

# INPP4B Is a PtdIns(3,4,5)P<sub>3</sub> Phosphatase That Can Act as

Cancer Discovery

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Precise control of localized signals. <i>Nature</i> , 2015, 522, 38-40.	13.7	4
2	INPP4B Is a Tumor Suppressor in the Context of PTEN Deficiency. <i>Cancer Discovery</i> , 2015, 5, 697-700.	7.7	17
3	Phosphoinositide signaling in cancer: INPP4B Akt(s) out. <i>Trends in Molecular Medicine</i> , 2015, 21, 530-532.	3.5	26
4	PIPPing on AKT1: How Many Phosphatases Does It Take to Turn off PI3K?. <i>Cancer Cell</i> , 2015, 28, 143-145.	7.7	9
5	A Kinase Divided. <i>Cancer Cell</i> , 2015, 28, 145-147.	7.7	2
6	The extended human <i>PTP</i> ome: a growing tyrosine phosphatase family. <i>FEBS Journal</i> , 2016, 283, 1404-1429.	2.2	90
7	Emerging evidence of signalling roles for PI(3,4)P <sub>2</sub> in Class I and II PI3K-regulated pathways. <i>Biochemical Society Transactions</i> , 2016, 44, 307-314.	1.6	96
8	Regulation of PtdIns(3,4,5)P <sub>3</sub> /Akt signalling by inositol polyphosphate 5-phosphatases. <i>Biochemical Society Transactions</i> , 2016, 44, 240-252.	1.6	53
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15	Regulation of PI3K effector signalling in cancer by the phosphoinositide phosphatases. <i>Bioscience Reports</i> , 2017, 37, .	1.1	91
16	INPP4B and PTEN Loss Leads to PI-3,4-P <sub>2</sub> Accumulation and Inhibition of PI3K in TNBC. <i>Molecular Cancer Research</i> , 2017, 15, 765-775.	1.5	26
17	<i>INPP4B</i> overexpression suppresses migration, invasion and angiogenesis of human prostate cancer cells. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2017, 44, 700-708.	0.9	14
18	IRF2-INPP4B axis participates in the development of acute myeloid leukemia by regulating cell growth and survival. <i>Gene</i> , 2017, 627, 9-14.	1.0	13

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19	MicroRNA-590-3p promotes cell proliferation and invasion by targeting inositol polyphosphate 4-phosphatase type II in human prostate cancer cells. <i>Tumor Biology</i> , 2017, 39, 101042831769594.	0.8	11
20	PTEN Regulates PI(3,4)P2 Signaling Downstream of Class I PI3K. <i>Molecular Cell</i> , 2017, 68, 566-580.e10.	4.5	149
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