Joint Event 4th EUROPEAN BIOPHARMA CONGRESS &

& 6WK, QWHUQDWLRQDO &RQIHUHQFH DQG (PHARMACOLOGY AND ETHNOPHARMACOLOGY

November 09-11, 2017 Vienna, Austria

Perspective of TRAIL and PEGylated TRAIL

its cognate receptors in cancer cells without apparent toxicity to normal cells. TRAIL has been considered as an anticar
drug due to its unique ability to selectively induce DR-mediated apoptosis in transformed cells. To date, recombinant huma
TRAIL and antibodies directed against TRAIL-R1 or TRAIL-R2 have been tested clinically. However, these have be
disappointing, showing a very limited bene t as an antitumor agent basically due to their poor agonistic activity of thes
agents. And in recent years, the physiological importance of TRAIL has expanded beyond being a tumoricidal molecule
one critical for a number of clinical settings - ranging from brosis and autoimmunity to cardiovascular anomalies. In an
attempt to overcome the poor agonistic activity and also low stability and solubility of rTRAIL, iww ideveloped a delivery
system by using PEGTd [(, .a (o o)1n(p)c)-3 (a)-4.9 (I si (d)0.s4)8 csi(419 (t)-(p)c)-3 (a)-4.9 t we d.d d b i (em)19 ()0

Notes: