation 8 (250 μg per sample) was incubated with increasing



concentrations of test compounds and [^3H](+)-pentazocine (2 nM) in a final volume of 0.5 mL. The K_d value of [^3H](+)-pentazocine was 2.9 nM. Unlabeled (+)-pentazocine (10 μM) was used to measure non-specific binding. Bound and free radioligand were separated by fast filtration under reduced pressure using a Millipore filter apparatus through Whatman GF/6 glass fiber filters, presoaked in a 0.5% poly(ethyleneimine) water solution for 120 min. Each filter paper was rinsed three times with 3 mL ice-cold Tris buffer (50 mM, pH 7.4), dried at r.t., and incubated overnight with 3 mL scintillation cocktail into pony vials. The bound radioactivity has been determined using a liquid scintillation counter (Beckman LS 6500).

4.2.1.5. $\sigma 2\text{R}$ ligand binding assays. *In vitro* $\sigma 2\text{R}$ ligand binding assays were carried out in Tris buffer (50 mM, pH 8.0) for 120 min at rt. The thawed membrane preparation of rat liver (250 μg per sample) (~21 g, male Sprague Dawley rats – ENVIGO RMS S.R.L., Udine, Italy) was incubated with increasing concentrations of test compounds and [^3H]DTG (2 nM) in the presence of (+)-pentazocine (5 μM) as $\sigma 1\text{R}$ masking agent in a final volume of 0.5 mL. The K_d value of [^3H]DTG was 17.9 nM. Non-specific binding was evaluated with unlabeled DTG (10 μM). Bound and free radioligand were separated by fast filtration under reduced pressure using a Millipore filter apparatus through Whatman GF/6 glass fiber filters, presoaked in a 0.5% poly(ethyleneimine) water solution for 120 min. Each filter paper was rinsed three times with 3 mL ice-cold Tris buffer (10 mM, pH 8), dried at r.t., and incubated overnight with 3 mL scintillation cocktail into pony vials. The bound radioactivity has been determined using a liquid scintillation counter (Beckman LS 6500).

4.2.1.6. Data analysis. The K_i -values were calculated using GraphPad Prism® 7.0 (GraphPad Software, San Diego, CA, USA). The K_i -values are given as mean value \pm SD from at least two independent experiments performed in duplicate.

4.2.2. Radioligand binding assays for opioid receptors. The radioligand binding assays and the data analysis were performed as previously reported.¹⁹

4.2.2.1. Materials. [^3H]-DAMGO (48.4 Ci mmol⁻¹), [^3H](2-D-Ala)-[tyrosyl-3,5-] DELTORPHIN II (54.7 Ci mmol⁻¹), and [^3H]-U69,593 (49.3 Ci mmol⁻¹) were purchased from PerkinElmer (Zaventem, Belgium). Unlabeled naloxone hydrochloride, DAMGO, (-)-U50,488, and naltrindole hydrochloride were purchased from Sigma-Aldrich (St. Louis, MO, USA). The Ultima Gold MV Scintillation cocktail was from PerkinElmer (Milano, Italy). A 10 mM stock solution was obtained by dissolving the test compound in DMSO and then diluting it with the assay buffer to obtain the required test concentrations for the assay (from 10⁻⁵ to 10⁻⁹ M). All experiments were performed using ultrapure water obtained with a Millipore Milli-Q Reference Ultrapure Water Purification System (Millipore, Burlington, MA, USA). The bound radioactivity has been determined using a Beckman LS 6500 liquid scintillation counter (Beckman Coulter, Brea, CA, USA).

4.2.2.2. Preparation of membrane homogenates from Sprague–Dawley rat brains for MOR and DOR binding assays or guinea pig brains for KOR binding assays. Sprague–Dawley rat brains (for MOR and DOR binding assays – ENVIGO RMS S. R.L., Udine, Italy) or guinea pig brains (for KOR binding assays – 200–300 g, male Dunkin–Hartley guinea pigs – Harlan Laboratories, S.Pietro al Natisone (UD)) were homogenized in ice-cold Tris buffer (50 mM, pH 7.4) by using a Dounce glass homogenizer (Wheaton, Millville, NJ, USA) with a loose inner tolerance pestle first and a tight inner tolerance pestle later in a cylindrical glass tube of 40 mL volume. The suspension was centrifuged at 40 000 $\times g$ for 20 min at 4 °C (Beckmann J2-20 centrifuge and a JA-21 rotor). The pellet was resuspended in ice-cold Tris buffer and then incubated at 37 °C for 30 min to remove endogenous ligands. After incubation, the suspension was centrifuged at 40 000 $\times g$ for 20 min at 4 °C, and the final pellet was resuspended in ice-cold Tris buffer and frozen at -80 °C in ~1 mL portions containing about 10 mg protein mL⁻¹.

