

Allosteric activation of apicomplexan calcium-dependent protein kinases

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Calcium-dependent protein kinases (CDPKs) comprise the major group of Ca^{2+} -regulated kinases in plants and protists. It has long been assumed that CDPKs are activated, like other Ca^{2+} -regulated kinases, by derepression of the kinase domain (KD). However, we found that removal of the autoinhibitory domain from *Toxoplasma gondii* CDPK1 is not sufficient for kinase activation. From a library of heavy chain-only antibody fragments (VHHs), we isolated an antibody (1B7) that binds *TgCDPK1* in a conformation-dependent manner and potently inhibits it. We uncovered the molecular basis for this inhibition by solving the crystal structure of the complex and simulating, through molecular dynamics, the effects of 1B7-kinase interactions. In contrast to other Ca^{2+} -regulated kinases, the regulatory domain of *TgCDPK1* plays a dual role, inhibiting or activating the kinase in response to changes in Ca^{2+} concentrations. We propose that the regulatory domain of *TgCDPK1* acts as a molecular splint to stabilize the otherwise inactive KD. This dependence on allosteric stabilization reveals a novel susceptibility in this important class of parasite enzymes.

calcium-dependent protein kinase | kinase activation | VHH

Calcium-dependent protein kinases (CDPKs) are major transducers of Ca^{2+} signaling in plants and protists. CDPKs regulate many essential processes in medically important parasites, including the apicomplexans responsible for malaria (*Plasmodium* spp.), toxoplasmosis (*Toxoplasma gondii*), and cryptosporidiosis (*Cryptosporidium parvum*). In these parasites, CDPKs control Ca^{2+} -dependent secretion, parasite motility, gametogenesis, as well as invasion and egress from infected cells (reviewed in ref. 1). Their absence from mammalian genomes makes CDPKs attractive drug targets. Inhibition of *TgCDPK1* (*T. gondii* CDPK1) phenocopies a conditional knock-down of the kinase by blocking Ca^{2+} -regulated secretion (2). The closest paralog of *TgCDPK1*, *TgCDPK3*, regulates the initiation of *T. gondii* egress from host cells (3–5), and its homolog in *Plasmodium falciparum*, *PfCDPK1*, is essential (6).

CDPKs comprise an N-terminal kinase domain (KD), linked by a junction region to four EF hands that form a calmodulin-like domain (CaM-LD) and bind Ca^{2+} . Studies of plant CDPKs suggest that the key mechanism of regulation is inhibition of the KD by the junction region preceding the EF hands, predicted to act as a pseudosubstrate (7, 8). The junction region interacts with the CaM-LD, and Ca^{2+} binding alters this interaction, presumably to relieve inhibition of the KD (9, 10). Structures of *TgCDPK1* corroborated these predictions (11, 12) and showed the extreme Ca^{2+} -dependent rearrangement of the CaM-LD, which warranted its distinction as a new regulatory domain, dubbed the CDPK activation domain (CAD) (11). In the absence of Ca^{2+} , the junction domain and the first α -helix of EF hand I form a single long α -helix that extends from the base of the KD, occupies the substrate-binding site, and occludes the catalytic pocket (11). This structure is reminiscent of the autoinhibited state of mammalian Ca^{2+} /CaM-dependent protein kinases (CaMKs) (13, 14), reinforcing the notion that autoinhibition is the major mechanism of CDPK regulation.

Here we show that removal of the regulatory domain of *TgCDPK1* is not sufficient to activate the kinase. To identify new tools to probe the mechanism of kinase activation, we turned to single domain antibody fragments, or VHHs, derived from alpaca (*Vicugna pacos*) heavy chain-only antibodies. In screening phage display libraries of VHHs from alpacas naturally infected with *T. gondii*, we isolated a single domain antibody (VHH 1B7) against *TgCDPK1*. This antibody recognizes the kinase in a conformation-dependent manner and selectively inhibits the Ca^{2+} -bound form. CocrySTALLIZATION of the antibody with *TgCDPK1* has yielded new insights into the activation of these enzymes and revealed a novel mechanism for CDPK inhibition.

Results

Removal of the Regulatory Domain of *TgCDPK1* Is Not Sufficient for Kinase Activation. By analogy to other Ca^{2+} -regulated kinases, it has been assumed that apicomplexan CDPKs are activated by derepression of the KD upon Ca^{2+} binding to the regulatory domain (CAD) (11, 12, 15). We reasoned that expression of *TgCDPK1* without the predicted repressive domains would yield a constitutively active kinase. We expressed two constructs that either removed the CAD completely (residues 1–314) or retained the autoinhibitory α -helix (residues 1–336). Like the full-length enzyme,

Significance

The activation of kinases by Ca^{2+} represents a vital class of signaling interactions that regulates many biological processes. The mechanism of activation for these enzymes is conserved and characterized by removal of an inhibitory element from the kinase domain. We report a previously unidentified mechanism for the activation of essential apicomplexan calcium-dependent protein kinases (CDPKs). Using *Toxoplasma* CDPK1 as a representative, we demonstrate that the kinase domain is intrinsically inactive and requires stabilization for activity. This distinct mechanism of activation reveals a susceptibility in CDPKs, which we exploit to effectively inhibit them. When viewed in the context of the entire protein kinase family, our results emphasize the remarkable adaptability of the kinase fold to diverse forms of regulation.

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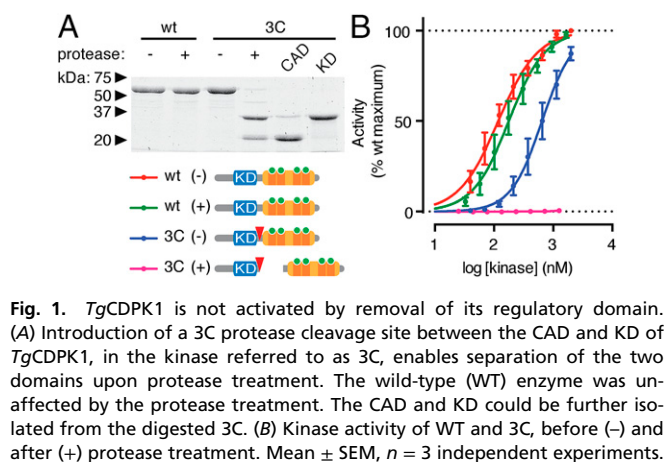
Data deposition: The atomic coordinates have been deposited in the Protein Data Bank, www.pdb.org (PDB ID code 4YGA).

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both constructs expressed well as soluble proteins, but neither showed measurable catalytic activity (Fig. S14).

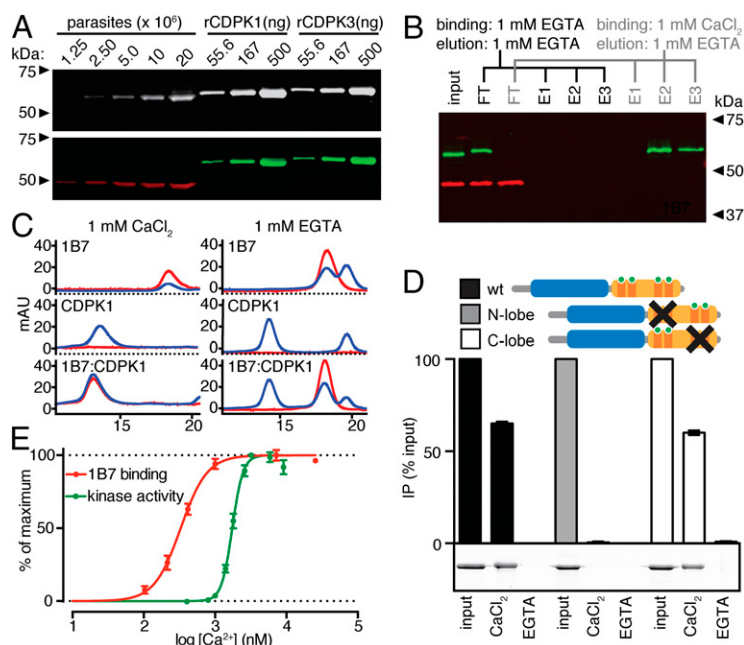
To examine the effect of severing the regulatory domain from the active kinase, we introduced the human rhinovirus 3C protease cleavage site at the junction of the KD and the CAD of *TgCDPK1* to generate the kinase referred to as “3C.” Treatment of the mutant enzyme with the protease separated the two domains of *TgCDPK1*-3C but did not cleave the wild-type enzyme (Fig. 1A). Although introduction of the protease site decreased activity somewhat, cleaving the connection between the CAD and the KD abolished kinase activity (Fig. 1B). Separating the KD from the CAD by size exclusion chromatography (SEC) or inclusion of a fivefold molar excess of the CAD failed to yield

appreciable kinase activity (Fig. S1B). We conclude that tethering the CAD to the KD is necessary for *TgCDPK1* activity.

