A divergent protein kinase A in the human pathogen *Leishmania* is associated with 

developmental morphogenesis

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Summary

Parasitic protozoa of the genus *Leishmania* cause human leishmaniasis. They cycle between the phagolysosome of mammalian macrophages, where they reside as round intracellular amastigotes, and the mid-gut of female sand flies, which they colonize as elongated extracellular promastigotes. Shifting promastigotes to a lysosome-like environment (pH 5.5 and 37C, 5% CO) initiates their development into amastigotes. Previous studies suggested a role for protein kinase A (PKA) in this differentiation process. Here, we describe a new, divergent, regulatory subunit (PKAR3) found only in a limited member of the family Kinetoplastidae. In *L. donovani*, phosphorylation of PKAR3 was regulated by the differentiation signal and coincided with parasite morphogenesis during stage development. LdPKAR3 was bound to the subpellicular microtubules cell cortex via a formin homology (FH2)-like domain at the tip of a large and divergent N-terminal domain. Immunoprecipitation, fluorescence resonance energy transfer (FRET), proteomics analyses, and structural modeling showed that PKAR3 selectively binds the C3 isoform of the PKA catalytic subunit in a holoenzyme complex. In promastigotes, PKAR3 recruited PKAC3 to the subpellicular microtubules at the central cortex. After exposure to a differentiation signal, PKAR3 homogenously distributed across the entire cortex, in concert with cell rounding. Deleting the genes encoding either the R3 or C3 subunit resulted in premature rounding of the promastigote population, indicating that PKA determines the normal elongated shape. Regulation of *Leishmania* developmental morphogenesis by interaction with the subpellicular microtubule corset is a novel function for a unique PKA complex not present in the host cell.

**Keywords:** differentiation, protein kinase A, signaling, microtubule cortex, morphogenesis, *Leishmania*, parasitism
Introduction

*Leishmania donovani* is a protozoan, trypanosomatid parasite that causes kala azar, a fatal form of visceral leishmaniasis in humans\(^1\). These organisms cycle between the midgut of female sand flies, where they reside as flagellated extracellular promastigotes, and the phagolysosomes of mammalian macrophages, where they live as aflagellate amastigotes\(^3\). Infective metacyclic promastigotes are introduced into the host during the sand fly blood meal, after which they are phagocytosed by resident macrophages near the bite site\(^5,6\). Once inside the host phagolysosome, promastigotes encounter two physical cues that distinguish this environment from that of the vector gut, i.e., acidic pH (~5.5) and elevated temperature (~37ºC). Promastigotes process these cues into a signal that initiates development into amastigotes\(^7,8,9\). However, the molecular details of this signaling pathway are still largely unknown.

Development of host-free systems using axenic parasites has advanced our understanding of the molecular mechanisms underlying *Leishmania* intracellular development. *L. donovani* differentiation can be induced in culture, by exposing promastigotes to the high temperature and acidity (37ºC, pH 5.5, 5% CO\(_2\)) typically found in the phagolysosome\(^10,11,12\). Differentiation to mature amastigotes takes 5 days, similar to the time it takes *in vivo\(^13*\). Among the earliest events during *L. donovani* promastigote-to-amastigote differentiation are changes in the phosphorylation status of many proteins\(^14\). Significantly, protein kinase A (PKA)-driven phosphorylation is common in promastigotes, with most of the associated phosphoproteins being dephosphorylated within a few minutes after initiation of promastigote-to-amastigote differentiation\(^7\). This suggests that regulation of PKA activity is important for initiation of *Leishmania* development.
PKA-mediated signaling pathways are ubiquitous in eukaryotic cells, and are implicated in growth control, development, and metabolism. The canonical PKA regulatory system in vertebrates involves assembly of two catalytic (PKAC) and two regulatory (PKAR) subunits into an inactive holoenzyme complex, mediated by dimerization and docking (D/D) domains in the N-terminal portion of PKAR. When cAMP binds the cyclic nucleotide-binding domains (cNBDs) near the C-terminal of each PKAR, the resultant conformational change triggers dissociation of the PKAC-R complex and release of active PKAC subunits. Compartmentalization to microdomains of PKA activity is achieved (at least in higher eukaryotes) through interaction of the D/D domain of PKAR with a diverse family of A kinase-anchoring proteins (AKAPs) that vary according to biological context. However, PKARs from many lower eukaryotes (including trypanosomatids) appear to lack the D/D domain.

Most eukaryote genomes, with the notable exception of plants and algae, encode one or more PKAC subunits. Trypanosomatids are no exception, usually bearing two subunits (C1 and C2) that are practically identical and a third (C3) that is more divergent. Phosphoproteomic analyses of *L. donovani* revealed that all three PKAC subunits contain at least two phosphorylation sites. The one within the kinase active site loop, matches the canonical PKA phosphorylation motif (RXXpS/pT) and is only phosphorylated in promastigotes. The second is located close to the C-terminus within a canonical ERK1/ERK2 substrate motif (pSP). LdPKAC1/C2 is phosphorylated at this site in promastigotes, but the site in LdPKAC3 is only phosphorylated after exposure to the differentiation signal. In other eukaryotes, phosphorylation of both a threonine (T197) in the activation loop and a serine (S338) near the C-terminus, is necessary for full activation of PKAC. Following the phosphorylation pattern of
these motifs in \textit{L. donovani} indicate that C1 and C2 are active in promastigotes, while C3 is activated only upon receiving the differentiation signal.

Most eukaryotes also possess one or more PKAR subunits, whose number and evolution are more dynamic than for PKAC\textsuperscript{18}. All trypanosomatids have a regulatory subunit (PKAR1) that localizes to the flagellum of \textit{Trypanosoma brucei}\textsuperscript{22,23,24} and \textit{L. donovani}, where it is expressed only in the promastigote stage\textsuperscript{7,25}. However, in contrast to higher eukaryotes, trypanosomatid PKAR1 appears to be monomeric and unable to bind cAMP\textsuperscript{17,19}.

Interestingly, \textit{Leishmania} and some related genera, including \textit{Trypanosoma cruzi}, but not \textit{T. brucei}, have a second regulatory subunit (PKAR3)\textsuperscript{20}, which in \textit{L. donovani} contains >10 sites that are dynamically phosphorylated and dephosphorylated during differentiation\textsuperscript{14}. However, until now, little information has been available concerning the potential role of PKAR3. Here, we show that the PKAR3-PKAC3 holoenzyme complex is anchored to the subpellicular microtubule at the cell cortex \textit{via} a formin homology (FH2)-like domain. Upon exposure to the differentiation signal, the C3-R3 holoenzyme dissociates and R3 relocates along the subpellicular microtubules, while in parallel promastigotes round up to form amastigote-shaped cells.

