

Real-time dynamics of *Plasmodium* NDC80 reveals unusual modes of chromosome segregation during parasite proliferation

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Abstract

Eukaryotic cell proliferation requires chromosome replication and precise segregation to ensure daughter cells have identical genomic copies. The genus *Plasmodium*, the causative agent of malaria, displays remarkable aspects of nuclear division throughout its lifecycle to meet some peculiar and unique challenges of DNA replication and chromosome segregation. The parasite undergoes atypical endomitosis and endoreduplication with an intact nuclear membrane and intranuclear mitotic spindle. To understand these diverse modes of *Plasmodium* cell division, we have studied the behaviour and composition of the outer kinetochore NDC80 complex, a key part of the mitotic apparatus that attaches the centromere of chromosomes to microtubules of the mitotic spindle. Using NDC80-GFP live-cell imaging in *Plasmodium berghei* we observe dynamic spatiotemporal changes during proliferation, including highly unusual kinetochore arrangements during sexual stages. We identify a very divergent candidate for the SPC24 subunit of the NDC80 complex, previously thought to be missing in *Plasmodium*, which completes a canonical, albeit unusual, NDC80 complex structure. Altogether, our studies reveal the kinetochore as an ideal tool to investigate the non-canonical modes of chromosome segregation and cell division in *Plasmodium*.

Keywords:

Malaria, *Plasmodium*, kinetochore, NDC80 complex, chromosome segregation, atypical cell division, endomitosis, meiosis.

Summary Statement

The dynamic localization of kinetochore marker NDC80 protein complex during proliferative stages of the malaria parasite life cycle reveals unique modes of chromosome segregation.

Introduction

Mitosis and meiosis are fundamental processes in cell division that enable DNA replication and chromosome segregation, and allow eukaryotic organisms to proliferate, propagate and survive. During these processes, microtubular spindles form to facilitate an equal segregation of duplicated chromosomes to the spindle poles. Chromosome attachment to spindle microtubules (MTs) is mediated by kinetochores, which are large multi-protein complexes assembled on centromeres located at the constriction point of sister chromatids (Cheeseman, 2014; McKinley and Cheeseman, 2016; Musacchio and Desai, 2017; Vader and Musacchio, 2017). Each sister chromatid has its own kinetochore, oriented to facilitate movement to opposite poles of the spindle apparatus. During anaphase, the spindle elongates and the sister chromatids separate, resulting in segregation of the two genomes during telophase. The NDC80 complex is the major component of the kinetochore and mediates its attachment to spindle MTs. In most model organisms, it is a member of the network of conserved Knl1, Mis12 and NDC80 complexes (KMN) (McKinley and Cheeseman, 2016; Petrovic et al., 2016). The ~170-190 kDa NDC80 complex has two globular domains at either end of a ~57 nm elongated coiled-coil, forming a dumb-bell shape. It is a heterotetramer comprising a 1:1:1:1 ratio of NDC80 (also known as HEC1 in humans), NUF2, SPC24 and SPC25 sub-complexed as two heterodimers: NDC80 with NUF2 and SPC24 with SPC25 (Ciferri et al., 2005; Farrell and Gubbels, 2014; Wei et al., 2005). The C-terminal end of the SPC24-SPC25 dimer anchors the complex to the kinetochore; whereas the NDC80-NUF2 dimer mediates plus-end MT binding through its calponin homology domain (CHD) (Alushin et al., 2010; Farrell and Gubbels, 2014; Sundin et al., 2011).

Malaria, caused by the apicomplexan parasite *Plasmodium* spp., remains one of the most prevalent and deadly infectious diseases worldwide, with 219 million clinical cases and 435,000 deaths in 2017 (WHO, 2018). *Plasmodium* has several morphologically distinct proliferative stages during its life cycle that alternates between vertebrate host and mosquito vector (**Fig.1**) (Francia and Striepen, 2014; Sinden, 1991a; Sinden, 1991b). A malaria parasite-infected female anopheles mosquito inoculates haploid sporozoites into the mammalian host during a blood meal. Sporozoites travel through the blood stream to the liver and infect hepatocytes, where the parasite replicates and develops into a multinucleated schizont. At the end of this exo-erythrocytic schizogony the host cell is ruptured to release haploid merozoites, which infect erythrocytes. In the intra-erythrocytic phase, an initial ring stage form develops into a trophozoite and then into a schizont where multiple rounds of asexual multiplication occur (erythrocytic schizogony). At the end of schizogony, host cell rupture releases further merozoites that infect new erythrocytes.

