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CORRECTION
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Corrigendum: Metabotropic Glutamate Receptor 7: A New Therapeutic Target in Neurodevelopmental Disorders

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A Corrigendum on

Metabotropic Glutamate Receptor 7: A New Therapeutic Target in Neurodevelopmental Disorders

by Fisher, N. M., Seto, M., Lindsley, C. W., and Niswender, C. M. (2018). Front. Mol. Neurosci. 11:387. doi: 10.3389/fnmol.2018.00387

In the original article, there was a mistake in **Figure 1** as published. The chirality of L-AP4 and LSP1-2111 was incorrect. pEC_{50} values have also been corrected for LSP1-2111 in **Table 1**. The authors apologize for these errors and state that they do not change the scientific conclusions of the article in any way. The original article has been updated.

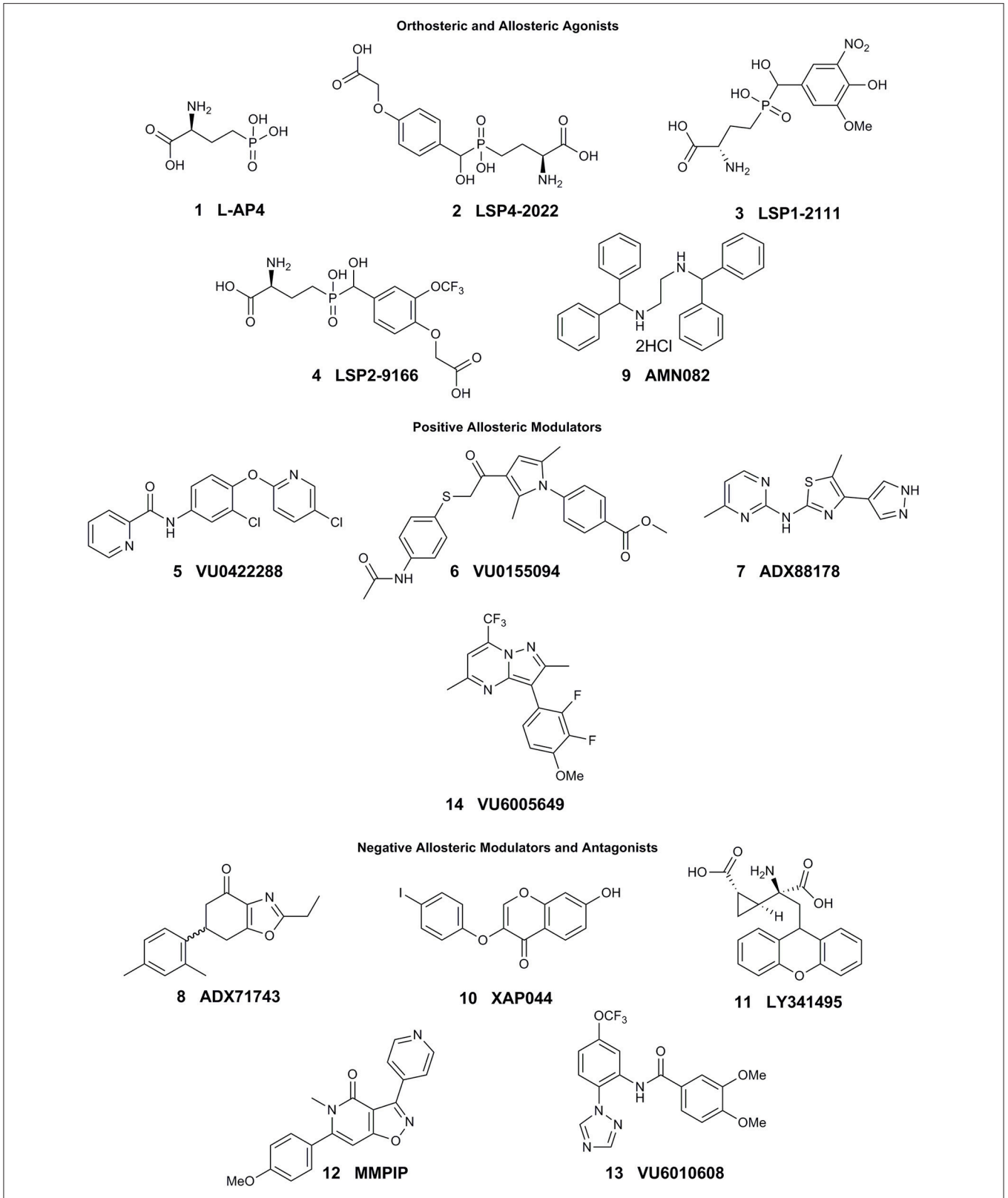

FIGURE 1 | Current tool compounds used to study $mGlu_7$.

