

# KRICT | Reversible BTK-non-ITK inhibitor

Target	
<b>Mechanism of Action</b>	<ul style="list-style-type: none"> <li>• Bruton's tyrosine kinase inhibitor</li> </ul>
<b>Indication - Primary</b>	<ul style="list-style-type: none"> <li>• Hematological cancers (mantle cell lymphoma, chronic lymphocytic leukemia)</li> </ul>
<b>Indication - Expansion</b>	<ul style="list-style-type: none"> <li>• Rheumatoid arthritis (reversible BTK inhibitor)</li> <li>• Parkinson's disease (Abl inhibitor)</li> </ul>
<b>Route of Administration</b>	<ul style="list-style-type: none"> <li>• PO, QD</li> </ul>
<b>Competitive Advantage</b>	<ul style="list-style-type: none"> <li>• Reversible BTK inhibitor</li> <li>• Neither ITK nor EGFR inhibition → ADCC &amp; improved adverse effect profiles</li> <li>• More active than ibrutinib in a murine xenograft model using TMD-8 cells</li> </ul>
<b>Data Files</b>	<ul style="list-style-type: none"> <li>• In vitro: enzymes (BTK, BTK C481S, Abl), cell (TMD-8, hPBMC) kinase profiling</li> <li>• In vivo: murine xenograft model using TMD-8 cells, CIA model</li> <li>• DMPK: CYP, metabolic stability, Patch clamp, solubility, PK, BBB</li> </ul>
<b>IP Status</b>	<ul style="list-style-type: none"> <li>• Patent filing underway</li> </ul>
<b>Collaboration Model</b>	<ul style="list-style-type: none"> <li>• License-out</li> <li>• Collaborative research under funding</li> </ul>
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