

# Efficacy of the IRAK4 Inhibitor CA-4948 in Patient-Derived Xenograft Models of Diffuse Large B Cell Lymphoma

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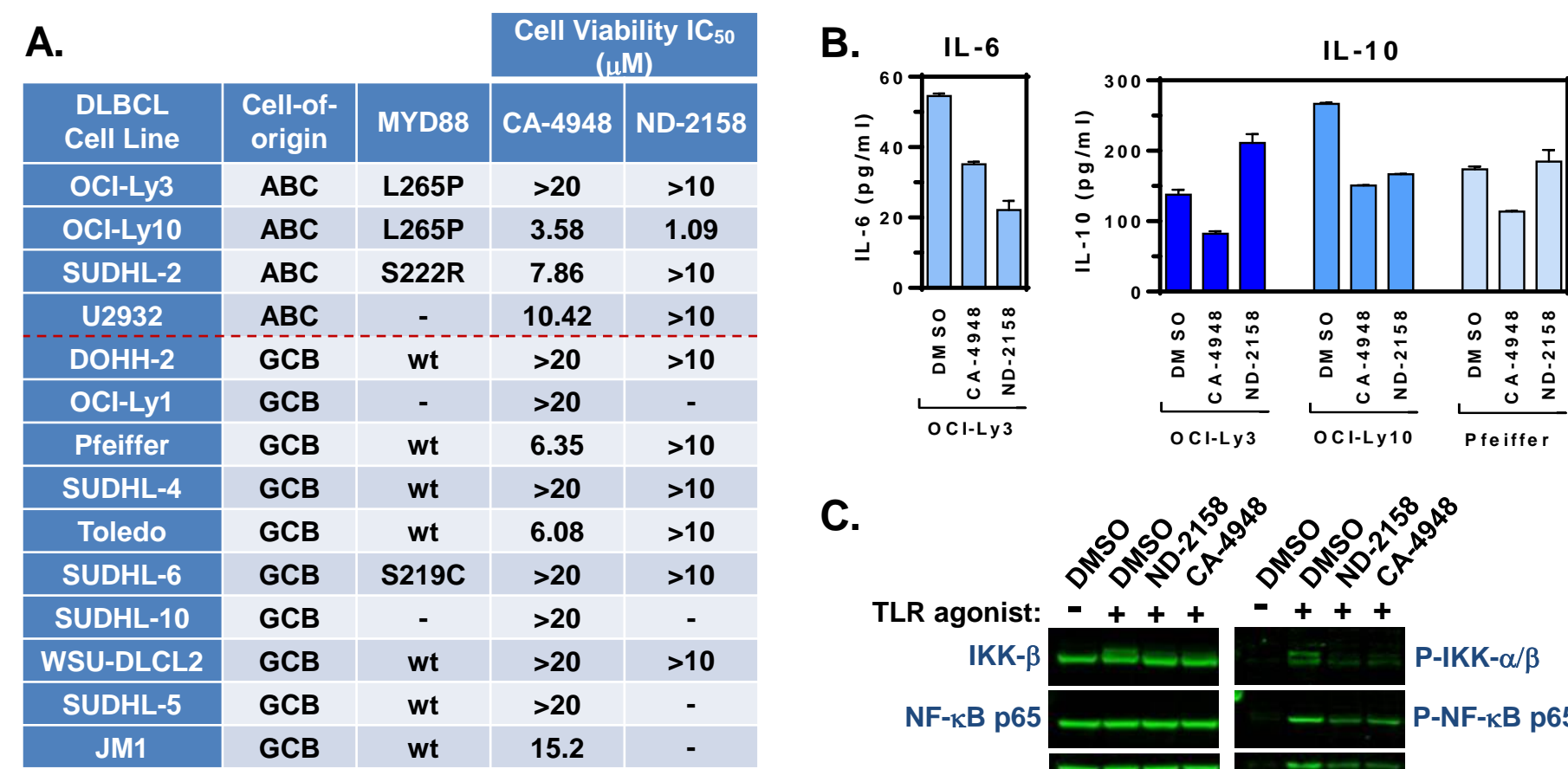
## Abstract

IRAK4 kinase activity is required for toll-like receptor (TLR) and interleukin-1 receptor (IL-1R) signaling in a variety of myeloid and lymphoid cell types. Recruitment of IRAK4 to these receptors and its subsequent activation is facilitated by the MYD88 adaptor protein, which is mutated in ~22% of DLBCL cases. The MYD88 L265P activating mutation is found in ~30% of the activated B-cell (ABC) and ~6% of germinal center B-cell (GCB) subtypes of DLBCL and leads to constitutive activation of NF- $\kappa$ B signaling that is associated with worse prognosis. Thus, the development of small molecule inhibitors targeting IRAK4 is an attractive anticancer strategy for MYD88 mutation-containing cancers such as DLBCL.

We are developing an IRAK4 inhibitor, CA-4948, as a therapeutic agent for hematological cancers with dysregulated TLR/MYD88/IRAK4 signaling. CA-4948 (previously AU-4948) is a selective and potent IRAK4 kinase inhibitor with in vivo activity in a TLR4-induced cytokine release model. CA-4948 exhibits favorable DMPK properties, oral bioavailability, and is well tolerated in mice. Furthermore, CA-4948 was previously shown to exhibit dose-dependent efficacy in ABC-DLBCL MYD88-L265P xenograft tumor models using cell lines OCI-LY3 and OCI-LY10.

Here, we report the efficacy results from testing CA-4948 in a panel of well characterized, patient-derived DLBCL tumor xenograft (PDX) mouse models. CA-4948 exhibited the greatest efficacy in four of the five ABC-DLBCL PDX models tested as compared to GCB-DLBCL and ABC/GCB DLBCL PDX models. Furthermore, CA-4948 was efficacious in ABC-DLBCL PDX tumors containing activating mutations in both TLR/IL-1R and BCR signaling pathways (MYD88 and CD79B double mutants). Interestingly, the one ABC-DLBCL PDX model that failed to respond to CA-4948 treatment contained a MYD88 L265P mutation as well as a BCL6 translocation. While this particular PDX model was resistant to CA-4948, and showed a weak anti-tumor response to single-agent ibrutinib, the combination treatment of ibrutinib and CA-4948 exhibited a synergistic tumor growth inhibition effect. In summary, CA-4948 exhibited anti-tumor activity in ABC-type DLBCL PDX tumor models including those containing combinations of activating mutations in the TLR/IL-1R and BCR signaling pathways. These results underscore the therapeutic potential of IRAK4 kinase inhibition by CA-4948, as a single-agent or in combination with BCR inhibitors, for the treatment of DLBCL.

## CA-4948 In Vitro Effect on DLBCL Cell Lines

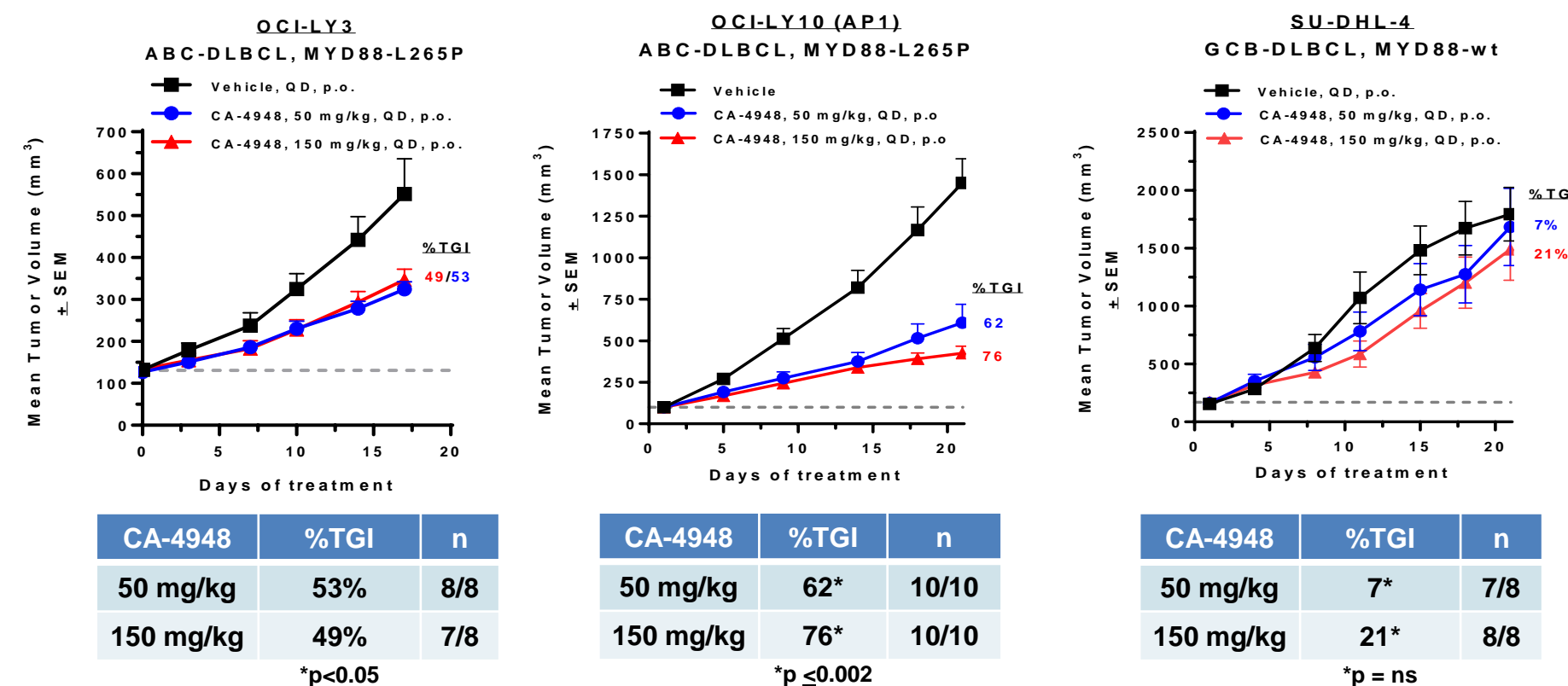


**A.** Viability IC<sub>50</sub> values determined after 72 hr treatment (CellTiter-Glo) with IRAK4 inhibitors CA-4948 and ND-2158

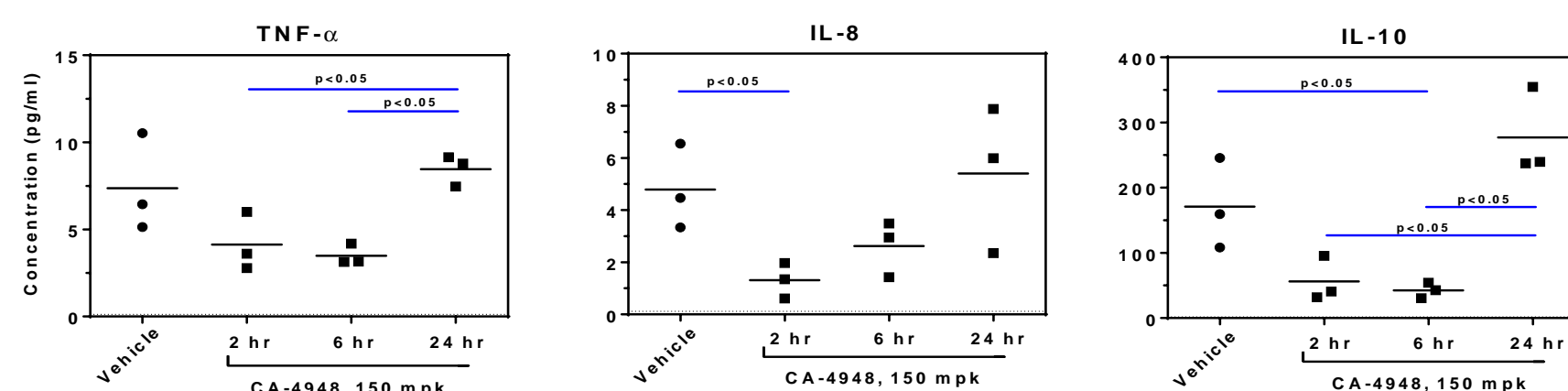
**B.** Level of cytokines present in media after 21 hr treatment with 10 μM of IRAK4 inhibitors