TABLE 1 | Summary of current tool compounds used to study $mGlu_7$.

Name (#)	Type	$mGlu_7$ pEC_{50}/pIC_{50}	$mGlu_8$ pEC_{50}/pIC_{50}	$mGlu_4$ pEC_{50}/pIC_{50}	$mGlu_6$ pEC_{50}/pIC_{50}	Source
L-AP4 **(1)**	Orthosteric agonist	3.47 (PIH)	6.53 (PIH)	7.00 (PIH)	5.62 (PIH)	Acher et al., 2012; Selvam et al., 2018
		3.61 (Ca^{2+})	6.53 (Ca^{2+})	6.89 (Ca^{2+})	6.00 (Ca^{2+})	
LSP4-2022 **(2)**	Orthosteric agonist	4.34 (Ca^{2+})	4.54 (Ca^{2+})	6.96 (Ca^{2+})	5.36 (Ca^{2+})	Acher et al., 2012; Goudet et al., 2012; Selvam et al., 2018
LSP1-2111 **(3)**	Orthosteric agonist	4.28 (PIH)	4.18 (PIH)	5.66 (PIH)	5.77 (PIH)	Selvam et al., 2018
		4.00 (Ca^{2+})	4.71 (Ca^{2+})	6.05 (Ca^{2+})	5.49 (Ca^{2+})	
LSP2-9166 **(4)**	Orthosteric agonist	5.71 (Ca^{2+})	4.25 (Ca^{2+})	7.22 (Ca^{2+})	not reported	Acher et al., 2012
VU0422288 **(5)**	Group III PAM	6.85 (Ca^{2+})	6.93 (Ca^{2+})	6.98 (Ca^{2+})	not reported	Jalan-Sakrikar et al., 2014
VU0155094 **(6)**	Group III PAM	5.80 (Ca^{2+})	6.07 (Ca^{2+})	5.48 (Ca^{2+})	not reported	Jalan-Sakrikar et al., 2014
ADX88178 **(7)**	$mGlu_{4/8}$ PAM	>4.52 (Ca^{2+})	5.66 (Ca^{2+})	8.46 (Ca^{2+})	>5	Le Poul et al., 2012
ADX71743 **(8)**	$mGlu_7$ NAM	7.20 (human, Ca^{2+})	inactive	inactive	inactive	Kalinichev et al., 2014
		7.06 (rat, Ca^{2+})	inactive	inactive	inactive	
AMN082 **(9)**	Allosteric agonist	6.59 (GTPγS)	>5 (GTPγS)	>5 (GTPγS)	>5 (GTPγS)	Mitsukawa et al., 2005
XAP044 **(10)**	Antagonist	5.26 (cAMP)	4.48 (cAMP)	inactive	inactive	Gee et al., 2014
		5.55 to 5.46 (GTPγS)				
LY341495 **(11)**	Orthosteric antagonist	6.00 (cAMP)	6.76 (cAMP)	4.66 (cAMP)	not reported	Kingston et al., 1998
MMPIP **(12)**	$mGlu_7$ NAM	6.66 (cAMP)	>5 (cAMP)	>5 (cAMP)	not reported	Suzuki et al., 2007
		7.15 (Ca^{2+})				Niswender et al., 2010
		6.14 (Thallium)				Niswender et al., 2010
VU6010608 **(13)**	$mGlu_7$ NAM	6.12 (Ca^{2+})	>5 (Ca^{2+})	>5 (Ca^{2+})	inactive (>5)	Reed et al., 2017
VU6005649 **(14)**	$mGlu_{7/8}$ PAM	6.19 (Ca^{2+})	5.59 (Ca^{2+})	>5 (Ca^{2+})	inactive	Abe et al., 2017

NAM, negative allosteric modulator; PAM, positive allosteric modulator; EC_{50}, effective concentration 50; IC_{50}, inhibitory concentration 50. Assay type is indicated in parentheses: PIH, Phosphatidylinositol hydrolysis; cAMP, cAMP accumulation; Ca^{2+}, Calcium mobilization; GTPγS, GTPγS binding.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.