Association between PKAR3 and PKAC3 is necessary for maintenance of the elongated shape of \textit{Leishmania} promastigotes, as evidenced by the abnormal shape of mutants lacking either PKAR3 or PKAC3. This study tie together an evolutionary divergent class of PKA and the microtubule cytoskeleton of \textit{Leishmania} with developmental morphogenesis. This study sheds new light on the process initiating intracellular parasitism.
Results

*Leishmania PKA regulatory subunits are evolutionarily divergent*

Phylogenetic analyses of the PKAR subunits of a wide range of organisms revealed that they have undergone dynamic and divergent evolution during eukaryote history\(^\text{18}\). Analysis of the genomes available from 39 different species in the phylum *Euglenozoa* revealed that all taxa (except for *Perkinsela*, which is an obligate endosymbiont) contain 2-5 PKA catalytic subunits and one or two regulatory subunits (see Table S2), with the latter being quite different from other eukaryotic PKARs in both sequence and domain structure (Fig. 1). Phylogenetic analysis revealed three distinct groups of PKAR sequences, two of which were restricted to the *Euglenozoa* (Fig. 1A). Both the trypanosomatid sequences (PKAR1 and PKAR3) diverged widely from those found in other eukaryotes (including the RI and RII classes found in animals), including those of their closest relatives (*Trichomonas, Giardia* and *Naegleria*) in the Excavata supergroup. While both trypanosomatid PKAR1 and PKAR3 contain a conserved inhibitor/pseudo-inhibitor sequence motif (in the middle of the protein sequence) and two cyclic nucleotide-binding domains (cNBDs) in their C-terminal portion, they differ substantially from each other (and from PKARs in other organisms) in their N-terminal portion (Fig. 1B). Instead of the canonical dimerization and docking (D/D) domain found in many (but not all) other eukaryotes, all trypanosomatid PKAR1s contain an RNI-like domain, possibly explaining their inability to form a homodimer \(^\text{17}\). However, the leucine-rich repeats (LRRs) in the RNI-like domain may mediate different protein-protein interactions. *Diplonema papillatum* and *Euglena gracilis* (both of which are free-living flagellates outside of the Kinetoplastea class), as well as *Bodo saltans* (a free-living kinetoplastid) and *Paratrypanosoma confusum* (an early branching trypanosomatid parasite of mosquitoes) \(^\text{26}\) all contain a second paralogue of this gene (PKAR2),
with the *Euglena* version containing three additional pairs of cNBDs. Interestingly, Pcon_PKAR2 segregates outside the other Euglenozoan PKARs and lacks the RNI-like domain, suggesting it may represent a more “ancestral” version. While many trypanosomatids contain second, more divergent, PKAR paralogues (PKAR3), it is conspicuously absent from the Salivarian (*Trypanozoon* and *Nannomonas* sub-genera) *Trypanosoma* that lack an intracellular amastigote stage, as well as *Phytomonas* and *Herpetomonas*, which are the insect and/or plant pathogens in the Phytomonadinae subfamily. PKAR3 has a slightly longer C-terminal extension (following the cNBDs) and a much longer N-terminal region, with no sequence similarity to other PKARs (Fig. 1B). This region contains no obvious protein domains or motifs, although HHpred analysis (https://omictools.com/hhpred-tool) identified weak structural similarity at the N-terminus of LdPKAR3 (but not those from other species) with a formin homology (FH2)-like domain.

*Cytoskeletal localization of LdPKAR3 to cortical subpellicular microtubules*

Polyclonal antibodies raised in rabbits against recombinant LdPKAR3 reacted with a 72 kDa protein in wild-type *L. donovani* (+/+ in Fig. S1A). The protein was absent in null mutants (-/- in Fig. S1A) but was detected after ectopic expression of a full-length copy of the *LdPKAR3* gene in the null mutant (-/-+ in Fig. S1A), thereby confirming its identity. Confocal microscopy using dual labeling with antibodies against LdPKAR3 and the plasma membrane of *L. donovani* promastigotes revealed that LdPKAR3 localizes at the cell cortex (Fig. 2). In promastigotes, LdPKAR3 was restricted to the cortex between the nucleus and flagella pocket, a region representing only 45% (±10%) of the cell length (n=47 cells). In contrast, LdPKAR3 covered the entire cortex of both axenic and intracellular amastigotes (Fig. 2).
The *Leishmania* cell cortex contains an abundant rigid cytoskeleton comprised of stable subpellicular microtubules\(^\text{27,28,29}\). Previous studies have shown that non-ionic detergents can solubilize the trypanosomatid surface membrane, without impacting the subpellicular microtubules\(^\text{30,31,32}\). *Leishmania* cells treated with the zwitterion detergent n-dodecyl β-D-maltoside (DDM, 0.5%, which is neutral at pH 7) retained LdPKAR3 in the non-soluble pellet of promastigotes collected before, and 5 hours after exposure to the differentiation signal (Fig. 2A). As expected, β-tubulin was also enriched in the DDM-insoluble pellet (Fig. 3A and Fig. S2A). Indirect immunofluorescence indicated that this fraction contained the subpellicular microtubules, which retained the shape of promastigotes and amastigotes (Fig. S2B). Efficient solubilization of the parasite plasma membrane was confirmed by the absence of the integral membrane protein LdAAP24\(^\text{33}\) and the cytosolic HSP83\(^\text{11}\) in the enriched microtubules (Fig. S2C). Thus, we conclude that LdPKAR3 associates with the subpellicular microtubules cytoskeleton in *L. donovani* promastigotes.

Binding of PKARIβ to microtubules has been previously described in neuronal axons of mammalian cells\(^\text{34}\), where binding was mediated by interaction between the dimerization and docking (D/D) domain of R1β and a microtubule-associated protein 2 (MAP2)\(^\text{35}\). As indicated above (Fig. 1B), LdPKAR3 lacks the canonical D/D domain but instead contains an FH2-like domain at its N-terminus. FH2 is one of three cytoskeleton-binding domains found in formins, a diverse family of proteins that regulate cytoskeleton rearrangement in many eukaryotes\(^\text{36}\), which led us to speculate that this domain might be responsible for LdPKAR3 binding to the subpellicular microtubules. To test this hypothesis, a truncated version of LdPKAR3 (lacking the N-terminal 90 amino acids that contains the FH2 domain) was ectopically expressed in mutant (*Δldpkar3*) promastigotes. As shown in Fig. 3B, add-back of full-length (FL) LdPKAR3 resulted in restoration
of LdPKAR3 binding to the DDM-insoluble subpellicular microtubules, whereas the truncated version (Δ90) did not bind to the microtubules and remained in the supernatant. Hence, binding of LdPKAR3 to the microtubule cytoskeleton is mediated through the N-terminal FH2-like domain. Interestingly, the truncated LdPKAR3 protein appears to undergo proteolytic cleavage in the supernatant, suggesting that as shown in T. equiperdum\textsuperscript{17}, binding to the subpellicular microtubules is essential for its stability.

\textit{LdPKAR3 associates with a catalytic subunit to form a PKA holoenzyme}

To verify that LdPKAR3 can be classified as regulatory subunit of PKA, we tested its interaction with the PKA catalytic subunits. Because no antibodies are available for the \textit{L. donovani} PKAC isoforms, we expressed LdPKAR3-Ty1 under control of a tetracycline repressor in a \textit{T. brucei Δpkar1} cell line\textsuperscript{7,19}. The absence of TbPKAR1 in this cell line eliminates the possible competition with LdPKAR3 for interaction with TbPKAC subunits. As expected, expression of LdPKAR3-Ty1 increased in the presence of tetracycline (Fig. 4A), although some “leaky” background expression was also present in the absence of tetracycline. Upon detergent fractionation of \textit{T. brucei} (Fig. 4B), LdPKAR3 was found in the DDM-soluble supernatant as well as in the DDM-insoluble pellet, indicating that LdPKAR3 preserves its association with the cell cortex cytoskeleton as seen in \textit{L. donovani} (Fig. 3A). TbPKAC3 was enriched in the DDM-insoluble fraction, suggesting that an R-C complex may be associated with the cell cortex cytoskeleton. Affinity purification of LdPKAR3-Ty1 from the DDM supernatant using anti-Ty1 antibody coupled to magnetic beads, and western blotting with antibodies against the TbPKAC subunits, indeed showed stable interaction of LdPKAR3-Ty1 with TbPKAC3 (Fig. 4C), but not with TbPKAC1/2 (Fig. 4D). Interestingly, upregulation of LdPKAR3-Ty1 expression was
accompanied by increased TbPKAC3 levels, possibly due to its stabilization in a complex with LdPKAR3 (Fig. 4A).

To further confirm the interaction between homologous LdPKAC3 and LdPKAR3, both proteins were co-expressed in *L. tarentolae* using the LEXSY expression system (Jena Bioscience) and the putative complex was purified by tandem affinity chromatography using the N-terminal His6 tag on LdPKAR3 and Strep tag on LdPKAC3. The proteins were well expressed and LdPKAC3 co-purified with LdPKAR3 (Fig. 4E). As positive control, the tagged LdPKAR1 and LdPKAC3 were co-expressed and co-purified in a parallel experiment (Fig. 4F) since TbPKAR1/TbPKAC complexes were previously reported in *T. brucei* 19. Thus, we conclude that the LdPKAR3 forms a stable PKA catalytic subunit kinase, as expected for a regulatory subunit.

Western blot analysis of transgenic *L. donovani* WT and Δldpkar3 promastigotes detected HA-tagged LdPKAC3 was detected (at about the same level) in cell lysates of both the WT and mutant cell lines (Fig. 5A). In contrast, it was present at only near-background levels in the microtubule fraction of Δldpkar3. These results were confirmed by proteomic analysis of the DDM-insoluble fraction of WT and Δldpkar3 promastigotes (data are available via ProteomeXchange with identifier PXD025222) (Fig. 5B). The relative abundance of LdPKAC3 (calculated from spectral counts for LinJ.18.1090) was ~30-fold higher in WT compared to Δldpkar3 microtubules (Fig. 5B). Similar results were obtained upon analysis of promastigotes before (0 h), and 5 h after exposure to the differentiation signal, although LdPKAR3 and LdPKAC3 levels were usually somewhat lower at 5 h. In contrast, other proteins, not involved in the interaction with R3 showed little difference in relative abundance between samples, as illustrated by actin (ACT) levels, shown in Fig. 5B. These experiments confirm that LdPKA-R3 is essential for recruitment of LdPKAC3 to the subpellicular microtubules.
The molecular proximity of LdPKAR3 and LdPKAC3 within transgenic *L. donovani* promastigotes expressing HA-tagged LdPKAC3 was further demonstrated using fluorescence resonance energy transfer (FRET). Control experiments using cells labelled separately with each of the secondary antibodies showed only background fluorescence, but no FRET (Fig. S3). In contrast, FRET was clearly observed when both antibodies were added together (Fig. 5C). Each dot represents a point of interaction between LdPKAR3 and LdPKAC3, with the color scale reflecting the intensity of interaction (*i.e.* proximity between the two molecules). These results further confirming the LdPKAR3-mediated recruitment of PKAC3 to form a (putative) holoenzyme at the subpellicular microtubules of the parasite cell cortex.

**Structural analysis of the interaction between PKAR3 and PKAC3**

To date, we have been unable to experimentally determine the structure of PKAR3, although an X-ray structure is available for *T. cruzi* PKAR1. However, the mammalian PKAR and PKAC subunits have been extensively studied at the atomic level, both alone and in complex, enabling homology modeling. The N-terminal sequence of LdPKAR3 protein is divergent from that of human (and other) PKAR subunits (see Fig. 1B) and is predicted to be mostly low-complexity and disordered, with the exception of the N-terminal 80 amino acids; this region was omitted, and models were built using only the C-terminal portion of LdPKAR3, starting from the inhibitor/pseudo-inhibitor sequence. The equivalent region of *Bos taurus* (Bt) PKAR1α (*i.e.* lacking a DD domain) forms a homodimer where the R/R interface acts as a cAMP sensor. The residues responsible for the Bt R/R interaction are divergent in LdPKAR3 (Fig. 6A), preventing the formation of critical hydrogen bonds between the Y120K121 side-chains of PKAR1α (Q350E351 in LdPKAR3) and R144S145 (I374A375 in LdPKAR3), suggesting that LdPKAR3 does not form a homodimer, reminiscent of the monomeric R subunit of
trypanosomes\textsuperscript{19}. In addition, the cAMP-binding sites of BtPKARI\(\alpha\) are not conserved in LdPKAR3, and the strictly conserved residues in the mammalian cNBDs (R\textsubscript{210} and R\textsubscript{334} in BtPKARI\(\alpha\)) that stabilize the cAMP phosphate are replaced by N\textsubscript{443} and A\textsubscript{567} (Fig. 6B).

Furthermore, a change from R\textsubscript{239}\textsuperscript{R1} to D\textsubscript{473}\textsuperscript{R3} likely prevents the formation of a critical regulatory-salt bridge with an upstream residue (E\textsubscript{143}\textsuperscript{R1}/D\textsubscript{373}\textsuperscript{R3}) that strains the B/C helix between the cNBDs and favors the formation of the binding interface\textsuperscript{42}. Similar non-conservative amino acid changes are also found in LdPKAR1 (A\textsubscript{319} and T\textsubscript{443}), likely explaining why \textit{T. equiperdum} and \textit{T. brucei} PKAR1 bind neither cAMP nor cGMP\textsuperscript{17,19}.

Based on homology modeling based on the mammalian PKARI\(\alpha\)-C complexes, the salt-bridge triad (R\textsubscript{240}\textsuperscript{R1}-D\textsubscript{266}\textsuperscript{R1}-R\textsubscript{194}\textsuperscript{C}) that stabilizes interaction between the regulatory and catalytic subunits appears to be conserved in \textit{L. donovani} PKAR3 and PKAC3 (R\textsubscript{475}\textsuperscript{R3}-E\textsubscript{501}\textsuperscript{R3}-R\textsubscript{166}\textsuperscript{C3}), as are the PKAC activation loop (T\textsubscript{169}\textsuperscript{C3}) and the regulatory salt-bridge (Q\textsubscript{495}\textsuperscript{R3}-R\textsubscript{601}\textsuperscript{R3}) responsible for PKA activation (Fig. 6C). However, other residues involved in regulation of PKA catalytic activity by BtPKARI\(\alpha\) are not conserved. For example, non-conservative replacement (R\textsubscript{95}\textsuperscript{R1}/S\textsubscript{324}\textsuperscript{R3}) of the first residue in the predicted LdPKAR3 inhibitor/pseudo-inhibitor site is predicted to disrupt interaction with negatively charged and polar residues (T\textsubscript{23}, E\textsubscript{99}, E\textsubscript{142} and Y\textsubscript{302}) in LdPKAC3 (Fig. 6D).

\textit{Polar localization of LdPKAR3 is developmentally regulated and associated with morphogenesis}

To better characterize the changes in LdPKAR3 localization along the cortex subpellicular microtubules during development from promastigotes to amastigotes, DDM-insoluble fractions obtained at various times following exposure to the differentiation signal were
analyzed (Fig. 7). In promastigotes before differentiation, counter-staining of β-tubulin showed that LdPKAR3 was primarily localized in the central region of the cytoskeleton (Fig. 7, left column). In contrast, LdPKAR3 was uniformly distributed over the entire subpellicular microtubule network as early as 12 h after exposure to the differentiation signal (Fig. 7, middle column), and uniformly labelled subpellicular microtubules of fully differentiated axenic amastigotes (120 hours) (Fig. 7, right column). Thus, it appears that LdPKAR3 mobilization correlates with the timing of parasite rounding, which begins ~7 hours and is nearly complete by 12 hours, as we have previously described during phase II of differentiation.