4.2.2.3. Protein determination. The protein concentration was determined using the Bradford method. The calibration curve was built with bovine serum albumin as the standard compound at 7 concentrations ranging from 60 μg mL⁻¹ to 210 μg mL⁻¹ with blank correction. In a 96-well plate, 30 μL of the calibration solution or 30 μL of the membrane receptor preparation were mixed with 240 μL of the Bradford solution (10 mg of Coomassie Brilliant Blue G 250 in 5 mL of 95% ethanol, 10 mL of 85% phosphoric acid, and water up to 100 mL; with ultrapure water). After 5 min of incubation at RT, the UV absorbance was measured at $\lambda = 595$ nm using a microplate spectrophotometer reader (Synergy HT, BioTek, Winooski, VT, USA).

4.2.2.4. Opioid receptor ligand binding assays. MOR and DOR binding experiments were carried out by incubating 400 μg per sample and 500 μg per sample of rat brain membranes, respectively, for 45 min at 35 °C with 1 nM [^3H]-DAMGO or 2 nM [^3H]-Deltorphin II (2-D-Ala)-[tyrosyl-3,5-3H] in 50 mM Tris-HCl (pH 7.4). For KOR binding assays, guinea pig brain membranes (400 μg per sample) were incubated for 30 min at 30 °C with 1 nM [^3H]-U69,593. Test compounds were added to a final volume of 1 mL. The K_d values of [^3H]-DAMGO, [^3H]-Deltorphin II (2-D-Ala)-[tyrosyl-3,5-3H] and [^3H]-U69,593 were 1.0, 1.5, and 2.3 nM, respectively. Nonspecific binding was assessed in the presence of 10 μM unlabelled naloxone. The reaction was terminated by filtering the solution under reduced pressure using a Millipore filter apparatus through Whatman glass fiber filters, GF/C for MOR and DOR and GF/B for KOR, presoaked for 1 h in a 0.1% poly(ethyleneimine) solution. Filters were washed with 50 mM ice-cold Tris-HCl buffer (2 \times 4 mL), dried at r.t., soaked overnight in 4 mL of scintillation cocktail into 6 mL pony vials, and counted on a liquid scintillation counter.

4.2.2.5. Data analysis. K_i values were calculated using nonlinear regression analysis to fit a logistic equation to the competition data using GraphPad Prism 9.3.0 (GraphPad Software Inc., San Diego, CA, USA).



4.3. Molecular modeling studies

4.3.1. Structures preparation and minimization. All the molecules used in this study were built using Marvin Sketch (18.24, ChemAxon Ltd.). The PM6-D3H4 Hamiltonian, implemented in the MOPAC package (MOPAC2016 v.18.151, Stewart Computational Chemistry, Colorado Springs), was then used to further optimize the 3D structures before the alignment for the docking calculations.

4.3.2. Docking and molecular dynamics studies. Flexible ligand docking experiments were performed using AutoDock, as implemented in the YASARA suite. The molecular dynamics (MD) simulations of the complexes were performed with the YASARA structure package according to our previously reported procedures.⁴³

4.4. Chemical stability. *In vitro* assay

First, a standard calibration curve (concentration range, 10–250 μM ; in methanol) was performed to calculate unknown sample concentrations (Fig. S3†). Samples were prepared as 10 mM solution in DMSO of compound 7, added to the medium previously heated at 37 °C to have a final concentration of 200 μM . The resultant solution was incubated at 37 \pm 0.5 °C, and at proper time intervals, an amount of 150 μL of the reaction mixture was withdrawn and added to the same amount of acetonitrile to obtain a final concentration of 100 μM . The sample was vortexed, filtered by 0.22 μm filters, and analyzed by HPLC. The experiment was performed in duplicate. The stability of compound 7 was evaluated over 300 minutes (Fig. S4†). The data were acquired with a HITA903-0337 Chromaster System, equipped with an autosampler for direct injection, a column oven, a quaternary gradient pump, and a diode array detector. The system was equipped with a Kinetex C18 5 μ (4 \times 3.0 mm ID) column, maintained at 25 °C and connected to a SecurityGuard Cartridge. A 30 min gradient run was performed, with a mobile phase A (acetonitrile 0.1% in TFA), a mobile phase B (H₂O 0.1% in TFA), a work flow at 1 ml min^{−1}, an injection volume of 10 μL and the detector wavelength of λ = 285 nm. The half-life ($t_{1/2}$) of compound 7 was calculated by matching the data with one phase exponential decay equation using Prism software 9.3.0 (Graph Pad, San Diego, CA, USA).