Identification of Single Domain Antibodies to Probe *TgCDPK1* Structure.

Single domain antibodies derived from the heavy chain-only antibodies of alpacas (VHHs) are useful crystallization chaperones that often stabilize transient conformations (16). We identified VHHs that bind *TgCDPK1* from alpacas in our cohort that had antibodies against *T. gondii* proteins (Fig. S24). VHH phage display libraries from the seropositive animals were pooled and panned against recombinant *TgCDPK1*. VHH clones that showed reactivity against *TgCDPK1* by ELISA were enriched for the sequence represented by clone 1B7 (Fig. S2B). 1B7 was subcloned into a bacterial cytoplasmic expression vector, produced in *Escherichia coli*, and purified to homogeneity (Fig. S2C and D). By immunoblot, 1B7 recognized both endogenous and recombinant versions of *TgCDPK1* and its closest paralog *TgCDPK3* (Fig. 2A), with a limit of detection below 15 ng (Fig. 2A and Fig. S3). Measured against a standard curve of recombinant proteins, these kinases are expressed at ~ 4 –7 femtograms per cell, equivalent to 40,000–80,000 copies.

1B7 Dynamically Senses *TgCDPK1* Conformation. To determine whether 1B7 could sense Ca^{2+} -dependent conformational changes (11, 12), we performed immunoprecipitations (IPs) in the presence of either CaCl_2 or the cation chelator EGTA. 1B7 required Ca^{2+} to deplete lysates of its cognate antigens (Fig. 2B). Chelation of Ca^{2+} with EGTA readily eluted the bound CDPKs from the 1B7 beads, which could be reversed by further Ca^{2+} addition, demonstrating the reversibility of the interaction (Fig. S4A). To identify parasite proteins retrieved by 1B7, the eluted samples were analyzed by mass spectrometry. The two paralogs, *TgCDPK1* and *TgCDPK3*, were prominently represented with 26 unique



peptides each (52% and 41% sequence coverage, respectively) and were the only CDPKs found (Table S1). 1B7 thus specifically recognizes the Ca^{2+} -bound form of *TgCDPK1* and *TgCDPK3*.

We next determined whether 1B7 could form a stable complex with *TgCDPK1*. Using 1B7 labeled with a carboxytetramethylrhodamine (TAMRA) fluorophore at its C terminus, we tracked 1B7 binding to recombinant *TgCDPK1* by SEC. When preincubated with *TgCDPK1* in the presence of Ca^{2+} , 1B7-TAMRA coeluted with the kinase, well-resolved from 1B7-TAMRA alone (Fig. 2C). Thus, 1B7 binds a conformation of *TgCDPK1* that appears in the presence of Ca^{2+} and releases the kinase upon Ca^{2+} chelation.

1B7 Recognizes an Intermediate Conformation of *TgCDPK1*. To explore the conformational changes imposed by Ca^{2+} on *TgCDPK1* and their influence on 1B7 binding, we mutated the Ca^{2+} -binding EF hands of the kinase. The four EF hands of canonical CDPKs are segregated into two lobes, proposed to have differing Ca^{2+} affinities (10). To isolate the contribution of each lobe, we mutated the first invariant Asp in each EF hand loop in either lobe (N-lobe Asp368Ala/Asp415Ala and C-lobe Asp451Ala/Asp485Ala). Both the C-lobe mutant and wild-type kinase bound immobilized 1B7 in a Ca^{2+} -dependent manner with indistinguishable affinities, whereas binding was abolished by mutation of the N lobe (Fig. 2D). The N-lobe mutations also prevented 1B7 recognition by immunoblot (Fig. S4B). The N-lobe EF hands are thus crucial for 1B7 binding.

We next compared the precise $[\text{Ca}^{2+}]$ necessary for 1B7 binding and kinase activation. Using buffers with defined free $[\text{Ca}^{2+}]$, we measured the amount of kinase bound by 1B7 and, separately, the phosphorylation of a heterologous substrate. Binding of *TgCDPK1* by 1B7 occurred at a low free $[\text{Ca}^{2+}]$, with an EC_{50} of 300–350 nM (95% confidence interval), fivefold lower than that required for activation (Fig. 2E). The discrepancy between the $[\text{Ca}^{2+}]$ required for binding and that required for activation suggests that 1B7 binds a *TgCDPK1* conformation that is distinct from the active state of the kinase.

1B7 Is a Specific Inhibitor of *TgCDPK1* and Related Kinases. We wondered whether 1B7-bound CDPKs would retain activity in the presence of Ca^{2+} . We measured kinase activity in increasing concentrations of 1B7, or an irrelevant VHH. 1B7 inhibited both kinases—*TgCDPK1* and *TgCDPK3*—with an IC_{50} of ~40 nM and achieved complete inhibition at approximately equimolar concentrations of antibody and enzyme (Fig. 3A).

Although ATP γ S is a substrate of many parasite kinases, 6-Fu-ATP γ S is used exclusively by *TgCDPK1* in parasite lysates (17). Specificity is conferred by the unusually large ATP-binding pocket of *TgCDPK1* that harbors a Gly at a key position typically occupied by a bulky residue, which prevents other kinases—like *TgCDPK3*—from using 6-Fu-ATP γ S as a substrate. Thiophosphorylation can be detected following alkylation using the monoclonal antibody 51-8 (17). We measured thiophosphorylation in total cell extracts treated with increasing concentrations of 1B7. The presence of 1B7 robustly inhibited *TgCDPK1*-mediated thiophosphorylation but did not affect overall kinase activity (Fig. 3B).

We also probed lysates from erythrocytic stages of two different *P. falciparum* strains. A single band of the correct molecular weight for the homologs of *TgCDPK1* and *TgCDPK3* was detected in each lysate (Fig. 3C). Accordingly, 1B7 also inhibits recombinant *P. falciparum* CDPK1 (Fig. 3A). 1B7 is thus a potent and specific inhibitor of *TgCDPK1* and related apicomplexan kinases.

1B7 Interacts Directly with the Regulatory Domain of *TgCDPK1*. To understand the Ca^{2+} dependency of 1B7 binding and kinase inhibition, we obtained crystals of the *TgCDPK1*:1B7 complex that diffracted to 2.94 Å (Table 1). We solved the structure by molecular replacement (MR) and single anomalous dispersion (SAD) phasing using selenomethionine-derivatized *TgCDPK1*. The structure was refined to a final R_{work} of 22.6% and an R_{free} of 26.2%. We used

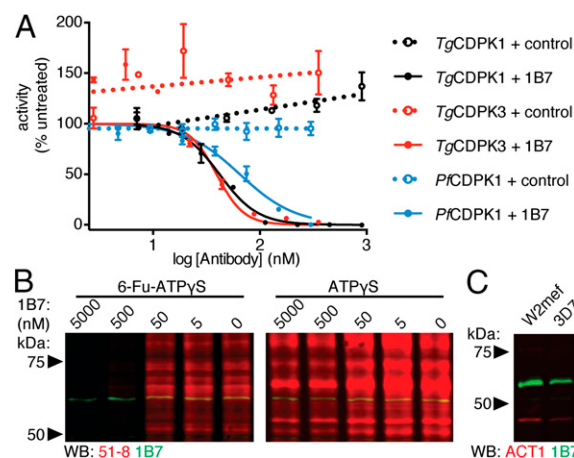


Fig. 3. 1B7 inhibits *TgCDPK1* and related kinases. (A) In vitro kinase assay with increasing concentrations of 1B7. Mean $\pm$ SEM, $n = 3$ independent experiments. (B) *TgCDPK1*-dependent (6-Fu-ATP γ S) and total (ATP γ S) thiophosphorylation in parasite lysates incubated with varying 1B7 concentrations. Thiophosphorylation is visualized by Western blot with rabbit mAb 51-8 (red); 1B7 staining is shown as a loading control (green). (C) Immunoblot of *P. falciparum* lysate from strain 3D7 or W2mef probed with 1B7-IRDye800 (green) or an antibody recognizing parasite actin (ACT1; red).

the structures of *TgCDPK1* previously solved in the Ca^{2+} -bound [Protein Data Bank (PDB) code 3HX4] and Ca^{2+} -depleted (PDB ID codes 3KU2 and 3V51) states (11, 12) for comparison.

The antibody engages the N lobe of the CAD (Fig. 4A and Fig. S5A), consistent with loss of antibody binding to the N-lobe mutant (Fig. 2D and Fig. S4B). All four EF hands are occupied by Ca^{2+} in the Ca^{2+} /1B7-bound structure (Fig. S5B). The Ca^{2+} dependency of the interaction is explained by 1B7's engagement of residues that participate in Ca^{2+} coordination and form the Ca^{2+} -binding regions of EF hands I and II. The Ca^{2+} -binding loop of EF hand II is ordered in the presence of 1B7, in contrast to the structures derived in the absence of Ca^{2+} (11, 12). Asn53, Asn55, and Ser58 of 1B7 CDR2 form a hydrogen-bonding network with the carbonyl oxygens of Gly371 and Asp372 in the loop segment of EF hand I. CDR3 of 1B7 contacts $\alpha 2$ of the N lobe via Thr100–Lys350 and Arg101–Ser349 hydrogen bonds and $\alpha 3$ (the first helix of EF hand I) via an Arg101–Thr361 hydrogen bond, a Glu102–His365 salt bridge, and a Phe103–Phe364 aromatic ring stack. CDR3 also contacts the loop segment and second helix of EF hand II ($\alpha 6$) through Tyr107, which forms a bidentate hydrogen bond to Asp417 and Glu426 on the CAD and a hydrogen bond between the amide of Ala106 and the Glu423 side chain (Fig. 4B; see Fig. S5 C–E for density maps). There are additional Van der Waals contacts that result in a total buried surface area of 790 Å² between 1B7 and *TgCDPK1*.

Having defined the interactions between *TgCDPK1* and 1B7, we explored the basis of 1B7's specificity. We generated a sequence alignment of known 1B7 binders (*TgCDPK1*, *TgCDPK3*, and *PfCDPK1*) together with CDPKs from *T. gondii*, *P. falciparum*, and *C. parvum*. Of the six residues involved in sequence-specific interactions, Lys350 and Thr361 are conserved in the 1B7-binding CDPKs and variable in the others (Fig. S6).

Allosteric Inhibition of *TgCDPK1* by 1B7. We aligned the three solved *TgCDPK1* structures by their KDs to determine the mechanism through which 1B7 inhibits the kinase. The CADs from the Ca^{2+} -depleted and Ca^{2+} /1B7-bound conformations occupy the opposite face of the KD from where the CAD lies in the Ca^{2+} -bound state (Fig. 5A). The first helix of the CAD (CH1) sits in the same cleft of the KD in the Ca^{2+} /1B7-bound and Ca^{2+} -depleted structures, although the helix terminates earlier

Table 1. Data collection and refinement statistics

Protein	TgCDPK1-1B7 native	TgCDPK1-1B7 SeMet derivative
Organism	<i>T. gondii</i>	<i>T. gondii</i>
PDB ID code	4YGA	
Data collection		
Space group	P1	P2 ₁
a, b, c, Å	76.92, 91.07, 106.64	77.61, 91.92, 107.90
$\alpha, \beta, \gamma, ^\circ$	88.7, 108.3, 90.3	90.0, 108.5, 90.0
Wavelength, Å	0.9792	0.9792
Resolution range, Å	73.04–2.94 (3.05–2.94)	102.3–4.09 (4.24–4.09)
Total reflections	215,441	126,898
Unique reflections	53,985	21,729
Completeness, %	93.5 (94.8)	98.2 (95.6)
Redundancy	3.9 (3.8)	5.8 (4.3)
Anomalous completeness, %		96.4
R _{sym} , %	11.0 (94.3)	30.1 (74.4)
R _{p.i.m.} , %	6.1 (53.8)	13.4 (37.7)
I/σ	10.3 (1.4)	6.0 (1.8)
CC _{1/2} , %	99.4 (69.6)	99.1 (75.0)
Refinement		
Resolution range, Å	70.67–2.94	
R _{work} , %	22.6	
R _{free} , %	26.2	
Coordinate error, Å	0.46	
Number of reflections		
Total	53,984	
R _{free} reflections	1,715	
Nonhydrogen atoms	17,348	
Protein atoms	17,332	
Calcium atoms	16	
rmsds		
Bond lengths, Å	0.003	
Bond angles, °	0.78	
Average B factors, Å ²		
Protein	113.3	
Calcium	84.4	
Ramachandran, %		
Favored, %	94.9	
Allowed, %	4.3	
Outlier, %	0.8	
Clashscore	9.05	
Molprobity percentile	97th	

Values in parentheses are for the highest-resolution shell.

in the $\text{Ca}^{2+}/1\text{B7}$ -bound structure. Superposition of the CAD domains shows that the $\text{Ca}^{2+}/1\text{B7}$ -bound CAD closely resembles the Ca^{2+} -bound conformer (Fig. 5B).

The Ca^{2+} -depleted conformation of TgCDPK1 has been proposed to be inhibited by an autoinhibitory triad between Lys338 in CH1 and the catalytic Glu135 and Asp138 and steric occlusion of the substrate binding site (11, 12). In contrast, Lys338 in our Ca^{2+} /1B7-bound structure participates in interactions within the CAD. We observed the formation of a disulfide bond linking Cys247 in the KD to Cys505 in the CAD in our Ca^{2+} /1B7-bound structure (Fig. S7A). Addition of a reducing agent did not affect 1B7 inhibition of kinase activity (Fig. S7B), although crystals were not obtained in the presence of 1 mM DTT. It is therefore likely that the Ca^{2+} /1B7-bound CAD is sampling many conformations in solution, including the state captured by the oxidized crystal structure, making steric occlusion of the substrate-binding pocket an unlikely mechanism for 1B7-mediated inhibition of TgCDPK1. The Ca^{2+} /1B7-bound kinase may be inactive because 1B7 and the KD bind to the same face of the CAD in the active conformation (Fig. 5C). This would imply an unprecedented role for the interaction of the CAD with the KD during activation.

Molecular Dynamics Show That the KD of TgCDPK1 Is Intrinsically Unstable. Long-timescale molecular dynamics (MD) simulations have been used to assess the intrinsic stability of various tyrosine kinases and inform their mechanisms of activation (18). To model the interaction of the CAD and the KD, we performed MD simulations using the Ca²⁺-bound conformation of TgCDPK1 as the starting point. Simulations performed in the absence of the CAD showed that the KD diverged from that of the Ca²⁺-bound, active structure (Fig. 6A). The major changes were in the rearrangement of the activation loop and in an approximate 30° turn in the αC helix of the KD to a so-called “αC-out” conformation that dissolves the catalytically important salt bridge between Lys80 and Glu99 and marks the inactive state of many protein kinases. We noted a clear preference for the inactive state when the CAD was removed from the KD (Fig. 6B and C).

The “regulatory” spine (R-spine) is a series of noncontiguous hydrophobic residues (Leu103, Leu114, His172, and Phe196 in TgCDPK1) that are aligned in the active conformations of all protein kinases (19). Misalignment of the R-spine residues is a key feature of the inactive states found in the MD simulation in the absence of the CAD and the $\text{Ca}^{2+}/1\text{B7}$ -bound structure

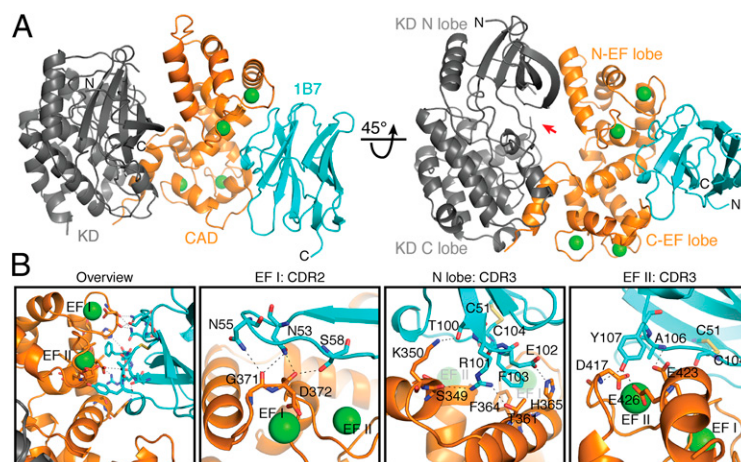


Fig. 4. 1B7 stabilizes a novel conformation of *TgCDPK1*. (A) Cartoon depicting the structure of the complex with the catalytic KD (gray), the CaM-like CAD (orange), bound calcium ions (green spheres), and 1B7 (blue). The position of the active site pocket is indicated (red arrow). (B) Specific interactions within the CAD:1B7 binding interface. Hydrogen bonds and salt bridges are depicted as black dotted lines. Interacting residues are shown as sticks for the CAD and 1B7. Cys51, in the beta strand immediately preceding CDR2, and Cys104, in CDR3, form a stabilizing, intrachain disulfide in 1B7. CDR1 on 1B7 makes no contacts with *TgCDPK1*.

(Fig. 6D). The R-spine was similarly disrupted in the Ca^{2+} -depleted structure, in contrast to its perfect alignment in the Ca^{2+} -bound conformation (Fig. 6D). The Ca^{2+} -bound CAD appears to impart an active conformation on the KD by stabilizing the position of the αC helix required for alignment of the R-spine and catalysis.

The CAD Acts as a Molecular Splint to Stabilize the KD of *TgCDPK1*.

During activation, the CAD imposes a 15° rotation in the relative orientations of the N and C lobes of the KD, suggesting that points of attachment to both kinase lobes are needed to generate torque upon Ca^{2+} binding. Having established that integrity of the tether between the KD and the CAD is necessary (Fig. 1B), we investigated whether the flexibility or length of the tether would affect kinase activity. We replaced nine residues from the tether with a flexible linker of equal length (9AA) or one 11 residues longer (20AA). We observed a loss of kinase activity proportional to the

length and presumed flexibility of the tether (Fig. 6E). The greater loop entropy associated with the longer linkers presumably disfavors proper positioning of CAD and KD (20).

Interaction between the CAD and an N-terminal extension of the KD, also visible in the Ca^{2+} -bound structure (11), could provide the proposed tether to the N lobe of the KD. Superposition of the Ca^{2+} -bound and Ca^{2+} /1B7-bound structures showed that the same groove in the Ca^{2+} -bound CAD was engaged by both the N-terminal extension and CDR3 of 1B7 (Fig. S8). We therefore asked whether the N-terminal extension of *TgCDPK1* participates in kinase activation. Of the 50 residues in the N-terminal extension, only the final 20 are observed in the Ca^{2+} -bound structure, and only the final six in both inactive structures. Although truncation of the first 35 residues minimally affected kinase activity, elimination of 44 residues from the N-terminal extension led to a complete loss of function (Fig. 6F).

We identified two key residues important for this interaction: Phe39 and Val40. We mutated Phe39 and Val40 individually and in combination to Ala and measured their effects. Val40Ala shifted the EC_{50} of the kinase by ~ 10 -fold, whereas Phe39Ala strongly inhibited kinase activity, as did the double mutation (Fig. 6G). The N-terminal extension is therefore crucial for the activation of *TgCDPK1*.

Allosteric Activation Is Required for *TgCDPK1* Function in Vivo.

To examine the role of the N-terminal extension in vivo, we used a conditional knockout (cKO) of *TgCDPK1* (2). This strain uses a tetracycline transactivator to express an HA9-tagged allele of *TgCDPK1*, in the absence of the endogenous gene (Fig. 7A). Growth of the cKO strain in the presence of anhydrotetracycline (ATc) blocks expression of *TgCDPK1*, which was used to demonstrate the essentiality of the kinase in the *T. gondii* lytic cycle (2). The cKO can be complemented with an allele of the kinase constitutively expressed under the endogenous promoter of *TgCDPK1* (Fig. 7A). We generated strains complemented either with wild type or with the Phe39Ala mutant incapable of allosteric activation. We confirmed regulation of the HA9-tagged allele and constitutive expression of the complementing, Myc-tagged alleles by immunoblot (Fig. 7B). To determine whether the expressed enzymes were active, we performed thiophosphorylation reactions, using 6-Fu-ATP γS as a substrate, with lysates from parasites grown for 48 h in the presence of ATc. We observed robust Ca^{2+} -dependent thiophosphorylation in the wild-type

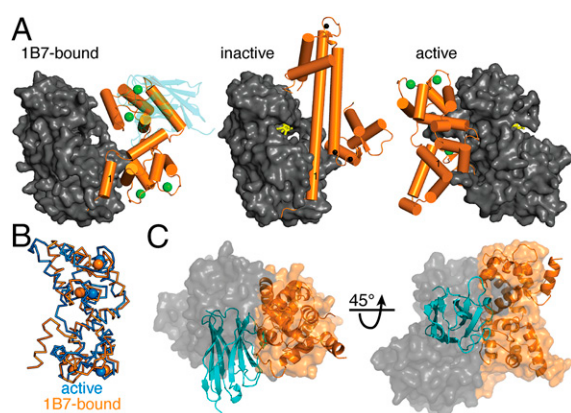


Fig. 5. 1B7 prevents reorganization of the CAD to its activation-associated conformation. (A) Comparison of 1B7-bound *TgCDPK1* to the inactive (3KU2) and active (3HX4) conformations. The three *TgCDPK1* conformations are aligned by their KDs (gray), depicting the CAD (orange), 1B7 (blue), bound calcium (green), and unknown cations (black). ANP (phosphoaminophosphonic acid-adenylate ester) is built as yellow sticks to indicate the active site. The first helix of the CAD (CH1) is labeled with "1." (B) Aligned carbon trace of the CADs from the active (blue) and 1B7-bound structures (orange). (C) Superposition of 1B7 and the bound CAD as ribbon diagrams over a space-filling model of the active structure.

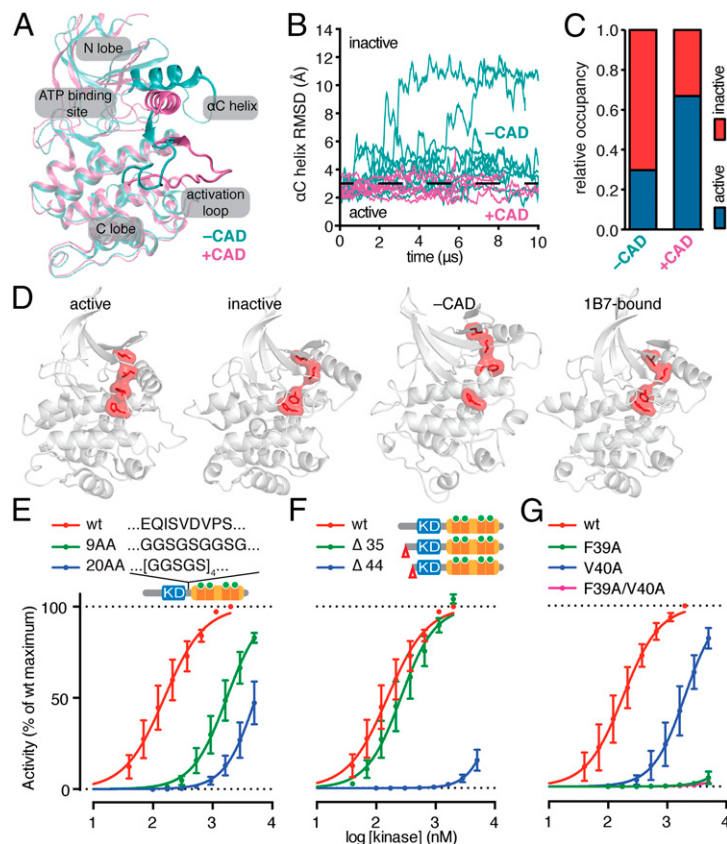


Fig. 6. The KD of *TgCDPK1* is intrinsically inactive and stabilized by the Ca^{2+} -bound CAD. (A) Superposition of the active conformation of the *TgCDPK1* KD with an inactive conformation obtained from a simulation without the CAD, in which the KD readily departed from the (starting) active conformation. The superposition is obtained using the C-lobe backbone atoms. (B) The rmsd trace of the αC helix relative to the (starting) active conformation in simulations of the *TgCDPK1* KD with or without the CAD. Conformations with an rmsd less than 3 Å are considered active. (C) The relative occupancy of the catalytically active conformation in simulations of the *TgCDPK1* KD with or without the CAD. As shown, the presence of the CAD stabilizes the active conformation. (D) Structure of the KD highlighting the integrity of the R-spine (red) in the Ca^{2+} -bound (active) conformation, and its disruption in the Ca^{2+} -depleted (inactive), the simulation without the CAD (-CAD), and the Ca^{2+} /1B7-bound (1B7-bound) conformations. (E) Kinase activity of *TgCDPK1* mutants carrying different flexible linkers between the KD and CAD domains. (F) Kinase activity of different N-terminal truncations removing the first 35 ($\Delta 35$) or 44 ($\Delta 44$) amino acids preceding the KD. (G) Kinase activity of point mutants in the N-terminal extension, alone or in combination. All kinase experiments plotted as means $\pm$ SEM, $n = 3$ independent experiments.

complemented strain. However, consistent with our analysis of the recombinant enzymes, no *TgCDPK1* activity was observed in the Phe39Ala mutant strain (Fig. 7C).

We next measured plaque formation to assess functional complementation of the cKO. All strains formed plaques of similar size when grown in the absence of ATc (Fig. 7D). However, plaque formation was impaired when the cKO was grown in the presence of ATc (2). This defect was complemented by constitutive expression of the wild-type allele, but not the Phe39Ala mutant (Fig. 7D). Taken together, these data demonstrate that the Phe39 residue of *TgCDPK1*, which mediates allosteric activation, is necessary for the function of the kinase in vivo.

