

# A Study on the Development of USP1 inhibitor candidate as a New Synthetic Lethality Target for the Treatment of BRCA (-) and HRD (+) Cancer

AIGEN Sciences



<b>Disease Area</b>	<b>Oncology</b>
<b>Product Type</b>	Small molecule
<b>Indication</b>	BRCA (-) cancers, HRD (+) cancers, CCNE-1 (+) cancers
<b>Target</b>	USP1
<b>Mechanism of Action</b>	Synthetic lethality by the ubiquitination of PCNA
<b>Competitiveness</b>	KSQ4279: phase 1 ISM-3091: phase 1
<b>Development Stage</b>	<b>Lead optimization (Lead to candidate)</b>
<b>Route of Administration</b>	oral

<b>Key Data</b>	<p><b>AIG07016: lead compound</b></p> <ul style="list-style-type: none"> <li>- Good selectivity for DUB panel</li> <li>- Good selectivity for BRCA wild type cancer cell</li> <li>- Synergism with PARP inhibitor</li> </ul>
	<p>Selectivity for DUB panel</p> <p>Cell growth inhibition: BRCA1 mt vs. wt</p> <p>Combination with Olaparib</p>

<b>IP</b>	Filed
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