4.5. Stability in mouse plasma

The stability of compound 7 to the esterase was evaluated by incubating it in mouse C57 plasma. The solution of compound 7 (10 mM in DMSO) was added to plasma preheated at 37 °C; thus, obtaining a final concentration of the compound of 200 μM . The solution was incubated at 37° \pm 0.5 °C, and at appropriate time intervals (0, 1, 5, 20, and 60 minutes) an aliquot of 200 μL of reaction mixture was withdrawn and added to 200 μL of acetonitrile containing 0.1% HCOOH, to deproteinize the plasma. The final concentration was of 100 μM . The sample was vortexed and centrifuged for 15 minutes 13 000 rpm. The supernatant was filtered through 0.22 μm filters and analyzed by RP-HPLC. All

experiments were performed in duplicate. The HPLC analyses were performed as described above. The half-life ($t_{1/2}$) in mouse plasma of compound 7 was calculated by matching the data with one phase exponential decay equation with semi-log plots. The plateau was fixed to a constant value of '0'. All statistical analyses were performed with GraphPad Prism 8.0 (GraphPad Software, Inc., San Diego, CA).

4.6. *In vivo* pharmacology

4.6.1. Animals. Experiments were performed on male CD1 mice (Envigo Laboratories, Indianapolis, IN, USA) aged between 8 and 12 weeks. Mice were housed in 5 animals per cage under a 12/12 h light/dark cycle at a constant temperature (23–25 °C) with free access to food and water and were allowed to acclimate for at least one week upon arrival before starting all experiments, which were conducted between 9:00 a.m. and 3:00 p.m. This study was executed according to the European Communities Council directive and Italian regulation (EEC Council 2010/63/EU and Italian D.Lgs. no. 26/2014) to replace, reduce, and refine the use of laboratory animals. All procedures were approved by the ethical committee of the University of Catania (OPBA) and by the Italian Ministry of Health (authorization no. 385/2021-PR).

4.6.2. Mouse formalin test. Before starting every experiment, mice were randomly assigned to each experimental group and allowed to acclimate in the room for 20 min. The results were analyzed by a blind researcher to the experimental protocol. Formalin solution (5%, 10 μL) was administered subcutaneously into the plantar surface of the right hind paw (i.pl.), monitoring the nociceptive behavior, such as licking and flicking or shanking the injected paw, for 1 h and recording every 5 min, as previously reported.⁴⁴ The formalin injection induces a rapid pain response (phase I), which lasts about 5 min, due to direct nociceptors' activation, characterized by acute pain. After a short quiescent period, another behavioral pain form (phase II) occurs, considered more important clinically, characterized by an inflammatory component and persistent pain due to nociceptive sensitization in the dorsal horn of the spinal cord.

4.6.3. Drugs. Formalin was purchased from Merck. Compound 7 was dissolved in pyrogen-free isotonic saline and dimethyl sulfoxide (DMSO 0.5% v/v) and was administrated i.p. 20 min before formalin injection.

4.6.4. Data analysis. Results are expressed as mean \pm S.E. M. (n = 4–11 per group). *** p < 0.05. Multiple comparison testing was performed using ordinary two-way ANOVA followed by Bonferroni's multiple comparison test. Statistical analyses were performed using GraphPad Prism version 9.3.1 (GraphPad Software, San Diego, CA, USA).

Data availability

The data supporting my article have been included as part of the ESI† that I included in my manuscript (see file attached).



Author contributions

G. C., L. P. and C. P. were responsible for the study design, analysis, and interpretation of the data. G. C. synthesized, purified, and characterized the compounds. G. C. and G. Co. performed *in vitro* binding experiments. E. N. and G. M. P. performed HPLC analysis. R. M. performed chiral HPLC analysis. V. P., A. C., and A. R. conceived the computational molecular modeling experiments. M. G., S. Z., and C. P. conducted *in vivo* pharmacology experiments and described and discussed the relative results. G. C., L. P., C. P., V. P., and A. R. drafted the main text. M. A., E. A., and A. M. contributed to the writing of the manuscript. All authors have participated in the refinement of the writing and have approved the final version of the manuscript.

Conflicts of interest

The authors declare no competing financial interest.

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