Discussion

Unlike many other Ca^{2+} -regulated kinases, the KD of *T. gondii* CDPK1 is intrinsically inactive. We identified an alpaca-derived single domain antibody (1B7) that binds to and inhibits *TgCDPK1* in a reversible, Ca^{2+} -dependent manner. Solving the structure of *TgCDPK1* in complex with 1B7 showed a common binding site on the Ca^{2+} -bound CAD for both the antibody and the *TgCDPK1* N-terminal extension. 1B7 binding prevents the stabilization of the KD by blocking the interaction of the N-terminal extension and the CAD. We refer to this mode of stabilization as a molecular splint because anchoring of the KD to the CAD at both the N and C termini is required for activation.

Ca^{2+} -regulated kinases are activated by removal of an inhibitory element from the KD (7, 8, 13, 14). Animal Ca^{2+} /CaMKs possess intrinsically active KDs, kept inactive by an autoinhibitory region (13, 14). A similar autoinhibitory region is part of the CAD of CDPKs (11). Peptide mimetics of this region inhibit both plant (7, 8) and parasite kinases (6, 21). Based on observations of plant CDPK truncations that appeared constitutively active and Ca^{2+} -independent, CDPK KDs—like those of CaMKs—were thought to be intrinsically active (7, 21). These truncations consisted of N-terminal fusions with GST, which can dimerize and stabilize the conformation of its fusion partners. For phosphatidylinositol 3'-kinase and the insulin receptor KD, fusion to GST is sufficient for activation (22, 23). In contrast, the KD of *TgCDPK1* is inactive on its own. Removal of the CAD, either genetically or proteolytically, fails to activate *TgCDPK1*. MD simulations showed that the KD adopts an inactive conformation in the absence of the CAD. The KDs of different CDPKs might well differ in stability, but our work suggests that instability is a conserved feature of the CDPKs most closely related to *TgCDPK1*.

Ca^{2+} binding to the CAD imposes rigidity to its fold. When connected to the N and C termini of the KD, the CAD acts as a molecular splint that supports and stabilizes the KD in its active conformation (Fig. 7E). The CAD therefore exerts a far greater degree of regulation over the KD than assumed previously. Perturbation of either point of contact between the CAD and

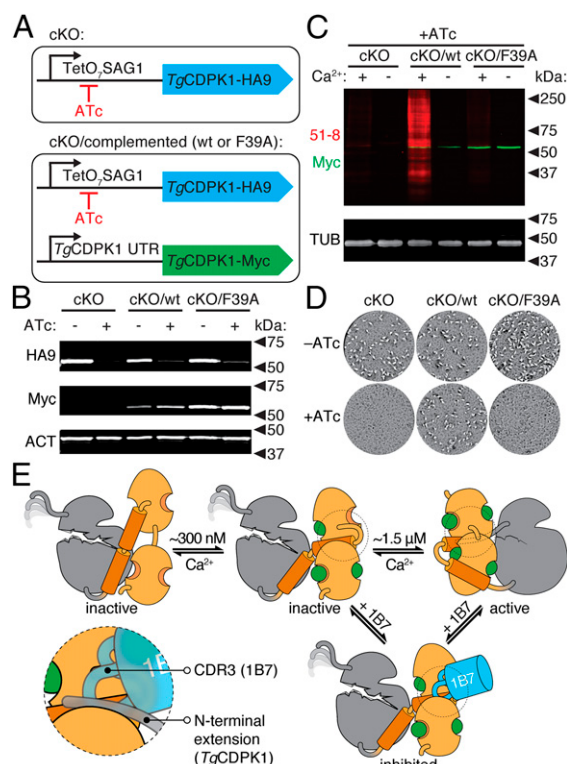


Fig. 7. Allosteric activation is required for *TgCDPK1* activity in vivo. (A) Strategy depicting the *TgCDPK1* cKO carrying an HA9-tagged allele that can be shut down by addition of ATc. The complemented strains additionally carry a constitutive Myc-tagged allele (WT or F39A), expressed under the endogenous *TgCDPK1* promoter. (B) Immunoblot of regulatable (HA9) or constitutive (Myc) *TgCDPK1* alleles in the different strains grown in the presence or absence of ATc for 48 h. Parasite actin (ACT) is included as a loading control. (C) *TgCDPK1*-dependent thiophosphorylation (51-8; red) in lysates from the various parasite strains following growth in the presence of ATc. Reactions were performed in the presence or absence of Ca²⁺. Tubulin is included as a loading control. (D) Plaque formation by the various strains in the presence or absence of ATc. (E) Model for the activation of *TgCDPK1* and its inhibition by 1B7. The KD (gray) is intrinsically inactive and its catalytic site is occluded by the CAD (orange). At ~300 nM Ca²⁺, the CAD is partially occupied by Ca²⁺ and can bind 1B7 (blue). Higher Ca²⁺ concentrations are required for full Ca²⁺ occupancy of the CAD and stabilization of the KD, which leads to activation. The region of the CAD competitively bound by 1B7 and the N-terminal extension is circled and highlighted in the *Inset*.

KD impairs kinase activity. Disruption of the N-terminal extension (Fig. 6 F and G) strongly inhibited kinase activity, congruent with studies of *PfCDPK1* (24). We identified Phe39 and Val40 in the N-terminal extension as crucial for kinase activation. The presence of Phe39 is required for *TgCDPK1* function in vivo, because the Phe39Ala mutant, when expressed at wild-type levels, fails to complement a cKO of the kinase (Fig. 7). The similarity of these anchor residues among diverse CDPKs suggests that the corresponding regions may function like the N-terminal extension of *TgCDPK1*.

KDs have evolved diverse modes of positive and negative regulation, often through acquisition of additional protein domains or noncovalent interacting partners. The diversity of these regulatory systems often converges on similar themes, including regulation of the α C helix position via a hydrophobic patch in the N lobe of the KD (25). Regulatory elements can interact directly with this hydrophobic patch, as is the case with the nonreceptor tyrosine kinases Fes and Abl (26). In the case of Abl, the same regulatory domain can also act to lock the kinase in an inactive conformation (27). Only a few domains are known to exert dual

functions in the regulation of kinase activity, by serving both as repressors and activators. In a remarkable convergence, the CAD has adopted such a role in the regulation of *TgCDPK1*.

The interaction of 1B7 with its target is characteristic of VHHs, which often present large convex paratopes that can access buried epitopes less accessible to conventional antibodies (28). The 1B7 CDR3 protrudes into a groove that forms upon Ca²⁺ binding to the CAD, displacing the N-terminal extension in the active conformation of the kinase (11). Features associated with autoinhibition are notably absent from the Ca²⁺/1B7-bound structure. 1B7 may bind an intermediate state in the activation of *TgCDPK1* as inferred from the lower [Ca²⁺] required for 1B7 binding, versus kinase activation (Fig. 2E). Preventing the stabilization of the KD is the primary mechanism for the inhibition of *TgCDPK1* by 1B7, a trait generalizable at least to the clade of CDPKs to which *TgCDPK1* belongs.

CDPKs are obvious therapeutic targets. Development of small molecule inhibitors against them has focused exclusively on ATP analogs, which often suffer from off-target effects against other host and parasite kinases. Targeting the atypical ATP-binding pocket of *TgCDPK1* and related kinases has afforded some measure of specificity. However, a single mutation suffices to confer resistance to these compounds (3). We identify a vulnerability in this family of parasite kinases, exposed by their need for stabilization. Through the use of 1B7, we provide proof of principle that this feature can be exploited to inhibit *TgCDPK1* and related kinases, including *PfCDPK1*, a kinase essential for the erythrocytic stages of the malaria parasite (6). Conservation of the residues that mediate allosteric activation suggests that this mechanism may extend to *Cryptosporidium* CDPK1 and *Plasmodium* CDPK4. Inhibition of these kinases has already shown promising activity in murine infection models, in the case of *CpCDPK1* (29), and in blocking the transmission of malaria, in the case of *PbCDPK4* (30). We expect that the binding of 1B7 to *TgCDPK1* can be used to guide the design of small molecules with the therapeutic potential to allosterically inhibit these essential parasite kinases.

Methods

Protein Purification. Vectors encoding *TgCDPK1*-6xHis and *TgCDPK3*-6xHis were generated as previously described (3). Mutations in the EF hands or the N-terminal extension were introduced using the QuikChange II Site-Directed Mutagenesis Kit (Agilent Technologies), with specific primers designed according to the manufacturer's instructions. N-terminally truncated versions of *TgCDPK1* were generated using primers P1 and P2 (Δ 35) or P1 and P3 (Δ 44). Mutant kinases with insertions in the junction between KD and CAD were ordered as gBlocks from Integrated DNA Technologies (B1 for 9AA, B2 for 20AA, and B3 for 3C) and cloned into pET22b+ (Novagen). Primer and gBlock sequences are listed in Table S2.