Following erythrocyte invasion, some parasites differentiate into male (micro) and female (macro) gametocytes to initiate the sexual phase of the life cycle, which occurs inside the mosquito. These haploid parasites are arrested at the G0/G1 phase of the cell cycle (Arnot and Gull, 1998). Ingestion by a mosquito activates gametogenesis. Male gametogenesis is very rapid with three rounds of genome replication from 1N to 8N and the release of eight motile haploid microgametes within 15 min. The activated female gametocyte rounds up and the macrogamete egresses from the red blood cell. Gametogenesis can be studied *in vitro* using a culture medium that mimics the mosquito midgut environment (Billker et al., 1998; Tewari et al., 2005).

After fertilisation the zygote differentiates into a motile ookinete. The ookinete invades the mosquito midgut wall where it develops into an oocyst. At this stage multiple rounds of endomitotic division occur in a process similar to schizogony, which is followed by cytokinesis to form thousands of motile sporozoites (Francia and Striepen, 2014; Gerald et al., 2011). The sporozoites are released from the oocyst and migrate to the mosquito's salivary glands for transmission to the vertebrate host.

The life cycle of *Plasmodium* is characterized by two unique processes of mitosis and a single stage of meiosis. The first mitotic process occurs during schizogony within mammalian hepatocytes and erythrocytes, and during sporogony in oocysts in the mosquito (Sinden, 1991a; Sinden, 1991b) (**Fig. 1 A**). This mitotic division is atypical, for example no clear G2 cell cycle phase has been observed during schizogony (Arnot and Gull, 1998; Doerig et al., 2000). Furthermore, this asexual proliferation is characterised by multiple rounds of asynchronous nuclear division without chromosome condensation and in the absence of cytokinesis. Mitosis is closed, occurring without dissolution and reformation of the nuclear envelope, and the spindle-pole body (SPB)/microtubule-organising centre (MTOC), also known as the centriolar plaque (Arnot et al., 2011; Francia et al., 2015; Sinden, 1991a), is embedded within the nuclear membrane. The asynchronous nuclear division precedes cell division, leading to a multinucleate syncytium. The last round of nuclear division in these cells is synchronous and it is only after this final round of mitosis that cytokinesis occurs to form the haploid daughter merozoites or sporozoites, respectively.

The second type of mitotic division occurs during male gametogenesis following activation in the mosquito midgut (**Fig. 1 B**). Three rounds of rapid genome duplication (from haploid to octoploid) without concomitant nuclear division (endoreduplication) are followed by chromosome condensation and nuclear budding into the male gametes during exflagellation, all within 12 to 15 min of activation (Arnot and Gull, 1998; Janse et al., 1988; Sinden, 1983).

The resultant eight flagellated microgametes each contain a haploid genome (Guttery et al., 2015; Sinden et al., 2010). Fertilization of the female gamete results in a diploid zygote, which develops in the mosquito gut and differentiates over a 24-hour period into a motile ookinete (**Fig. 1 C**). It is in this stage that meiosis occurs. The DNA is duplicated once to form a tetraploid cell and then two rounds of chromosome segregation result in four discrete haploid genomes prior to nuclear division and ookinete maturity. Reductive division to haploidy presumably occurs in the subsequent oocyst during sporozoite formation (Guttery et al., 2015; Sinden, 1991a; Sinden, 1991b). Collectively, these different stages of cell division and proliferation indicate that the parasite has evolved alternate modes of chromosome replication, condensation and segregation, as well as nuclear and cell division at different stages during its life cycle.