These "coordinated" events prompted us to hypothesize that an intact LdPKAR3-C3 holoenzyme at the central region of the subpellicular microtubules is necessary for maintaining the elongated shape of promastigotes. Promastigotes of wild type and mutants lacking R3 (\( \Delta ldpkac3 \)) or C3 (\( \Delta ldpkac3 \)) were subjected to Image Streaming \(^{43,44,45} \) and confocal fluorescence analysis. Indeed, while the majority (53%) of WT promastigotes were elongated, with only 16% round (Fig. 8A), the proportion of round cells increased to 44% in \( \Delta ldpkac3 \) mutants, and to 49% in \( \Delta ldpkac3 \) mutants. The proportion of elongated cells in these mutant populations decreased to 20% and 18%, respectively; while the proportion of “ovoid” cells (36% and 33%, respectively) remained similar to that in WT samples (31%). Ectopic expression of the full-length \( LdPKAR3 \) or \( LdPKAC3 \) genes in the null mutants partially restored the ratio of elongated to rounded cells (38%:26% and 31%:30%) to WT levels (53%:16%). Confocal fluorescence analysis using the anti-membrane antibodies confirmed the shape changes enforced by both C3 and R3 gene deletion (Fig. 8B). Taken together, it is the interaction between LdPKAR3 and LdPKAC3 and localization of the holoenzyme to the central region of the subpellicular microtubules that maintain the elongated cell morphology in \textit{Leishmania} promastigotes.
Discussion

Upon entry into their mammalian host, *Leishmania* promastigotes are internalized into macrophage phagolysosomes, where lysosome-specific environmental signals, such as acidic pH (~5.5) and body temperature (~37°C) act as cues for differentiation into amastigotes\textsuperscript{11,46,47,48}. Our previous studies identified PKA as a potential key enzyme in this process\textsuperscript{7}. The present work identified and characterized a highly divergent regulatory subunit of *Leishmania* PKA (LdPKAR3) as another key player in promastigote-to-amastigote differentiation. In promastigotes, LdPKAR3 localized to subpellicular microtubules in the central region of the cell cortex, where it associated with the catalytic subunit isoform PKAC3. This localization proved essential for maintenance of the elongated morphology typical of promastigotes, as shown by cell rounding upon loss of either subunit. Upon exposure to the differentiation signal, LdPKAR3 relocated to the entire cell cortex, coinciding with the time into differentiation when promastigotes assume the rounded morphology of amastigotes.

To date, studies on the role of the cytoskeletal microtubules on *Leishmania* development and morphogenesis have focused mostly on the flagella\textsuperscript{49,50}. The complex axoneme structure and its dynamics during differentiation from the flagellated promastigotes to the aflagellated amastigotes attracted significant attention. The promastigotes axoneme has the typical 9+2 structure, but, during differentiation into amastigotes, the flagella shorten and then collapse to become a sensory flagellum with a 9+0 arrangement inside the flagella pocket\textsuperscript{51}. Because flagella shortening parallels change in cell morphology, it was believed that these processes are linked. However, using mutant promastigotes whose flagella shortened and transformed to the sensory 9+0 proved the opposite; flagella shortening and morphogenesis are not linked processes\textsuperscript{52}. In this study, we demonstrated that the differentiation signal activates the relocation of LdPKAR3
along the entire subpellicular microtubule cortex, in parallel to rounding to amastigote-shaped cells. The observation that a novel regulatory subunit of PKA plays a role in *Leishmania* morphogenesis supports the notion that this process is not evolutionarily conserved. In higher organisms, it is mainly actin together with myosin motor proteins that are responsible for cell morphogenesis\textsuperscript{53,54}. These observations set the stage for more in-depth studies into the role of microtubules in remodeling cell shape and differentiation-induced morphogenesis.

Four independent methodologies were used to demonstrate that LdPKAR3 forms a holoenzyme complex with LdPKAC3. Firstly, co-immunoprecipitation of LdPKAR3 and PKAC3 ectopically expressed in *T. brucei* and *L. tarentolae* showed their tight and stable interaction. Secondly, proteomic analysis of the DDM-enriched subpellicular microtubules showed that LdPKAR3 associates with LdPKAC3 in *L. donovani*. Thirdly, confocal immunofluorescence combined with FRET analysis confirmed the proximity of LdPKAR3 and LdPKAC3 at the parasite cell cortex of *L. donovani*. Finally, molecular modelling identified conserved amino acid residues at the interface between the mammalian PKA catalytic and regulatory subunits in LdPKAR3 and LdPKAC3. However, as shown for the *T. brucei* TbPKAR1\textsuperscript{17,19}, LdPKAR3 lacks a salt bridge that contributes to PKAR dimerization in higher eukaryotes and lacks the D/D domain and canonical cAMP binding sites. We conclude that *Leishmania* PKAR3 forms a R-C heterodimer whose activation is independent of cAMP. It is currently not known whether LdPKAC3 can bind an alternative (unknown) ligand similar to TbPKAR1\textsuperscript{19}.

An intriguing phenomenon described in this study is that LdPKAR3 is bound to the stable subpellicular microtubules cortex of *Leishmania*. The only other example of similar phenomenon is the mammalian PKARI\(\beta\) that associates with the axonal stable microtubules. This regulatory protein binding to the microtubules is carried out via the D/D domain and mediated by MAP2\textsuperscript{55}. 
However, LdPKAR3 lacks the canonical D/D, indicating that binding of this ancient regulatory subunit to the subpellicular microtubules differs from that of the mammalian PKARIβ. Moreover, Trypanosomatids genomes lack MAP2 orthologues, which only appear later in vertebrate evolution. The presence of a predicted FH2 domain at the N-terminus of LdPKAR3 and the importance of this domain for localization raised the intriguing possibility that it mediates binding to subpellicular microtubules within the cell cortex of Leishmania, thus suggesting evolution of an alternative PKA tethering mechanism distinct from the mammalian D/D domain to MAP2 interaction. In other organisms, FH2 is one of three domains found in formins, a family of proteins involved in cell shape determination, adhesion, cytokinesis, and morphogenesis, through their remodeling effect on actin or microtubule cytoskeletons. However, the absence of significant HHpred matches for this region in other PKAR3 orthologues (including most Leishmania species) challenges this hypothesis.

PKAR1 retains conserved amino acid interactions at the inhibitor/pseudo-inhibitor site and functions as a ligand-dependent PKAC inhibitor, as reported previously for T. brucei PKAR1 (Bachmaier et al., 2019). A non-conservative change in the LdPKAR3 inhibitor/pseudo-inhibitor site may weaken the ability of LdPKAR3 to act as a pseudo-substrate and inhibitor for the catalytic subunit. Conversely, both S329R3 and S332R3 are predicted to be equally distant in space from T169C3, thereby allowing phosphorylation of either to cause the same repulsion effect that phosphorylation of S101R (which corresponds to S329R3) exerts on T197C of the bovine PKA-R/C complex. This phosphorylation is postulated to enhance catalytic activity even without cAMP stimulation. Therefore, it remains possible that LdPKAR3 regulates PKA catalytic activity by a mechanism independent of ligand binding. Interestingly, S332 phosphorylation is
upregulated two-fold immediately after exposure to the differentiation signal\textsuperscript{14}, which may explain the increase in PKA-mediated phosphorylation at LdPKAR3 T\textsubscript{262} early in differentiation.

In summary, this study is the first to characterize a new, divergent PKA regulatory subunit (LdPKAR3) that appears to be restricted to trypanosomatids with an intracellular lifecycle stage. The observation that LdPKAR3 is bound to the subpellicular microtubule and migrates upon exposure to the differentiation signal suggests that it may be part of the lysosome sensing pathway involved in initiating parasite differentiation inside its host\textsuperscript{48}. We are currently investigating the mechanism by which LdPKAR3 and LdPKAC3 influence differentiation-related morphogenesis.
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Author contributions

RFW, SB, RD, MA, GBG, EP, PT and RNK conducted the experiments. IQP conducted the structure modeling analysis. EP conducted the bioinformatics and phylogenetics analyses. PJM and MB supervised all bioinformatics and phylogenetic analysis and analyses in the heterologous systems, respectively, and contributed to writing the manuscript. DZ managed the project and wrote the manuscript.

Declaration of interests

The authors declare no competing interests.
**Figure legends**

**Fig. 1: Phylogenetic analysis of PKA regulatory subunits.** (A) Maximum-likelihood phylogenetic trees were constructed using the RaxML module of Geneious on a MAFFT alignment of amino acid sequences of the PKAR orthologues from 39 Euglenozoan species and 14 other species in the Amoebozoa, Apusozoa, Archaeplastida, Excavata, Opisthokonta or SAR (Stramenopiles/Alveolata/Rhizaria) supergroups. Red dots indicate PKAR orthologues containing a canonical (red fill) or non-canonical (light red fill) dimerization/docking (D/D) domain. Green dots indicate PKAR orthologues containing an RNI-like domain. (B) Schematic representation of the protein domains within selected PKAR orthologues. The cAMP binding domains are indicated by blue boxes, the dimerization domain in red (canonical) or light red (non-canonical) and the RNI-like domain in green; with the inhibitory (pseudo-)phosphorylation sites marked by a brown or tan box.

**Figure 2: LdPKAR3 relocates from the central region of the cortex in promastigotes to the entire cell cortex during differentiation to axenic and intracellular amastigotes.** Indirect immunofluorescence of LdPKAR3 in promastigotes (left column), axenic (middle column) and intracellular amastigotes (48 h after THP1 macrophage infection, right column). Cells were stained with anti-LdPKAR3 (upper row, green) and anti-membrane (middle row, red) antibodies. Fluorescence was detected using a Zeiss LSM 700 inverted confocal microscope. Scale bar = 5 μm.
Figure 3: LdPKAR3 is bound to the subpellicular microtubules of *L. donovani* via a FH2 formin domain. (A) Subpellicular microtubules were extracted from wild type *L. donovani* promastigotes before and 5 h after exposure to a differentiation signal, using 0.5% detergent n-dodecyl β-D-maltoside (DDM; details in STAR Methods). Proteins in whole cell lysates, DDM-insoluble pellets (P) and DDM-soluble supernatants (S) were separated by 10% SDS-PAGE and then subjected to western blot, using anti-LdPKAR3 (upper row), anti-HSP83 (a soluble protein marker, middle row) and anti-β tubulin (microtubule-rich fraction, lower row) antibodies. (B) A 90-amino-acid peptide from the LdPKAR3 protein was truncated (Δ90 LdPKAR3). Plasmids expressing this construct and the full-length add-back were expressed in ∆ldpkar3. These and wild type promastigotes were subjected to subpellicular microtube enrichment, using 0.5% DDM and then subjected to western blot using anti-LdPKAR3 (upper row) and anti-β tubulin (lower row) antibodies. (*) C indicates whole cell lysates, P indicates DDM-insoluble pellet and S indicates DDM-soluble supernatant.

Fig. 4: LdPKAR3 interacts with the PKAC3 subunits. (A-D) LdPKAR3 with a C-terminal Ty1-tag was tetracycline-inducibly (+/- Tet) expressed in *T. brucei* MiTat 1.2 pkar1 null cells (Δtbkar1 LdPKAR3-Ty1)7. (A) Western blots probed with anti-PFR-A/C antibody as loading control; LdPKAR3-Ty1 and TbPKAC3 were detected using anti-LdPKAR3 and anti-TbPKAC3 antibodies, respectively, and showed expected molecular masses. (B) DDM-based extraction using the microtubule-associated protein TbSAXO30 as a marker for detergent resistance. Whole-cell lysate (WCL), soluble fractions 1 and 2 (sol) and detergent-resistant pellet (P) show detergent resistance of a fraction of LdPKAR3-Ty1 and enrichment of TbPKAC3 in the pellet fraction. (C-D) The DDM-soluble fractions of LdPKAR3-Ty1 expressing *T. brucei* and of the
parental Δtbpkar1 cell line were subjected to anti-Ty1 pull-down, followed by detection of LdPKAR3-Ty1, TbPKAC3 (C) and TbPKAC1/2 (D) on western blots with rabbit anti-LdPKAR3, anti-TbPKAC3 (C) and anti-TbPKAC1/2 (D) antibodies. IN = input; FT = flow-through; E = elution; M = molecular weight marker. (E-F) Purification of PKA holoenzyme complexes. (E) His6-LdPKAR3 and Strep-LdPKAC3 or (F) His6-LdPKAR1 and Strep-LdPKAC3 were co-expressed in L. tarentolae and tandem-affinity-purified. The soluble input (IN) and eluates (E1, E2) were analyzed by western blot and probed with mouse monoclonal anti-His (Bio-Rad), anti-Strep (Qiagen) antibodies and rabbit anti-LdPKAR3 or anti-TbPKAR1 (cross-reacting with LdPKAR1) antibodies7.

Figure 5: LdPKAC3 binding to the subpellicular promastigotes is mediated by LdPKAR3. (A) WT L. donovani promastigote and Δldpka-r3 (ΔR3) cell lines expressing HA-tagged LdPKAC3 (C3) were subjected to subpellicular microtubule enrichment (0.5% DDM). Proteins from these fractions were subjected to western blotting using anti-HA (upper row), anti-LdPKAR3 (middle row) and anti-β-tubulin (lower row) antibodies. (B) Proteins from DDM-enriched subpellicular microtubules of WT and Δldpka3 samples were loaded on a 10% SDS PAGE. A slice containing proteins at the molecular mass range of 35-42 kDa was excised and subjected to mass-spectroscopy (Orbitrap, Thermo, Ltd). The area below the graph of LdPKAC3 (LinJ.18.1090), LdPKAR3 (LinJ.34.2680) and LdACT (actin, LinJ.04.1250); their abundance relative to the abundance in WT (set to 7000) is illustrated in promastigotes before (■, ■) and 5 h after exposure to a differentiation signal (■, ■). C) Fluorescence resonance energy transfer (FRET) of LdPKAR3-C3 associations at the cell cortex. Promastigotes of WT L. donovani and of a cell line ectopically expressing HA-tagged LdPKAC3 were labeled with rabbit anti-
LdPKAR3 (endogenous protein) and mouse anti-HA antibodies. Subsequently, cells were labeled with Alexa fluor 568-conjugated anti-rabbit and Alexa fluor 647-conjugated anti-mouse antibodies. The emission spectrum of Alexa fluor 568 overlaps with the excitation spectrum of Alexa 647. The FRET emission color and strength were calculated using Axio Vision (details in Materials and Methods). The color scale ruler on the left of each panel indicates the strength of FRET emission.

**Figure 6: 3D modeling of LdPKAR3-C3 interactions.** Homology models of LdPKA-R3 were built based on mammalian structures of PKARIα and PKACα from Bos taurus. (A) Protein sequence alignment of LdPKA-R orthologues with Bos taurus (bovine) PKARIα (Bt_R_2qcs). (B) 3D modelling of the predicted LdPKAR3 cyclic nucleotide binding domains based on bovine PKARIα. (C) LdPKAR3-C3 interacting amino acids are conserved. Side chains of residues involved in the regulatory salt bridges and potential electrostatic interactions downstream of the pseudo-substrate are shown on the ribbon diagram of LdPKAR3 (magenta) and LdPKA-C3 (blue). (D) Side chains of critical residues involved in the PKAR-PKAC interaction for Bos taurus BtPKARIα-PKACα (left) and L. donovani LdPKAR3-PKAC3 (right) are depicted. Note the R95S substitution in the LdPKAR3 pseudo-inhibitor residue compared to BtPKARIα.

**Figure 7: LdPKA-R3 relocates on the subpellicular microtubules during promastigote to amastigote differentiation.** Promastigotes before and 12 h after exposure to a differentiation signal, and mature amastigotes were subjected to 0.5% DDM-driven enrichment of subpellicular microtubules. The extracted microtubules were then subjected to dual fluorescence staining using anti-LdPKAR3 (green, upper row) and anti-β-tubulin antibodies (red, middle panel). Merging the
two fluorescent channels indicates the location of R3 on the microtubules (lower row). Control experiments (Fig. S2D) indicated that subpellicular microtubules from the Δldpκar3 mutants showed no background fluorescence with antibody against LdPKAR3.

**Figure 8: Deleting the genes encoding R3 or C3 leads to promastigotes rounding.** Mid-log culture of WT, Δldpκar3, Δldpκar3 L. donovani promastigotes and their respective add-backs were subjected to image streaming (Amnis® ImageStream®X Mk II). A) A graph illustrating percentage of elongated (■) and round (■) cells. The number of ovoid cells was around 30% in all cell lines and therefore is not presented. B) Bright field (upper row) and anti-membrane immunofluorescence of the cell lines from (A). * indicates statistical significance of p < 0.05 (significantly different from WT) and ** indicates p < 0.01 (significantly different from WT), n = 3. The bars in (B) indicates 5 µm.
STAR Methods

_Phylogenetic analysis of PKA regulatory subunits._ The amino acid sequences of the PKAR orthologues from 39 species of Euglenozoa and 14 other eukaryotic species, including at least one in each of the Amoebozoa, Apusozoa, Archaeplastida, Excavata, Opisthokonta and SAR (Stramenopiles/Alveolata/Rhizaria) supergroups, were downloaded from the TriTrypDB, GenBank or Euglena International Network (EIN) databases (see Supplementary Table S2). The sequences were aligned using the MAFFT module of Geneious Prime 2021.1.1 (https://www.geneious.com) and a phylogenetic tree constricted using the RAxML (Randomized Axelerated Maximum Likelihood) version 8.2.11 module, implemented in rapid hill-climbing mode with joint branch length optimization and the BLOSUM62 Substitution Matrix. A GAMMA model of rate heterogeneity was used for ML estimate of alpha-parameter from 100 distinct randomized maximum parsimony (MP) trees. Protein domains were identified using the InterProScan module of Geneious and/or the HHpred website of the MPI Bioinformatics Toolkit.

_Cells and culture._ L. donovani MHOM/SD/00/1S axenic promastigotes were grown in Earle’s-based medium 199 (M199; Biological Industries, Ltd.), supplemented with 10% heat-inactivated fetal bovine serum (FBS; GIBCO, Ltd) and 1% penicillin-streptomycin solution (Biological Industries, Ltd.). Axenic differentiation of _L. donovani_ promastigotes to amastigotes was stimulated as described previously. Briefly, mid-log phase promastigotes were washed twice in EARL’s salt solution and finally suspended in amastigote medium containing M199 at pH 5.5, supplemented with 0.5 mM sodium succinate, 25% fetal calf serum and 1% penicillin-streptomycin solution (differentiation medium). Mature amastigotes developed within 5 days of exposure to the differentiation medium. Trypanosomes were cultivated at 37°C and 5% CO₂, in
modified HMI-9 culture medium, supplemented with 10% heat-inactivated FBS. Cell density was maintained below 1× 10^6 cells/ml by regular dilution.

**Cloning.** PCR products and the plasmid vectors (see Table S1) were digested by restriction enzymes for 30 min, at 37°C (FastDigest enzymes, Thermo Scientific) and ligated (T4 DNA ligase, Thermo Scientific) for up to 1 h, at 22°C. Ligation products were transformed into *E. coli* DH5α by heat-shock (42°C), which were then incubated for 1 h in Luria broth (LB) medium at 37°C and plated on LB-agar plates supplemented with ampicillin. LB-agar plates were incubated at 37°C overnight. Colonies were screened using PCR or the colony cracking method. Positive colonies were grown overnight in LB medium supplemented with ampicillin, and plasmids were extracted using a mini-prep kit (GeneJet Plasmid Miniprep Kit, Thermo Scientific). Extracted plasmids were sequenced to assure that no mutations were inserted during PCR or the cloning procedure.

**Generation of a polyclonal anti-PKAR3 antibody.** The full-length CDS of LdPKAR3 (encoded by *LinJ.34.2680*) was PCR-amplified from genomic DNA of *L. infantum* strain JPCM5 using primers (Table S1) designed to introduce an N-terminal His6 tag followed by a TEV protease cleavage site, and cloned via *Bam*HI and *Not*I restriction sites, into pETDuet-1 (Novagen, Merck Millipore). The protein was expressed in *E. coli* Rosetta and purified using Ni-NTA columns, followed by TEV protease cleavage. Recombinant LdPKAR3 (5 mg/ml) was injected into two rabbits. Rabbits were boosted four times with 5 mg/ml recombinant LdPKAR3. Injections and bleedings were carried out by a contracted service (Sigma-Aldrich, Ltd).

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1 To maintain consistency with our previous publications, we use the original GeneIDs for the *L. infantum* JPCM5 reference genome in TryTrypDB, rather than those for any of the *L. donovani* genomes.
Western blotting. Western blot analysis was performed as described previously\textsuperscript{64}, using a 1:1,000 dilution of polyclonal rabbit anti-LdPKAR3, -TbPKAR1\textsuperscript{7}, -TbPKAC1/C2\textsuperscript{19} and TbSAXO\textsuperscript{30} antibodies. Rabbit antibodies against TbPKAC3 were used at 1:250 dilution\textsuperscript{19}. For Western blot detection of proteins recombinantly expressed in \textit{L. tarentolae}, the primary antibodies mouse anti-His (Bio-Rad) and mouse anti-Strep (Qiagen) were used. The secondary antibodies IRDye680LT goat anti-rabbit (1:50000; LICOR) and IRDye800CW goat anti-mouse (1:10000; LICOR) were used for detection with the Odyssey\textsuperscript{TM} CLx imaging system (LICOR). Rabbit antiserum against \textit{L. donovani} HSP83\textsuperscript{11} or β-tubulin (cell signalling #2146) were used as a protein loading marker.

Gene knockouts. The gene encoding LdPKAR3 was deleted from the 1S-2D clone of \textit{L. donovani} using homologous recombination, as described previously by Inbar et al.\textsuperscript{33}. PCR analyses of genomic DNA were performed to confirm the absence of the targeted gene (\textbf{Fig. S1B}). Polyclonal antibodies raised in rabbits against recombinant LdPKAR3 reacted with a protein of the correct molecular mass (72 kDa) in Western blots of wild type cells, but not \textit{Aldpkar3} mutants (+/+ and -/-, respectively in \textbf{Fig. S1A}). The protein was present (at slightly elevated levels) in add-back mutants (-/-/+ectopically expressing full-length LdPKAR3.

\textit{LinJ.18.1090} (which encodes LdPKAC3) was deleted from the 1S-2D clone of \textit{L. donovani} using the same strategy as above\textsuperscript{33}. PCR analyses of \textit{L. donovani} genomic DNA extracted from single colonies confirmed that both copies of the gene were missing from \textit{Aldpkac3}, while the antibiotic resistance genes that were inserted to replace LdPKAC3 were in place (\textbf{Fig. S4A and B}).

\textit{Inducible overexpression of LdPKAR3 in Trypanosoma brucei.} The LdPKAR3 CDS was PCR-amplified from genomic DNA of \textit{L. donovani} promastigotes using primers
LdPKAR3_HindIII_fw and LdPKAR3_lr_Ty1_BamHI_rev (Table S1) to introduce a C-terminal Ty1 tag. The PCR product was cloned into the HindIII and BamHI sites of pHD615, which was then linearized with NotI for transfection of a previously described homozygous T. brucei PKAR deletion mutant that additionally expresses a double tetracycline repressor (pHD1313). Transfected cells were selected with 2 µg/ml blasticidin (pHD615) and 2.5 µg/ml phleomycin (pHD1313). LdPKAR3 expression was induced with 1 µg/ml tetracycline.