**C.** Western blot analysis of a DLBCL cell line during a 20 min TLR-agonist stimulation with 10 μM IRAK4 inhibitors ND-2158 and CA-4948

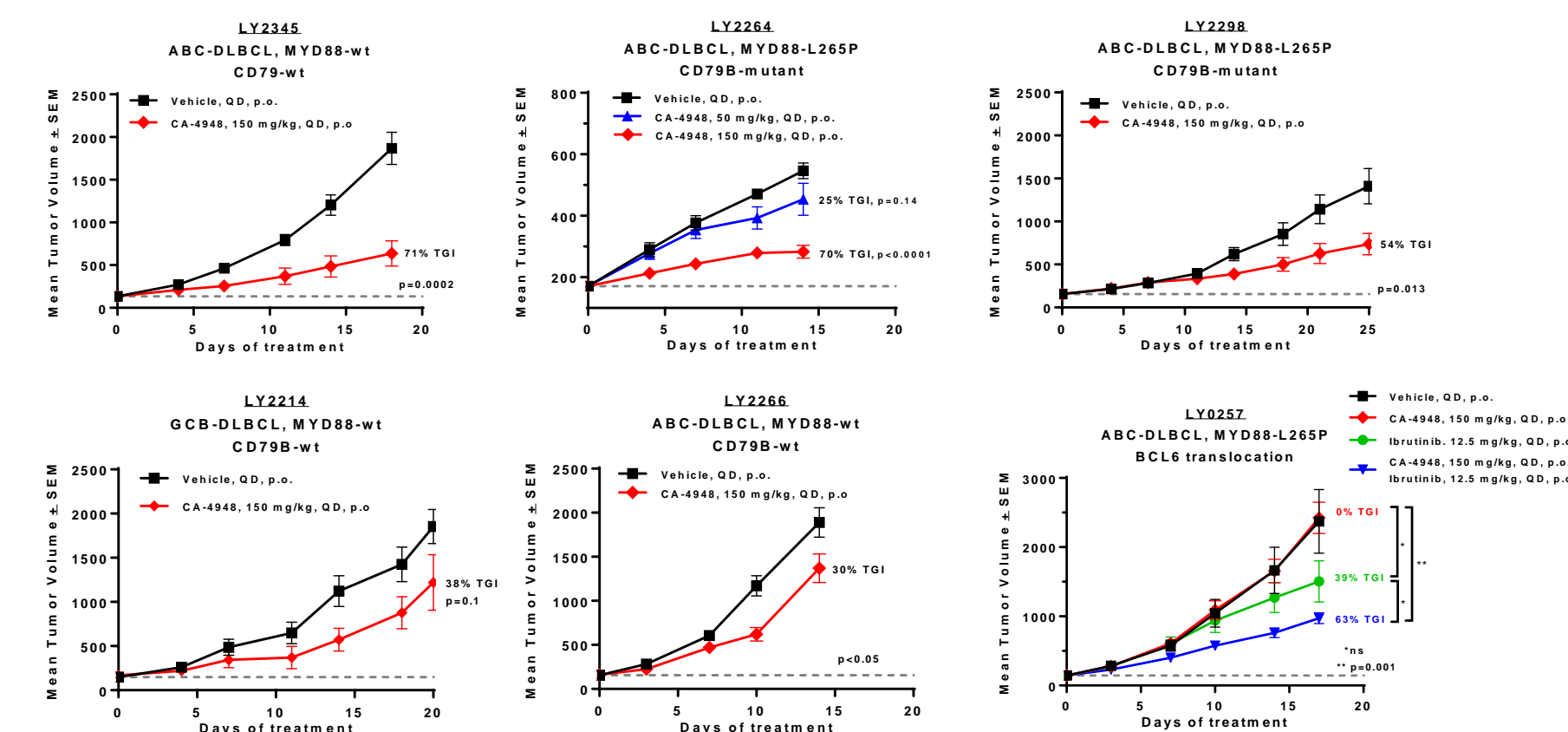
## CA-4948 Exhibits Enhanced In Vivo Efficacy in MYD88-L265P Mutant Versus MYD88-wt DLBCL Tumors



## CA-4948 Pharmacodynamic Response in DLBCL PDX Tumor-Bearing Mice



## CA-4948 Efficacy in 6 DLBCL PDX Tumor Models



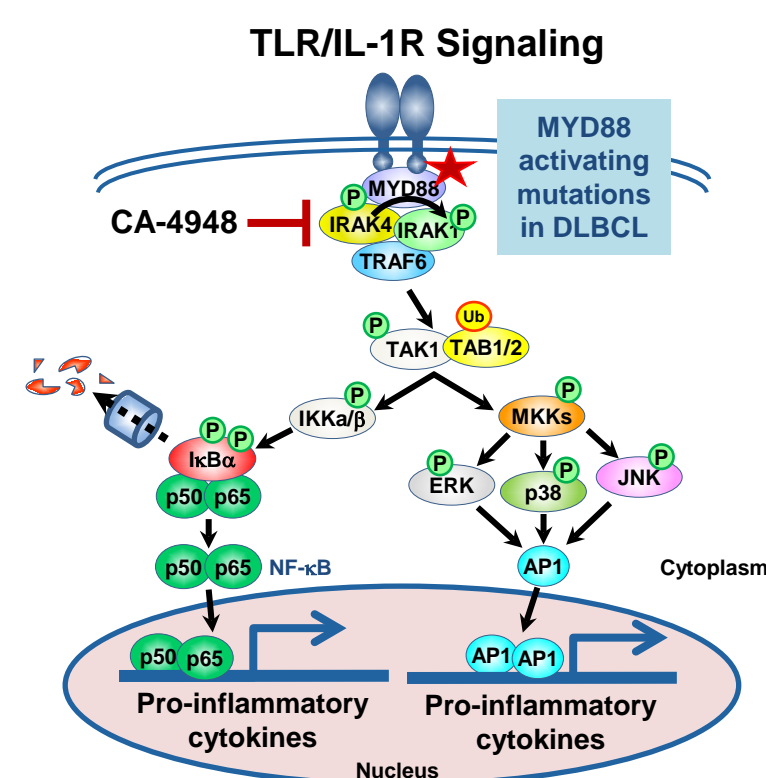
DLBCL PDX Model	COO Subtype Classification		Mutations/Expression			CA-4948 Efficacy
	Gene Signature	Hans IHC	MYD88 Mutation	B-Cell Receptor Mutation	BCL6 Expression (IHC)	150 mg/kg QD, %TGI
LY2345	ABC	non-GCB	WT	-	2%	71**
LY2264	ABC	non-GCB	L265P	CD79B	60%	70**
LY2298	ABC	non-GCB	L265P	CD79B	20%	54*
LY2214	GCB	non-GCB	WT	WT	95%	38
LY2266	ABC/GCB	non-GCB	WT	WT	0%	30*
LY0257	ABC	non-GCB	L265P	WT	95%	0