The process of chromosome segregation and associated kinetochore dynamics, which is the key role of the mitotic apparatus throughout the life cycle, is not well understood in *Plasmodium*. To date, analysis of *Plasmodium* mitotic/meiotic spindle assembly and chromosome segregation has been performed largely using transmission electron microscopy (TEM) (Sinden et al., 1978; Sinden et al., 1976), and biochemical analysis of microtubule markers including α-tubulin (Fennell et al., 2008), and centrin associated with the putative MTOC (Gerald et al., 2011; Roques et al., 2019). An analysis of a *Plasmodium* artificial chromosome (PAC) identified a putative centromere derived from chromosome 5 (*PbCEN5*), and highlighted the dynamics of chromosome segregation during both mitotic and meiotic stages of the parasite's life cycle (Iwanaga et al., 2010). However, there is no real-time analysis of chromosome segregation dynamics during the various proliferative stages, especially during stages that occur inside the mosquito vector. Here, we have analysed the real-time expression and spatiotemporal dynamics of NDC80, as a kinetochore marker. We generated a stable transgenic *P. berghei* line expressing NDC80 with a C-terminal GFP-tag by modifying the endogenous gene locus. Using this tool, we examined NDC80 expression and location to follow the spatiotemporal organisation of outer kinetochores during mitosis in schizogony, sporogony and male gametogenesis, and during meiosis in ookinete development. We observed unusual kinetochore dynamics as patterns of clustered foci adjacent to the nuclear DNA during endomitotic nuclear division in asexual stages, with the number of foci corresponding to the likely ploidy of individual nuclei. However, kinetochores formed an unusual bridge-like pattern during endoreduplication stages in male gametogenesis. We show there is likely a full complement of NDC80 complex subunits, with a highly divergent candidate for the previously undetected SPC24 subunit identified by a combination of proteomics and sensitive comparative sequence analysis. This transgenic parasite line expressing GFP-tagged NDC80 is a valuable resource for studying chromosome

segregation and dynamics, as well as identifying and characterising the various protein complexes involved in these cell division processes.

Results

To study kinetochore dynamics, we generated NDC80-GFP, a transgenic *P. berghei* line expressing NDC80 with a C-terminal GFP tag, by inserting an in-frame *gfp* coding sequence at the 3' end of the endogenous *Ndc80* locus using single homologous recombination (**Fig. S1A**). To complement our GFP-based imaging studies, we also generated a NDC80-mCherry line using the same strategy (**Fig. S1A**). Successful insertion was confirmed by PCR (**Fig. S1B**). Western blot analysis of a schizont protein extract using an anti-GFP antibody revealed the NDC80-GFP protein at the expected size of 96 kDa compared to the 29 kDa GFP alone (**Fig. S1C**).

Following successful generation of the NDC80-GFP transgenic line, the spatiotemporal profile of NDC80-GFP protein expression and location was examined during the parasite life cycle at the three asexual mitotic replicative stages (liver and blood schizogony in the vertebrate host and oocyst development (sporogony) in the mosquito vector) (**Fig. 1A**), the sexual mitotic stage (male gametogenesis) (**Fig. 1B**) and the meiotic stage (ookinete development) (**Fig. 1C**).

Real-time live cell imaging using NDC80-GFP reveals kinetochores aggregate as discrete foci during schizogony

In the asexual blood stage, no NDC80-GFP fluorescence was observed in the intra-erythrocytic ring stage, a non-replicative G1 phase of the cell cycle (Arnot and Gull, 1998; Arnot et al., 2011; Doerig et al., 2000) (**Fig. 2A**). A faint but discrete single focus of NDC80-GFP adjacent to the nuclear DNA was observed in the early trophozoite, which became more intense as the trophozoite developed (**Fig. 2A**). The late trophozoite stage marks the transition into early S phase of the cell cycle, when DNA synthesis starts. The NDC80-GFP focus then split into two foci that migrated away from each other but remained attached to the nuclear DNA (stained using Hoechst) that then separated into two nuclear masses. This is consistent with the separation of sister chromatids (anaphase) and then the first nuclear division (telophase) that marks the onset of schizogony. These observations indicate that kinetochores are grouped together in a tight focus throughout these mitotic stages of nuclear replication. Similar fluorescence patterns were also observed in the NDC80-mCherry line (**Fig. S2**). NDC80-GFP also revealed the asynchronous nature of nuclear division during early schizogony, as displayed by two or more nuclei with single and double distinct NDC80 foci

