



## ICR-1 AM

### Lot 10120a

Method	Specification	Analysis
<b>LCMS</b>	<b>Agilent 1220 Infinity II</b>	
Purity*	≥ 90%	94.3 %
Molecular Ion	<i>Common Peaks</i> 1190.28 ± 0.5 m/z (MH <sup>+</sup> ) 1212.26 ± 0.5 m/z (MNa <sup>+</sup> ) 595.65 ± 0.5 m/z (MH <sub>2</sub> <sup>2+</sup> )	<i>Detected Peaks</i> 1190.6 m/z 1212.6 m/z 595.9 m/z
<b>Absorbance Spectrum</b>	<b>Agilent Cary 60 UV-VIS Spectrophotometer</b>	
Longest-Wavelength Absorbance Maximum**	577 ± 3 nm	578 nm
<b>Fluorescence Spectrum</b>	<b>Horiba Jobin Yvon FluoroMax 4 Spectrofluorometer</b>	
Excitation Max.; Emission Max.**	585 ± 3 nm; 644 ± 3 nm	585 nm; 643 nm
<b><sup>1</sup>H NMR Spectrum</b>	<b>Bruker Avance 400</b>	
Peaks and Integrations	Only relevant product peaks — with appropriate chemical shifts and peak integrations — and solvent peaks present	Confirmed
<b>Cell Assay</b>	<b>BioTek Cytation 5 Imaging Reader</b>	
F/F <sub>0</sub> post-stimulus in relevant biological assay	≥ 1.2	1.5

\*Column: Agilent Infinity Lab Poroshell 120 ECC18, 3.0 x 50 mm, 2.7 μm C<sub>18</sub>, UV-Vis Diode Array Detector: 254 nm, Single Quad MS Detector: ESI Positive; \*\*solvent: High-Calcium Buffer, AM esters hydrolyzed to ion-sensing salt form prior to acquiring spectral data.

Approved by P. Rogelio Escamilla Oct 2019