Kinase constructs were transformed into BL21(DE3)V2RpAcYc-LIC+Lamp *E. coli*, which express the Lamp phosphatase as previously described (11). Cultures were grown to 0.6 OD₆₀₀ at 37 °C and induced with 1 mM isopropyl- β -D-thiogalactopyranoside (IPTG) for 2.5 h. Pelleted cells were lysed in Cel-Lytic B buffer (Sigma) containing 0.1 mg·mL⁻¹ lysozyme (Sigma), 650 units benzonase (Sigma), and HALT Protease Inhibitor Mixture (Thermo-Scientific). Lysates were clarified by centrifugation at 24,000 $\times$ g for 30 min at 4 °C. The soluble fraction was added to prewashed Ni-NTA agarose (Qiagen) and allowed to bind 1 h at 4 °C, rotating. The resin was washed with 50 mM sodium phosphate pH 8.0, 500 mM NaCl, 15 mM imidazole, and HALT Protease Inhibitor Mixture, and bound proteins were eluted by increasing imidazole to 500 mM. Eluates were concentrated using Amicon Centrifugal Filters (EMD Millipore) and loaded onto a Superdex 200 26/60 SEC column equilibrated with 50 mM Tris, pH 7.5, and 150 mM NaCl. Peak elution fractions were pooled and concentrated. Protein purity was assessed by SDS/PAGE, and concentrations were determined using the DC Protein Assay (Bio-Rad). Recombinant CDPKs were stored at -80 °C after addition of 25% (vol/vol) glycerol and 1 mM DTT.

For crystallization, the following changes were made. Proteins were expressed in BL21(DE3)-LOBSTR-RIL *E. coli*, which has been optimized for low background in Ni-affinity purifications (31). Cultures were shifted to 18 °C before induction with 0.2 mM IPTG for 16 h. Cells were lysed using a cell disruptor (Constant Systems) in 50 mM potassium phosphate, pH 8.0, 500 mM NaCl,

30 mM imidazole, 3 mM β -mercaptoethanol, 1 mM PMSF, and 250 units of Turbonuclease (Eton Bioscience). The soluble fraction was added to pre-washed Ni-Sepharose fast-flow beads (GE Healthcare) and agitated 20 min at 4 °C. Protein was eluted with 250 mM imidazole, pH 8.0, 150 mM NaCl, and 3 mM β -mercaptoethanol. SEC was performed in 10 mM Tris, pH 7.4, 150 mM NaCl, 1 mM DTT, and 1 mM CaCl_2 . SeMet-derivatized TgCDPK1 was purified identically, with the exception that 10 mM β -mercaptoethanol and 5 mM DTT were used in purification steps.

Upon VHH selection by phage display and ELISA, the 1B7 coding sequence was subcloned for expression. First, VHH coding sequences were amplified with primers P4 and P5, which include a sortase A recognition motif. The resulting product was subcloned into the pET30b+ (Novagen) expression vector using NdeI and XhoI sites for inclusion of a C-terminal 6xHis tag. BL21 (DE3) *E. coli* containing the plasmid were grown to midlog phase at 37 °C in 2YT plus kanamycin and induced with 1 mM IPTG overnight at 30 °C. Total soluble protein was collected by French press and cleared at 39,000 $\times$ g for 30 min, before loading onto Ni-NTA (Qiagen) in 50 mM Tris, pH 7.5, 150 mM NaCl, and 10 mM imidazole. Protein was eluted in 50 mM Tris, pH 7.5, 150 mM NaCl, 500 mM imidazole, and 10% (vol/vol) glycerol, and then loaded onto a Superdex 75 10/300 column in 50 mM Tris, pH 7.5, 150 mM NaCl, and 10% (vol/vol) glycerol. Peak fractions were pooled and concentrated. Purity was assessed by SDS/PAGE.

The TgCDPK1 variant carrying a human rhinovirus 3C protease cleavage site was digested with 3C protease for 36 h at 4 °C to achieve near-complete digestion. Wild-type kinase underwent the same treatment, to be used as a control. KD and CAD were separated from digested samples by SEC through a Superdex 75 16/60 column equilibrated with 50 mM Tris, pH 7.5, 150 mM NaCl, 1 mM CaCl_2 , and 1 mM DTT. Fractions for each peak were pooled and concentrated as described above.

Growth of Host Cells and Parasite Strains. *T. gondii* tachyzoites were maintained by growth in monolayers of human foreskin fibroblasts cultured in DMEM containing 10% (vol/vol) tetracycline-free FBS (HyClone), 2 mM glutamine, 10 mM Hepes pH 7.5, and 20 $\mu\text{g}\cdot\text{mL}^{-1}$ gentamicin, as described (2). The TgCDPK1 cKO was generated as previously described (2). The wild-type complementation plasmid was previously generated and contains a Myc-tagged version of TgCDPK1 downstream of the endogenous promoter sequence of the kinase (2). Phe39 was mutated to Ala in this construct using the QuikChange II Site-Directed Mutagenesis Kit (Agilent Technologies), according to the manufacturer's instructions. Mutagenesis and integrity of the entire ORF was confirmed by sequencing. The complementing plasmids were cotransfected with pDHFR-TS (32), and stable integration was selected for by growth in 3 μM pyrimethamine (Sigma). Clones were isolated by limiting dilution, and expression of the transgenes was confirmed by immunoblot. We added 1.5 $\mu\text{g}\cdot\text{mL}^{-1}$ ATc (Clontech) as indicated to suppress expression of the regulatable, HA9-tagged wild-type allele.

Plaque Assays. Assays were performed as previously described (2). Confluent monolayers of human foreskin fibroblasts in six-well plates were infected with 400–1,000 parasites per well in media with or without 1.5 $\mu\text{g}\cdot\text{mL}^{-1}$ ATc. At 24 h after infection, additional medium was added to decrease the concentration of ATc to 1 $\mu\text{g}\cdot\text{mL}^{-1}$. Monolayers were fixed 7 d after infection and stained with crystal violet.

Immunoblots. Parasites from fully lysed monolayers were lysed directly in SDS/PAGE loading buffer, or in a 50 mM Tris, pH 7.5, 150 mM NaCl, 1% Triton X-100, and HALT Protease Inhibitor Mixture, before measuring protein concentration. Serum was collected from each alpaca before immunization and stored at –80 °C (33). *T. gondii* lysates were run on SDS/PAGE and transferred to nitrocellulose. Membranes were blocked overnight at 4 °C in PBS, 5% (wt/vol) nonfat dry milk. Serum was diluted 1:5,000 in blocking solution and incubated 1 h with the respective section of the blot. Horseradish peroxidase-conjugated goat anti-llama IgG (Bethyl) was used as a secondary antibody at a 1:20,000 dilution in blocking solution. To compare *T. gondii* lysates with recombinant proteins, lysates corresponding to different numbers of parasites were loaded besides dilutions of the recombinant proteins. 1B7 was labeled using the LI-COR IRDye 800CW labeling kit according to the manufacturer's instructions and used at a concentration of 1 $\mu\text{g}\cdot\text{mL}^{-1}$ for immunoblots. Expression of the regulatable and complementing alleles of TgCDPK1 was similarly assessed by immunoblot following growth of the parasites in the presence or absence of ATc. Immunoblots were probed, as indicated, with Rabbit anti-TgACT1 (34) diluted 1:10,000, mouse anti-penta HIS (Qiagen) diluted 1:5,000, rabbit anti-HA9 (Invitrogen) diluted 1:10,000, mouse anti-Myc (9E10; Sigma) diluted 1:5,000, and mouse anti- α Tubulin (12G10; Developmental Studies Hybridoma Bank) diluted 1:10,000.

VHH Library Generation. Alpacas (*V. pacos*) were purchased locally, maintained in pasture, and immunized following a protocol authorized by the Tufts University Cummings Veterinary School Institutional Animal Care and Use Committee. For each alpaca, total RNA was isolated from $\sim 10^6$ fresh peripheral blood lymphocytes using the RNeasy Plus Mini Kit (Qiagen), following the manufacturer's instructions. First strand cDNA synthesis was performed using SuperScript III reverse transcriptase (Life Technologies) and a combination of oligo dT, random hexamer, or Ig-specific primers, ALCH2 and ALCH2.2, as previously described (33). Subsequent PCR amplification of VHH sequences and phage library generation followed an established procedure (33). This included the use of alpaca-specific primers for VHH gene amplification and the use of a phagemid vector adapted for VHH expression as a pIII fusion. Following transformation into *E. coli* TG1 cells (Agilent), the total number of independent clones for each library was estimated to range from 1×10^6 – 10^7 . The resulting phagemid libraries were stored at –80 °C.

Generation of M13 Phage Displaying Anti-*T. gondii* VHH Library. Several independent libraries were generated from animals immunized with different mixtures of antigens. None of the animals were deliberately immunized with *T. gondii*. Libraries were pooled and used to inoculate 100 mL SOC Medium with 50 $\mu\text{g}\cdot\text{mL}^{-1}$ ampicillin. The culture was grown to midlog phase and infected with 100 μL 10^{14} pfu $\cdot\text{mL}^{-1}$ VCSM13 helper phage. Following a 2-h incubation at 37 °C, the cells were harvested by centrifugation and resuspended in 100 mL 2YT, 0.1% glucose, 50 $\mu\text{g}\cdot\text{mL}^{-1}$ kanamycin, and 50 $\mu\text{g}\cdot\text{mL}^{-1}$ ampicillin. Cultures were incubated overnight at 30 °C, then centrifuged for 20 min at 7,700 $\times$ g, followed by phage precipitation from the resulting supernatant with 1% PEG-6000, 500 mM NaCl at 4 °C, and resuspension in PBS.

Selection of VHHS by Phage Display. VHHS were selected by panning against recombinant TgCDPK1. We biotinylated 100 μg recombinant TgCDPK1 by coupling Chromalink NHS-biotin reagent (Solulink) to primary amines for 90 min in 100 mM sodium phosphate, pH 7.4, 150 mM NaCl. Unreacted biotin was removed using a 10 kDa molecular weight cut-off concentrator (Millipore). Biotin incorporation was monitored, following the manufacturer's guidelines. We blocked 100 μL MyOne Streptavidin-T1 Dynabeads (Life Technologies) in PBS, 2% (wt/vol) BSA (Sigma) for 2 h at 37 °C. We added 20 μg biotinylated TgCDPK1 in PBS to the blocked beads and incubated it for 30 min at 25 °C, with agitation. The beads were then washed three times in PBS, and 200 μL of 10^{14} pfu $\cdot\text{mL}^{-1}$ M13 phage displaying the VHH library were added in PBS, 2% (wt/vol) BSA for 1 h at room temperature. The beads were then washed 15 times with PBS, 0.1% Tween-20. Phage was eluted by the addition of *E. coli* ER2738 (NEB) for 15 min at 37 °C, followed by elution with 200 mM glycine, pH 2.2, for 10 min at 25 °C. The eluate was neutralized; pooled with the *E. coli* culture; plated onto 2YT agar plates supplemented with 2% (wt/vol) glucose, 5 $\mu\text{g}\cdot\text{mL}^{-1}$ tetracycline, and 10 $\mu\text{g}\cdot\text{mL}^{-1}$ ampicillin; and grown overnight at 37 °C. A second round of panning was performed with the following modifications: 2 μg of biotinylated TgCDPK1 was used as bait and incubated with 2 μL 10^{14} pfu/mL M13 phage displaying the first-round VHH library for 15 min at 37 °C, followed by extended washes in PBS, 0.1% Tween-20.

ELISA. Following two rounds of phage panning, 96 colonies were isolated in 96-well round-bottom plates and grown to midlog phase at 37 °C in 200 μL 2YT, 10 $\mu\text{g}\cdot\text{mL}^{-1}$ ampicillin, and 5 $\mu\text{g}\cdot\text{mL}^{-1}$ tetracycline, induced with 3 mM IPTG and grown overnight at 30 °C. Plates were centrifuged at 12,000 $\times$ g for 10 min, and 100 μL of supernatant was mixed with an equal volume of PBS, 5% (wt/vol) nonfat dry milk. This mixture was added to an ELISA plate coated with 1 $\mu\text{g}\cdot\text{mL}^{-1}$ TgCDPK1. Following four washes in PBS, 1% Tween-20, anti-llama-HRP antibody (Bethyl) was added at a 1:10,000 dilution in PBS, 5% (wt/vol) nonfat dry milk for 1 h at 25 °C. The plate was developed with fast kinetic 3,3',5,5'-tetramethylbenzidine (Sigma) and quenched with 1 M HCl. Absorbance at 450 nm was determined in a plate reader (Spectramax; Molecular Devices).

IP of TgCDPK1 and TgCDPK3. We coupled 4 mg of 1B7 to 500 mg of cyanogen bromide-activated Sepharose (Sigma) according to the manufacturer's instructions. Coupling efficiency was monitored by SDS/PAGE. 1B7-Sepharose was stored in 50 mM Tris, pH 7.5, 150 mM NaCl at 4 °C. Approximately 10^9 parasites were harvested, resuspended in lysis buffer [50 mM Hepes, pH 7.4, 100 mM KCl, 1 mM MgCl_2 , 10% (vol/vol) glycerol, and 0.05% Triton X-100], and disrupted with a Dounce homogenizer. We prepared 0.25 g 1B7-Sepharose for each sample by washing with lysis buffer supplemented with either 1 mM EGTA or CaCl_2 . Half of the parasite lysate was diluted to 10 mL containing 1 mM EGTA or CaCl_2 and incubated with the respective beads for 1 h at 4 °C, with agitation.

The resin was then washed with 200 mL of the appropriate lysis buffer supplemented with EGTA or CaCl_2 and eluted with three consecutive 500 μL volumes of lysis buffer containing 1 mM EGTA. The eluted material was analyzed by SDS/PAGE followed by silver staining. Bands were excised, reduced, alkylated, and digested with trypsin at 37 °C overnight. The resulting peptides were extracted, concentrated, and injected by standard reverse-phase chromatography into a Thermo Orbitrap Elite mass spectrometer operated in a data-dependent manner. The resulting fragmentation spectra were correlated against the known database using SEQUEST. Scaffold Q+S (Proteome Software) was used to provide consensus reports for the identified proteins.

The IP of recombinant, wild-type, and double EF hand mutant kinases was performed similarly. We diluted 5 μg of kinase in 1 mL of the respective binding buffer (50 mM Tris pH 7.5, 150 mM NaCl, 1% Triton X-100, supplemented with either 1 mM EGTA or 1 mM CaCl_2 , respectively) for 1 h. For each binding condition, 1B7-Sepharose beads were washed twice with the respective binding buffer, followed by incubation of ~ 100 μg 1B7-Sepharose with 1 mL of diluted kinase for 30 min, with agitation. Following incubation, the beads were loaded onto 200 μL filter tips and washed with three column volumes of the respective buffer, before eluting with 40 μL 200 mM glycine, pH 2.2, followed by neutralization with 10 μL 1 M Tris, pH 9.0. The amount of immunoprecipitated kinase was determined by SDS/PAGE, SYPRO Ruby protein staining (Life Technologies), and fluorescent gel imaging using a Typhoon FLA 9500 laser scanner (GE Healthcare Life Sciences). Densitometry was conducted using ImageQuant TL software (GE Healthcare Life Sciences).

The IP of wild-type TgCDPK1 in the presence of specific $[\text{Ca}^{2+}]_{\text{free}}$ concentrations was performed correspondingly, but with distinct buffer compositions. These binding buffers were made using the Calcium Calibration Buffer Kit (Life Technologies), adjusting the supplied buffer composition to 30 mM Mops, pH 7.2, 100 mM KCl, 4 mM CaEGTA/ K_2EGTA , 1 mM MgCl_2 , and 0.05% Tween-20 to more closely mimic physiological conditions and to assure consistency between the conditions used to measure kinase activity and 1B7 binding. The $[\text{Ca}^{2+}]_{\text{free}}$ were determined using WEBMAX STANDARD (35). Before IP, wild-type TgCDPK1 was dialyzed against chelex-100 (Bio Rad)-treated buffer, to minimize contaminating the Ca^{2+} present in the buffers.

SEC. Sortase was used to label 1B7 by incubating 1 mg of purified 1B7 with 1 mg *Staphylococcus aureus* sortase A and 1 mM GGGK-TAMRA in 50 mM Tris, pH 7.5, 150 mM NaCl, and 10 mM CaCl_2 . The reaction proceeded overnight at 25 °C, and unreacted VHH and sortase A were removed by adsorption onto Ni-NTA agarose (Qiagen). Labeled 1B7 was isolated by SEC, concentrated, and its purity determined as described above. TgCDPK1 and purified TAMRA-labeled 1B7 were incubated at a 1:1 molar ratio in 50 mM Tris, pH 7.5, and 150 mM NaCl, supplemented with 1 mM CaCl_2 or 1 mM EGTA for 2 h at 25 °C. Samples were then loaded onto a Superdex 75 10/30 column in 50 mM Tris, pH 7.5, and 150 mM NaCl, supplemented with 1 mM CaCl_2 or 1 mM EGTA, and absorbance was monitored at both 280 nm and 550 nm.

