

## Ugr 9-1

<b>Product name:</b> Ugr 9-1	<b>Synonyms :</b> $\pi$ -AnmTX Ugr 9a-1
<b>Catalog # :</b> 13UGR001	
<b>Product description</b> <p>Ugr 9-1 (<math>\pi</math>-AnmTX Ugr 9a-1) was isolated from the venom of the sea anemone <i>Urticina grebelnyi</i>. Ugr 9-1 reversibly inhibits homomeric human ASIC3. Ugr9-1 fully inhibits transient currents with an IC<sub>50</sub> value of around 10 <math>\mu</math>M and also partially blocks sustained currents (IC<sub>50</sub> = 1.4 <math>\mu</math>M). The peculiarity of this peptide is its ability to block ASIC3 when activated at weak pH (pH 4.0). Ugr 9-1 does not block or activate ASIC1a, ASIC1b, ASIC2a and Kv1.3 at significant concentrations.</p> <p>Ugr 9-1 significantly reverses inflammatory and acid-induced pain in-vivo in mice. The antinociceptive effect is attributed to the block of the sustained inward Na<sup>+</sup> current in sensitive neurons.</p>	
<b>Product specifications</b> <p><b>AA sequence:</b> Ile-Ser-Ile-Asp-Pro-Pro-Cys<sup>7</sup>-Arg-Phe-Cys<sup>10</sup>-Tyr-His-Arg-Asp-Gly-Ser-Gly-Asn-Cys<sup>19</sup>-Val-Tyr-Asp-Ala-Tyr-Gly-Cys<sup>26</sup>-Gly-Ala-Val-OH</p> <p><b>Disulfide bonds:</b> Cys<sup>7</sup>-Cys<sup>19</sup>, Cys<sup>10</sup>-Cys<sup>26</sup></p> <p><b>Length (aa):</b> 29</p> <p><b>Formula:</b> C<sub>134</sub>H<sub>196</sub>N<sub>38</sub>O<sub>42</sub>S<sub>4</sub></p> <p><b>Appearance:</b> White lyophilized solid</p> <p><b>Molecular Weight:</b> 3135 Da</p> <p><b>CAS number:</b> NA</p> <p><b>Source:</b> Synthetic</p> <p><b>Counterion:</b> TFA salts</p> <p><b>Solubility:</b> Water or saline buffer, 5 mg/mL maximum (recommendation)</p>	
<b>Formulation</b> <p><b>Storage/Stability:</b> Shipped at ambient temperature under lyophilized powder. Store at -20°C (-4°F). Do not freeze-thaw. Aliquot sample if required and store at -80°C (-112°F).</p> <p><b>Expiry date:</b> One year</p> <p><b>Use restrictions:</b> For laboratory use only. Not for drug, household or other uses. Not for use in diagnostic or therapeutic procedures.</p>	
<b>Related products</b> <ul style="list-style-type: none"> <li>• <a href="#">Psalmotoxin 1 - #13PCT001</a>: selective ASIC1a blocker</li> <li>• <a href="#">APETx2 - #07APE002</a>: ASIC3 selective blocker</li> </ul>	
<b>References</b> <ul style="list-style-type: none"> <li>• Osmakov D., <i>et al.</i> (2013) Sea anemone peptide with uncommon <math>\beta</math> -hairpin structure inhibits acid-sensing ion channel 3 (ASIC3) and reveals analgesic activity. <i>JBC</i></li> </ul>	

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