

NEW PARP-1 INHIBITORS FOR IMAGING PARP-1 EXPRESSION IN VIVO

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[¹⁸F]WC-DZ-F is a derivative of [¹⁸F]FluorThanatrace and a radiotherapeutic agent. [¹⁸F]WC-DZ-F has suitable biodistribution profile, favourable logP and super metabolic stability in vivo. We have demonstrated that [¹⁸F]WC-DZ-F is capable of imaging PARP-1 expression in PC-3 prostate cancer xenograft tumor in small animal. [¹⁸F]WC-DZ-F can be a replacement of [¹⁸F]FluorThanatrace in the

imaging of PARP-1 expression.