**Immunofluorescence** Indirect immunofluorescence analysis was carried out following the methods described by Inbar et al. Briefly, mid-log phase promastigotes were washed twice in phosphate buffered saline (PBS) and then fixed in 1% formaldehyde/PBS on a slide for 10 min before permeabilization by exposure to 0.2% TritonX-100/PBS for 10 min. Cells were incubated with blocking solution [5% (v/v) non-fat dried skimmed milk powder/PBS-Tween (PBST)] for 30 min at room temperature, incubated with rabbit anti-LdPKAR3 (this study), rabbit anti-promastigote membrane proteins or mouse anti-HA tag antibodies (1:500 dilution) for 1 h. Subsequently, slides were washed three times with PBST and incubated with the fluorescent secondary antibody in the dark for 30 min. Slides were washed three times with PBST and a drop of DAPI Fluoromount G (Southern Biotech Ltd) was added. Slides were covered with slips, sealed, and then stored in the dark. Subsequently, they were examined using a Zeiss LSM 700 inverted confocal laser scanning microscope. Image processing was done using Zen Lite software, Zeiss. Control experiments (Fig. S2D) indicated that subpellicular microtubules from the ∆ldpkar3 mutants showed no background fluorescence with antibody against LdPKAR3.

**Image streaming.** Image Streaming analysis was carried out using Amnis® ImageStream®X Mk II. The Image Streamer was calibrated by feeding it with late log phase axenic promastigotes and amastigotes. For each cell, a snapshot was taken, and the
aspect ratio was calculated. The streamer divided these populations into three groups: elongated (Fig. S5A top, B top left), round (Fig. S5A bottom, B bottom), and intermediate shape, which we refer to as ovoid (Fig. S5A top and bottom, B top right). Note that round promatigotes are flagellated, whereas round amastigotes are aflagellated. These shapes were used as templates to analyze 10000 cells of each experimental group.

*Detergent-based enrichment of subpellicular microtubules.* Axenic *L. donovani* promastigote and parasites at various time points after exposure to the differentiation signal (20 ml of cells at 1×10^7 cells/ml), were washed twice with ice-cold PBS. The final pellet was suspended in 200 µl PBS containing protease inhibitors mix (Roche cOmplete protease inhibitor) and 0.5% n-dodecyl β-D-maltoside (DDM) in Eppendorf tubes. After a short vortex, the tubes were placed on ice for 15 min, and subsequently centrifuged (14,000 RPM, 15 min, 4°C). The pellet was washed twice with 1 ml of ice-cold PBS containing the protease inhibitors. For immunofluorescence analysis, the pellet was suspended in 100 µl PBS. For Western blot analysis, the pellet was suspended in 100 µl Laemmli buffer. *T. brucei* subpellicular microtubules were enriched employing the same protocol used for *Leishmania*.

*Immunoprecipitation.* Immunoprecipitation of LdPKAR3-Ty1 was performed by binding anti-Ty1 to magnetic protein A beads (Dynabeads, Invitrogen) followed by a 2-hour incubation with 1×10^8 trypanosomes lysed in lysis buffer (10 mM Tris/Cl pH 7.5; 150 mM NaCl; 0.5 mM EDTA; 0.5% NP-40; Roche cOmplete protease inhibitor) for 30 min at 4°C. Beads were washed 4x with lysis buffer and proteins were eluted by incubation with 50 µl 2x Laemmlsi sample buffer, for 5 min, at 95°C.

*Tandem affinity purification from Leishmania tarentolae.* LdPKAR1 and LdPKAR3 were N-terminally tagged with a hexa-histidine peptide by PCR amplification from *L. donovani*.
genomic DNA. The PCR products were inserted in the pLEXSY_I-ble3® vector (Jena Bioscience). LdPKAC3 was N-terminally strep-tagged and inserted into pLEXSY_I-neo3®. The vectors were then linearized with SwaI and stably co-transfected for holoenzyme expression (LdPKAR1 and LdPKAC3 or LdPKAR3 and LdPKAC3) into the LEXSY T7-TR® cell line (Jena Bioscience). Protein co-expression, parasite cultivation and protein purification were performed as described by Bachmaier et al.19 with a few modifications. Briefly, co-expression was induced with 10 µg/ml tetracycline for 48 h. The cells were harvested by centrifugation (2000Xg for 5 min), washed with PBS, and lysed in His binding buffer (50 mM NaH2PO4 pH 8, 300 mM NaCl, 10 mM imidazole, 1% Triton-X, protease inhibitor cocktail). The soluble fraction was loaded onto a gravity flow Ni-NTA column (Thermo Fisher Scientific). The column was washed twice with His wash buffer (50 mM NaH2PO4 pH 8, 300 mM NaCl, 20 mM imidazole) before elution of the kinase complex with His elution buffer (50 mM NaH2PO4 pH 8, 300 mM NaCl, 250 mM imidazole). The Ni-NTA eluate was loaded on to a gravity flow Strep-Tactin column (IBA), which was then washed with Strep wash buffer (50 mM NaH2PO4 pH 8, 300 mM NaCl). Lastly, the kinase complexes were eluted with Strep elution buffer (50 mM NaH2PO4 pH 8, 150 mM NaCl, 50 mM biotin). The integrity of the co-expressed holoenzyme complexes was analysed by western blot.

Fluorescence Resonance Energy Transfer (FRET). Mid-logarithmic phase L. donovani promastigotes ectopically expressing LdPKAC3-HA were fixed in PBS containing 4% paraformaldehyde, and allowed to settle on slides. LdPKAR3 was labelled using polyclonal anti-R3 antibody (1:500), followed by detection with goat anti-rabbit Alexa 568 antibody (1:500). LdPKAC3 was labelled with anti-HA antibody (1:1000, BioLegend#MMS-101P) and secondary goat anti-mouse Alexa 647 antibody (1:500). Confocal microscopy was performed using a Zeiss
LSM 700 inverted confocal laser scanning microscope. An excitation wavelength of 578 nm and an emission wavelength of 640 nm and below were used for Alexa568, whereas an excitation wavelength of 651 nm and an emission wavelength of 640 nm and above were used for Alexa647. FRET was assessed with Axiovision FRET software, using the mathematical approach - Youvan's technique. Youvan's technique (Fc) is the basis of all correction measurement techniques (out of 4 techniques) in Axiovision FRET. It refers to corrected FRET values when three FRET filter sets are used. The measured values were corrected for the background (bg) and for the crosstalk from the donor (don) and the acceptor (acc). Calculation of the intensity values for FRET follow the formula: Fc = (fretgv - bg) - cfdon (dongv - bg) - cfacc (accgv - bg), where gv is the fluorescence value in every pixel and cf is a constant which is calculated for each of the samples with only one secondary antibody.

*Pulldown of subpellicular microtubule-bound proteins (proteomics).* To determine the levels of LdPKAC3 binding to the subpellicular microtubules of *L. donovani* wild type (WT) and *Aldp*kar3, microtubules of WT and *Aldp*kar3 promastigotes, before and 5 h after exposure to a differentiation signal, were enriched as described above. Subsequently, proteins associated with this fraction were separated by 9% SDS-PAGE and a slice containing proteins at the molecular mass range of 35-42kDa was excised and subsequently subjected to mass spectrometry.

The proteins in the gel were reduced with 3 mM DTT (60ºC, 30 min), modified with 10 mM iodoacetamide in 100 mM ammonium bicarbonate (in the dark, room temperature, 30 min) and digested in 10% acetonitrile and 10 mM ammonium bicarbonate with modified trypsin (Promega) at a 1:10 enzyme-to-substrate ratio, overnight at 37ºC. The resulting peptides were desalted using C18 tips (Homemade stage tips) dried and resuspended in 0.1% formic acid.
The peptides were resolved by reverse-phase chromatography on 0.075 X 180-mm fused silica capillaries (J&W) packed with Reprosil reversed phase material (Dr Maisch GmbH, Germany). The peptides were eluted with linear 60 min gradient of 5-28%, 15 min gradient of 28-95% and 15 min at 95% acetonitrile with 0.1% formic acid in water at flow rates of 0.15 μl/min. Mass spectrometry (MS) was performed with a Q Exactive plus mass spectrometer (Thermo) in a positive mode, using repetitively full MS scan, followed by collision induced dissociation (HCD) of the 10 most dominant ions selected from the first MS scan.