concurrently (e.g. **Fig. 2A**, Sch-E). As alternating repeated S/M phases following the division of individual nuclei continued, these NDC80-GFP foci were duplicated several times into multiple foci and nuclei. Further analysis of NDC80-GFP localization by super resolution microscopy confirmed the asynchronicity of nuclear division during blood stage schizogony (see **Fig. 2B**, **Fig. S3A, B, C** and **Supplementary videos SV1, SV2, SV3**). This stage of DNA replication and nuclear division concludes with cytokinesis to produce haploid daughter merozoites that egress from the erythrocyte (Arnot and Gull, 1998; Doerig et al., 2000). The short-lived, extracellular merozoite represents part of the G1 phase of the cell cycle (Arnot and Gull, 1998; Doerig et al., 2000). During schizogony in the pre-erythrocytic asexual stage in the liver, discrete fluorescent foci next to nuclear DNA were observed, which is similar to the pattern observed in blood stage schizogony (**Fig. S3D**).

To complement the live-imaging analysis, erythrocytic schizogony was examined in ultrastructural studies. In the multinucleated schizont during merozoite formation, it was possible to identify nuclear/spindle poles directly adjacent to the nuclear envelope with radiating microtubules and attached kinetochores on the inside of the nucleus (**Fig. 2C**). These structures were not seen in the mature merozoite or the early intracellular ring stage. These observations are consistent with the live-imaging data.

Immunofluorescence imaging shows the arrangement of NDC80 (kinetochore) with centrin (putative MTOC/SPB) and α-tubulin (spindles)

To determine the relative position of NDC80-GFP with other mitotic markers, especially at the spindle and/or putative MTOC/SPB, we used immunofluorescence-based co-localization assays with anti-GFP antibodies, anti-α-tubulin and anti-centrin, as markers for the spindle and putative MTOC/SPB, respectively. We observed that NDC80-GFP is located adjacent to alpha-tubulin showing some overlap between the NDC80 and alpha-tubulin signals in most early schizonts (Sch-E) but not in late schizonts (Sch-L) (**Fig. 2D**). Similarly, using anti-GFP with anti-centrin antibodies revealed that NDC80-GFP is located in close proximity to, but does not co-localize with, centrin (**Fig. 2E**)

Rapid spindle dynamics of NDC80-GFP shows unusual kinetochore bridges during endoreduplication in male gametogenesis

Given the remarkable speed and organisation of nuclear replication and chromosome segregation during male gametogenesis, we investigated the live cell dynamics of NDC80-GFP throughout this 15 min process following gamete activation *in vitro*. The results are presented in **Fig. 3A**, and include time-lapse screenshots (**Fig. 3B and C, Supplementary**

videos SV4 and SV5.). In non-activated male gametocytes a single diffuse and faint NDC80-GFP focus was present (**Fig. S4A**), which intensified to a sharp single focal point 1 minute post-activation (mpa) (**Fig. 3A**). By 2 mpa, this focal point extended to form a bridge across one side of the nucleus, followed by the separation of the two halves of the bridge to produce two shorter linear rods that then contracted to two clear single foci by 3 mpa (**Fig. 3A-C**). A schematic diagram for this process (1-3 mpa) is shown in **Fig. 3D**. This unusual linear arrangement of NDC80 shows that the distribution of kinetochores extends to approximately the full width of the nucleus, often arched around the nuclear margin, and maintains a consistent thickness of NDC80-GFP signal, suggesting that kinetochores are evenly spaced along this single linear element. This process was repeated twice, although non-synchronously, resulting in 8 discrete NDC80-GFP foci (**Fig. 3A**). To study the association of NDC80 with the spindle we used immunofluorescence-based co-localization assays with anti-GFP antibodies, and anti- α -tubulin. This showed clear co-localisation of NDC80 with the microtubule marker, both on the bridge-like structure and the foci for NDC80 (**Fig. 3E**)

Ultrastructural analysis of the nucleus during male gametogenesis within 8 mpa showed typical nuclear spindles with microtubules radiating from the nuclear poles, to which attached kinetochores could be identified dispersed along the length of the spindle (**Fig. 3F**). The spindles with attached kinetochores were located on the inner side of the nuclear membrane, (**Fig. 3F**). TEM images showed that these kinetochores are dispersed along the length of the mitotic spindle from one spindle pole to the other (similar to the bridge observed in the fluorescence microscopy). These observations are consistent with the fluorescence microscopy, and this kinetochore configuration is different to that in canonical metaphase, in which there is a central metaphase plate perpendicular to the spindle axis.

