MedChemExpress

Cat. No.:	HY-N7830	
CAS No.:	155205-64-4	
Molecular Formula:	$C_{20}H_{28}O_4$	1. J
Molecular Weight:	332.43	
Target:	ERK	
Pathway:	MAPK/ERK Pathway; Stem Cell/Wnt	HO´
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	
	Anatysis.	

,OH

BIOLOGICAL ACTIVITY					
Description	7α ,15-Dihydroxydehydroabietic acid is a natural abietane-type diterpenoid with antiangiogenic effects ^[1] .				
In Vitro	 7α,15-Dihydroxydehydroabietic acid (3.125-100 μM; 24 hours) significantly decreases HUVEC cell viability^[1]. 7α,15-Dihydroxydehydroabietic acid (3.125-6.25 μM; 24 hours) significantly inhibits the promotion of angiogenesis in HUVECs. 7α,15-Dihydroxydehydroabietic acid inhibits angiogenesis through downregulation of the VEGF, p-Akt and p-ERK signaling pathways^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay^[1] 				
	Cell Line:	Human umbilical vein endothelial cells (HUVECs)			
	Concentration:	3.125 μΜ, 6.25 μΜ, 12.5 μΜ, 25 μΜ, 50 μΜ, 100 μΜ			
	Incubation Time:	24 hours			
	Result:	Significantly decreased HUVEC cell viability.			
	Western Blot Analysis ^[1]				
	Cell Line:	Human umbilical vein endothelial cells (HUVECs)			
	Concentration:	3.125 μΜ, 6.25 μΜ			
	Incubation Time:	24 hours			
	Result:	Human umbilical vein endothelial cells (HUVECs) ^[1] 3.125 μ M, 6.25 μ M24 hoursShowed downregulation of VEGF, p-Akt and p-ERK in HUVECs.			

REFERENCES

[1]. Tae Kyoung Lee, et al. 7α,15-Dihydroxydehydroabietic acid from Pinus koraiensis inhibits the promotion of angiogenesis through downregulation of VEGF, p-Akt and p-ERK in HUVECs. Bioorg Med Chem Lett. 2018 Apr 1;28(6):1084-1089.

Caution: Product has not been fully validated for medical applications. For research use only.

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA