www.bioss.com.cn
sales@bioss.com.cn
techsupport@bioss.com.cn
400-901-9800

PLX51107

产品编号：	D60376
CAS:	1627929-55-8
分子式：	C26H22N4O3
纯度：	98%
InChi Key:	InChIKey=AMSUHYUVOVCWTP-INIZCTEOSA-N
Smiles:	C[C@H](N1C=C(C2=C1C=C(C=N2)C1=C(C)ON=C1C)C1=CC=C(C=C1)C(O)=O)C1=CC=CC=N1
外观：	固体粉末
作用通路：	Chromatin/Epigenetic
作用靶点：	BRD2-BD1; BRD3-BD1; BRD4-BD1; BRDT-BD1; BRD2-BD2; BRD3-BD2; BRD4-BD2; BRDT-BD2); CBP; EP300; Epigenetic Reader Domain
溶解性：	Soluble in DMSO
保存条件：	-20°C
产品介绍：	PLX51107 is a potent and selective BET inhibitor, with Kds of 1.6, 2.1, 1.7, and 5 nM for BD1 and 5.9, 6.2, 6.1, and 120 nM for BD2 of BRD2, BRD3, BRD4, and BRDT, respectively; PLX51107 also interacts with the bromodomains of CBP and EP300 (Kd, in the 100 nM range).