Following exflagellation, the eight discrete foci associated with endoreduplication disappeared rapidly and no NDC80-GFP fluorescence was observed in flagellated motile microgametes (**Fig. S4A**). Furthermore, no NDC80-GFP fluorescence was observed in either non-activated or activated female gametocytes (**Fig. S4A**). To independently test for an association of NDC80 with microtubules, we examined the effects of an anti-tubulin inhibitor (taxol) on NDC80 organisation during male gametogenesis. Addition of taxol at 1 min post activation blocked the dynamic progression of NDC80 distribution in more than 80% of male gametocytes while DMSO-treated gametocytes showed normal mitotic progression and NDC80 distribution (**Fig. S4B**). This showed that NDC80 distribution and localization depend on spindle dynamics and can be blocked by taxol treatment, which binds tubulin and stabilises microtubules by preventing depolymerisation. To further investigate the location of NDC80 in relation to the position of the centromere, NDC80-mCherry and the centromere-localised

condensin core subunit SMC4-GFP parasite lines were crossed and used for live cell imaging of both markers to establish their spatiotemporal relationship, as shown previously (Pandey et al., 2019). The location of both NDC80 and SMC4 was next to the nucleus, showing co-localization (**Fig. S4C**). Similarly, the basal body/axoneme marker, kinesin-8B-mCherry line (Zeeshan et al., 2019) was crossed with the NDC80-GFP line and this showed that NDC80 is located away from the basal body (kinesin-8B) during the start of male gametogenesis and later when kinesin-8B arranges across the axonemes (**Fig. S4D**). To confirm the centromere-associated localisation of NDC80 in a genome-wide manner, we performed a ChIP-seq experiment for NDC80-GFP in activated gametocytes (**Fig. S4E**). We observed strong ChIP-seq peaks at the centromeres of all 14 chromosomes similar to what was shown previously (Pandey et al., 2019).

NDC80-GFP shows unusual dynamics throughout the meiotic stages during zygote to ookinete differentiation.

Meiosis in the malaria parasite occurs during zygote differentiation to ookinete. This process takes 24 h to complete and during this time the ploidy of the parasite increases from 2N to 4N. To examine the behaviour of kinetochores throughout this process, we investigated the spatiotemporal profile of NDC80-GFP during ookinete differentiation.

NDC80-GFP fluorescence was first observed 1 to 1.5 h post-fertilization, as a single faint but distinct focal point, which gradually increased in intensity over the period 2 to 3 h post-fertilization (**Fig. 4**). As in male gametogenesis, no nuclear division was observed while the NDC80-GFP focus enlarged and divided to form a pair of elongated rod-like features. During stages II to IV of ookinete development, these rods appeared to fragment into multiple foci, ultimately resolving as 4 discrete foci in the mature ookinete, which has a 4N genome, at 18 h post-fertilization (**Fig. 4A**). The ultrastructure of the mature ookinete clearly showed four kinetochore clusters representing the 4N, but already segregated, genome within an intact nuclear membrane, consistent with the live cell imaging (**Fig. 4B**).

NDC80-GFP is present as multiple foci during oocyst development and sporozoite formation

During oocyst development and sporozoite formation, live cell imaging revealed NDC80-GFP fluorescence at multiple foci adjacent to the nuclear DNA during various stages of oocyst development from 7 days post-infection (dpi) of the mosquito to 21 dpi, as well as a single

focus in mature sporozoites (**Fig. 5A and B**). Ultrastructure analysis of oocyst development revealed an enlarged nucleus that formed a large multiple lobed structure with multiple nuclear poles/centriolar plaques/SPB/putative MTOC, followed by the formation of large numbers of sporozoites at the plasmalemma of the oocyst (**Fig. 5Bi**), as described previously (Ferguson et al., 2014; Schrevel et al., 1977). Detailed examination showed nuclear poles/centriolar plaques with kinetochores directed toward the developing sporozoites (**Fig. 5Bii**). The endomitotic process of sporozoite formation during sporogony resembles that of merozoite formation within host red cells and hence is similar to schizogony but with many more nuclei.