The MS data were analyzed using Proteome Discoverer 1.4 software with Sequest (Thermo) algorithm against *L. donovani* and *L. infantum* proteomes from TriTryp database with mass tolerance of 20 ppm for the precursor masses and 0.05 Da for the fragment ions (TriTrypDB). Oxidation on Met, and phosphorylation on Ser, Thr, Tyr were accepted as variable modifications and carbamidomethyl on Cys was accepted as static modifications. Minimal peptide length was set to six amino acids and a maximum of two miscleavages were allowed. Peptide- and protein-level false discovery rates (FDRs) were filtered to 1% using the target-decoy strategy. Semi-quantitation was done by calculating the peak area of each peptide based on its extracted ion currents (XICs) and the area of the protein that is the average of the three most intense peptides from each protein.

The MS proteomics data were deposited in the ProteomeXchange Consortium via the PRIDE\textsuperscript{68} partner repository with the dataset identifier PXD025222.

**Molecular modelling.** LdPKAR3 homology models were built from each of the PDB templates 2QCS, 5JR7 and 4MX3 with Chimera Modeller\textsuperscript{69,70}. The pairwise alignments used as input for threading PKAR3 and PKAC3 were extracted from multiple sequence alignments that included PKAR1 and PKAC1/C2, respectively, and were produced with Clustal Omega\textsuperscript{71}. The
R3/C3 complex was obtained by docking C3 into R3 using the PDB template 2QCS as a guide, followed by energy minimization. Structure images were produced with UCSF Chimera.\textsuperscript{72}
Supplementary figure legends

Figure S1: Deleting the LdPKAR3 gene (A) and raising rabbit antibodies against LdPKAR3 protein. (B) PCR on genomic DNA extracted from promastigotes of WT (+/+) and a Δldpkar3 (-/-). LdPKAR3 ORF (upper panel) is seen only on DNA extracted from WT cells, while the mutant contains both the resistance, hygromycin and neomycin genes (middle and bottom panels, respectively). (B) Proteins extracted from L. donovani WT (+/+; left lane), Δldpkar3 (-/-; middle lane) and Δldpkar3 ectopically expressing the full-length LdPKAR3 (-/-/+; addback; right lane) were separated by 10% SDS-PAGE and subsequently subjected to Western blotting using the antibodies raised against LdPKAR3. Tubulin was used as a loading control.

Figure S2: A-B subpellicular microtubule enrichment using 0.5 % DDM (A) The DDM-insoluble fraction was enriched with tubulin. Western blot of promastigote whole-cell lysate (left lane) and 0.5% DDM-insoluble fraction followed by Ponceau red staining. Anti-β tubulin was used as a loading control. (B) DDM-insoluble subpellicular microtubules of axenic promastigotes (left panel) and amastigotes (right panel) were stained with anti-β tubulin mouse antibodies (red). (C) Purity assessment of the DDM-insoluble fraction from L. donovani promastigotes. Whole-cell lysates (left lane) and DDM-insoluble fraction were stained with anti-AAP24 (proline transporter, upper panels) and anti-HSP30 antibodies (lower panels). (D) Rabbit anti-LdPKAR3 antibody specificity in WT (upper panel) and Δldpkar3 mutant (lower panel).

Figure S3: FRET control experiment. To determine the fluorescence level of each of the Alexa Fluor dyes that were coupled to the secondary antibodies used in the analysis described in Fig. 5C, (A) only LdPKAR3 was stained with Alexa Fluor 568 (in green) or (B) only ectopic...
LdPKAC3 (HA-tagged) was stained with Alexa 647 (in red). FRET analysis was carried out using Axio Vision program (STAR Methods).

**Figure S4: Deleting LdPKAC3 from the L. donovani genome.** PCR of genomic DNA extracted from wild type (+/+) (A) or (B) Δldpkac3 (-/-) cells. The PKAC3 ORF was detected only in wild type cells, while Δldpkac3 cells possessed both resistance marker cassettes; hygromycin and neomycin.

**Figure S5. Image streamer calibration with cell shape of L. donovani promastigotes and amastigotes.** An Amnis® ImageStream®X Mk II image streamer was fed with either late log phase axenic promastigotes (A, upper panel) or axenic amastigotes (A, lower panel). For each life stage, 10000 cells were tested, an image snapshot was taken (B) and subsequently an aspect ratio was calculated. The streamer divided these populations into three groups, elongated (marked in orange), round (red) and ovoid (pink). Note that round promastigotes are flagellated, whereas amastigotes are aflagellated round.
## Supplemental Tables

### Table S1: Primers used in this study

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### Primer sequences

Uppercase letters denote sequence that matches the target, while lowercase letters represent sequences added to incorporate restriction sites (underlined) and epitope tags.

LdPKAR3_CDS-Fwd: gccagatccggaaacctgtatatctcaggatctATGTCCAGTTTCGAC
LdPKAR3_CDS-Rev: attatcgccgctCTACGCGCTCTCGTG
LdPKAC3_5’Fwd1: gtagctggctgTCGCCAGGCAGAGCG
LdPKAC3_5’Rev1: cttgaagctTGGTGTCTATATTGACTGTAAGG
LdPKAC3_3’Fwd1: gacgctggtTGATTTCTGCCTCCAAGG
LdPKAC3_3’Rev1: cgtacgtagAGAATGTGCGTTGATG
LdPKAC3_CDS-Fwd2: GCGCGAGCTTCAAGATGG
LdPKAC3_CDS-Rev2: GCGGTAACGATCGTCTTG
LdPKAR3_HindIII_fw: gccgaagctgATGTCCAGTTTCGAC
LdPKAR3_lrt_1 BamHI rev:
gcgacggcatctcgtaaatctgtggtaatggtggacctcagatcCGCGCTCTCGTGTTG
LdPKA_CDS-Fwd1: gacctacattacgtaaatctgtaaatctccagacgatcATGACCAACCGAAAG
LdPKA_CDS-Rev1: ctgacgtacCTACTCGTCCGTATACCTTGCC
PKAR' 5' UTR fr: aatctGTCAAGCgcatcattcgtaattc
pkar' 5' UTR rev: aatctAAGCTTaaagggcaaggacagg
PKAR' 5' UTR fr: aatctGGTACcatcgtaattctetec
PKAR' 3' UTR fr: aatctGGATCCagccacagttctacc
PKAR' 3' UTR rev: aatctTCTAGAaagcttctctcctcct
PKAR' fr: AATCAGGTACAGCAACTTTCAATACTACG
PKAR' rev: AATCAGGTACCCCTACGCGCTCTCGTGTTG
Hyg-Rev1: AATGTCAAGCAGTGTCGGTCCG
Neo-Rev1: CAGCTGCGCAGAGGACGC
Upstream to PKAC3 5': GGTGCTCTGGGCTTCGGCAG
Table S2: Organisms included in PKAR phylogenetic analysis
Bibliography


555–566.


42. Barros, E.P., Malmstrom, R.D., Nourbakhsh, K., Del Rio, J.C., Kornev, A.P., Taylor, S.S.,


Cellular landmarks of *Trypanosoma brucei* and *Leishmania mexicana*. Mol Biochem Parasitol. 230, 24-36.


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Figure 5

A

Cell lysate

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<th>Wild type+/+</th>
<th>ΔR3</th>
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| HA* (LdPKA-C3) |
| LdPKA-R3 |

DDM-enriched

| WT | ΔR3 |

| β tubulin |

B

Relative abundance

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C

Image of cells with green and black background.
Figure 6

A

Bovine PKA-RI

Leishmania PKA-R3

B

C

D

Bovine PKA-RI

Leishmania PKA-R3

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Figure 7

α LdPKAR3

β-tubulin

Merged