\*p < 0.05, \*\*p < 0.001

## CA-4948 IRAK4 Inhibitor for Treatment of DLBCL

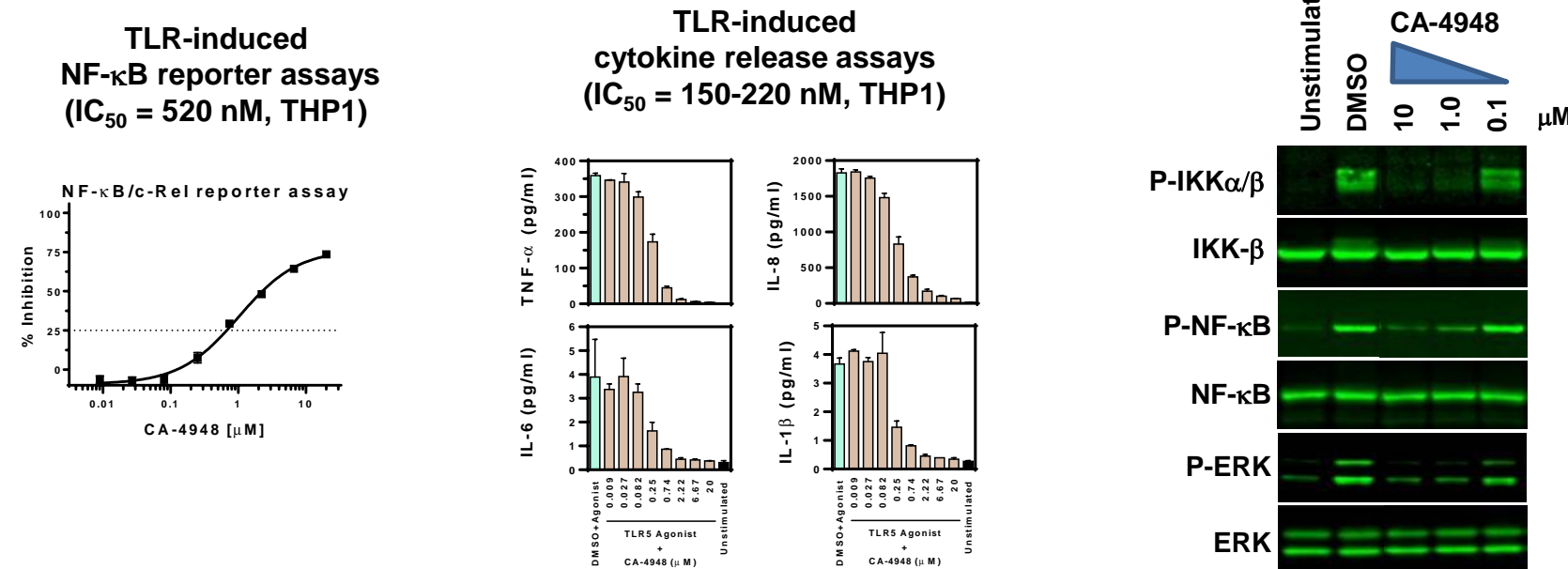
**CA-4948 Kinase Inhibitory Activity**

Kinase	Kd (nM)
IRAK4	23
IRAK1	>100



## CA-4948 Blocks the TLR/IL-1R Induced Canonical NF-κB Signaling Pathway

In vitro THP1 monocytic cell assays



## Summary

- CA-4948 is a potent IRAK4 Ser/Thr kinase inhibitor with >100-fold less activity against IRAK1
- CA-4948 exhibited weak to no anti-proliferative effect against DLBCL cell lines grown in 10% FBS. However, in ABC-DLBCL cells, CA-4948 repressed constitutive IL6 and IL10 cytokine secretion and TLR-agonist stimulation of the TLR/NF- $\kappa$ B signaling pathway
- CA-4948 exhibited greater efficacy in ABC-DLBCL, MYD88-L265P cell line xenograft models (OCI-LY3 and OCI-LY10) compared to a GCB-DLBCL, MYD88-WT model (SU-DHL-4)
- A single oral dose of 150 mg/kg CA-4948 repressed tumor-derived cytokine levels in the plasma of a DLBCL PDX model
- CA-4948 exhibited efficacy (>50% TGI) in 3-of-5 ABC-DLBCL PDX models, independent of MYD88 or B-cell receptor (BCR) mutations
- High BCL6 expression in the ABC-DLBCL, MYD88-L265P PDX tumor model LY0257 may be the cause of the tumor's resistance to CA-4948 treatment
- In conclusion, these results support the continued investigation of CA-4948, as a single-agent or in combination with BCR inhibitors, for the treatment of DLBCL

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