Kinase Assays. Kinase assays were conducted using a peptide-based ELISA, as previously described (3). We coated 96-well plates with Syntide-2 peptide (Calbiochem) diluted to 100 $\mu\text{g}\cdot\text{mL}^{-1}$ in carbonate coating buffer (pH 9.6) at 4 °C overnight. Following incubation, plates were rinsed with wash buffer (50 mM Tris, pH 7.5, and 0.2% Tween-20) and blocked with 3% (wt/vol) BSA in wash buffer for 2 h at 25 °C. Subsequent washes were performed with wash buffer. Reactions were conducted at 30 °C for 30 min in kinase buffer (20 mM Hepes, pH 7.0, 10 mM MgCl_2 , 1 mM CaCl_2 , 0.005% Tween-20, and 1 mM ATP). Phosphorylation was detected with mAb MS-6E6 (MBL), followed by HRP-conjugated goat-anti-mouse IgG, developed with the substrate o-phenylenediamine dihydrochloride (Thermo Scientific), and detected by absorbance at 492 nm. Where appropriate, each kinase preparation was individually tested to determine its half maximal effective concentration (EC_{50}). For VHH inhibition assays and Ca^{2+} titration experiments, TgCDPK1 was used at its EC_{50} . For the inhibition assays, 1B7 or a control VHH (2A7) was titrated into the reaction, before the addition of 1 mM ATP. Ca^{2+} titration experiments used reaction buffers with specific $[\text{Ca}^{2+}]_{\text{free}}$ prepared as in the IP experiments, with 1 mM ATP added. EC_{50} and IC_{50} values were calculated from a dose-response curve plotted in Prism (GraphPad). Duplicate or triplicate experiments were run for all assays.

Thiophosphorylation Assay. Thiophosphorylation was measured as previously described (17). Briefly, *T. gondii* parasites were lysed on ice in the reaction buffer (20 mM Hepes, pH 7.4, 137 mM KCl, and 10 mM MgCl_2), supplemented with 1% Triton X-100. Individual aliquots of the lysate were combined with an equal volume of the reaction buffer supplemented with 8 mM CaEGTA alone or in combination with different amounts of 1B7. To start the reactions, an equal volume of reaction buffer supplemented with 1 mM GTP,

0.1 mM ATP, and 50 μM of either ATP γS or 6-Fu-ATP γS (BioLog) was added to each sample. Reactions were allowed to proceed for 30 min at 30 °C, before alkylating with 2 mM p-nitrobenzyl mesylate (Epitomics) for 1 h. Samples were analyzed by Western blot using a rabbit-anti-thiophosphate ester (51-8; Epitomics) at a dilution of 1:10,000 and an appropriate loading control. Goat-anti-rabbit conjugated to IRdye 680 (Li-Cor) was used to visualize the signal from the thiophosphate ester antibody. To compare TgCDPK1-dependent thiophosphorylation in the different *T. gondii* strains, lysates were prepared from strains grown in 1.5 $\mu\text{g}\cdot\text{mL}^{-1}$ for 48 h. Reactions were initiated as described above, except CaEGTA was substituted for K_2EGTA in the samples without Ca^{2+} .

Crystallization. Recombinant TgCDPK1 and 1B7 were combined in a 1:1.5 molar ratio at room temperature, rotating for 1 h. The mixture was loaded on a Superdex 200 16/60 SEC column to separate complex from excess 1B7. The complex peak was pooled, concentrated, and dialyzed into 50 mM Tris, pH 7.4, and 150 mM NaCl for 16–20 h before crystal-tray setup. Initial crystals of TgCDPK1-1B7 complex grew via vapor diffusion at 4 $\text{mg}\cdot\text{mL}^{-1}$ in 25% (wt/vol) PEG 3350, 0.2 M $(\text{NH}_4)_2\text{SO}_4$, and 0.1 M Bis-Tris, pH 6.5 at 18 °C, after 1 d of incubation in a 96-well, sitting drop tray (IndexHT, Hampton Research). Protein and mother liquor were mixed in a 1:1 ratio. The use of 4% (vol/vol) 1-propanol, as an additive, produced optically superior crystals. Scaling-up to a 24-well, hanging-drop format and immediate microseeding of the crystallization drop produced diffraction-quality crystals that grew as thin, mechanically separable plate clusters. SeMet-derivatized protein crystals were obtained following this protocol, albeit they grew singular, were smaller, and took several days longer to grow. Individual crystals were serially cryoprotected in mother liquor supplemented with 20% (vol/vol) glycerol or 35% (wt/vol) PEG 3350 before cooling in liquid nitrogen.

Data Processing and Structure Determination. Complete native and SeMet datasets were collected at the Advanced Photon Source user end station 24-IDC. Data reduction was performed in *HKL2000* (36). MR and SAD phasing was performed using *phaser* in the PHENIX suite (37), available through *SBGrid* (38). An initial MR solution was obtained in the SeMet dataset using the Ca^{2+} -depleted KD of TgCDPK1 (PDB ID code 3KU2, residues 44–313) and a VHH with the complementarity determining regions (CDRs) removed (PDB ID code 1BZQ) as search models in *phaser-MR*. No solution was obtained with the active KD (PDB ID code 3HX4) or the CAD from either Ca^{2+} -bound or Ca^{2+} -depleted TgCDPK1. Even with weak anomalous data, an MR-SAD search strategy using the MR solution in *phaser-EP* yielded seven selenium sites of the possible 30 (2 copies per asymmetric unit in P2₁). Only one of these sites was in the placed KDs. We took advantage of the 14 known methionine positions in the two placed KDs by inserting selenium atoms at the side chain sulfur positions. Inputting these sites into the subsequent *phaser-EP* run resulted in the deletion of only a single site, the XYZ, B-factor, and occupancy refinement of all other input sites, and the identification of five new sites in the CAD. Application of the known noncrystallographic symmetry operator to the selenium sites resulted in a total of 23 sites. After solvent flattening and noncrystallographic symmetry map averaging in *parrot*, available through the *CCP4* suite (39), clear helical density could be seen corresponding to the missing CAD. Iterative rounds of model building in *Coot* (40), *phaser-EP* map calculation, and *parrot* density modification were performed until the map figure of merit was greater than 0.5. *Phenix.refine* (37) was used for model refinement in the SeMet dataset until it was clear that experimental phase restraints were hindering further model improvement. The partially built and refined model was placed into the native P1 unit cell (4 copies per asymmetric unit) using *phaser-MR*. Sequence assignment was aided by the use of Se sites as positional markers and structural homology to the high-resolution, active CAD structure (rmsd, 3.2 Å; 137 C α s aligned). Iterative building in *Coot* and refinement in *Phenix.refine* were performed in the native dataset to obtain the final, fully refined structure.

Structure Analysis. The CDPK1:1B7 interface was analyzed using PDBePISA (41). A multiple sequence alignment of TgCDPK1 homologs was created using ClustalW in JalView (42).

MD Simulations. Simulated systems were modeled from the X-ray crystal structure of active, Ca^{2+} -bound TgCDPK1 (PDB ID code 3HX4) (11). TgCDPK1 with the Ca^{2+} -bound CAD present (+CAD, residues 30–503) or removed (–CAD, residues 30–314) was solvated with 150 mM NaCl in cubic simulation boxes of 93 Å or 75 Å per side, respectively. There were $\sim 81,000$ atoms in each +CAD simulation and 43,000 atoms in each –CAD simulation. The systems were parameterized using the Amber ff99SB-ILDN (43–45) force field, combined with the ff99SB* backbone correction (46) for proteins, and a TIP3P water model

(47). After parameterization, the systems were energy-minimized and then equilibrated in a 10-ns simulation with an ensemble of constant number of atoms and constant temperature and pressure (NPT) at 310 °K and 1 bar. In the equilibration, 1 fs time step was used, and harmonic positional restraints of a force constant of 1 kcal/mol/Å² were initially applied to the protein backbone atoms. Production simulations were performed starting from the last frame of equilibration simulations with an NPT ensemble at 310 °K and 1 bar using the Nosé-Hoover thermostat (48) with a time constant of 0.05 ps. All hydrogen-containing bonds were constrained using an implementation (49) of the M-SHAKE algorithm (50). A reversible reference system propagator algorithm integrator (51) was used. Bonded, Van der Waals, and short-range electrostatic interactions were computed every 2 fs, whereas long-range electrostatic interactions were computed every 6 fs. The short-range electrostatic interactions were calculated at a cutoff of 9.0 Å, and the long-range electrostatic interactions were computed using Gaussian-split Ewald (52). All simulations were carried out on either Anton or Anton 2, special purpose supercomputers designed for MD (53).

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