Immunoprecipitation of NDC80-GFP recovers canonical members of the NDC80 complex and reveals a highly divergent SPC24-like candidate

Previous comparative genomics studies revealed evidence for the presence of three NDC80 complex members in Plasmodiidae: NDC80, NUF2 and SPC25, but did not identify any candidate SPC24 ortholog (van Hooff et al., 2017). In another apicomplexan lineage (*Cryptosporidium*), however, an SPC24 ortholog was identified, raising the question of whether SPC24 has been lost in Plasmodiidae or is an as-yet-unidentified highly divergent ortholog present. To determine the composition of the NDC80 complex in *P. berghei*, possibly including novel NDC80 interactors that might be responsible for the distinct kinetochore localisations in different life stages, we immunoprecipitated NDC80-GFP from lysates of schizonts following culture for 8-hours and from gametocyte lysates one minute after activation. Mass spectrometric analysis of these pulldowns identified NUF2 (PBANKA_0414300) as the main binding partner of NDC80, and we detected SPC25 (PBANKA_1358800) as part of a longer list of proteins identified with fewer unique peptides and recovered for the NDC80-GFP precipitate, but absent from the GFP-only control (**Fig. 6B, Table S2**). For further scrutiny of the list of candidate proteins, we assessed similarity in behaviour to NDC80 and/or SPC25 across the control and NDC80-GFP pulldown experiments using both principal component analysis (PCA), and Spearman rank correlation. We selected candidate proteins that showed similar variance to SPC25 based on the first two components of the PCA analysis, and those that had a correlation value of $R>0.7$ for both NDC80 and SPC25 (**Fig S5**). Of the resulting list, most proteins have functions in transcription/RNA-related processes, and one is the extracellular protein casein kinase 1 (PBANKA_0912100)(Dorin-Semblat et al., 2015), suggesting likely non-specific association within the cell lysate. However, one protein stood out: PBANKA_1442300, a protein with a long coiled-coil region

and a predicted C-terminal globular domain, which suggested that it might be a SPC24 ortholog (**Fig. 6B**).

To test whether this candidate was a genuine SPC24 ortholog, we first queried available databases for PBANKA_1442300 homologs using conventional sequence similarity detection approaches, but detected none outside of hematozoan lineages, consistent with these approaches having failed to discover Plasmodiidae SPC24 candidates in the past (Plowman et al., 2019; van Hooff et al., 2017). Therefore, we employed more sophisticated protein modelling approaches to discover PBANKA_1442300 orthologs. We constructed a large sequence database consisting of 64 apicomplexan and other eukaryotic genomes and transcriptomes (see **Table S3** for sources of the sequence database) and generated Hidden Markov Models (HMM) of automatically defined homologous groups of sequences (see methods). We then compared these models with HMM profiles of PBANKA_1442300-like homologs and bona fide eukaryotic SPC24 orthologs found in our dataset. This multi-step approach yielded candidate apicomplexan SPC24 orthologs (**Fig. 6C**). Hematozoan homologs (including PBANKA_1442300) were significantly similar ($E > 10^{-5}$) to the previously unannotated group of coccidian homologous sequences (including the *Toxoplasma gondii* gene TGME49_212800). The merged HMM profile of the C-terminal globular domain of these two groups, in turn, was significantly similar to that of eukaryote-wide SPC24 orthologs ($E > 10^{-5}$) (**Fig. 6C, Fig. S6**). These analyses provide strong credence to the idea that PBANK_1442300- and TGME49_212800-like sequences are divergent but bona fide homologs of SPC24. Given that no other SPC24 candidates were found in these taxa, and the proteomic evidence of association of PBANKA_1442300 with SPC25, NDC80 and NUF2, it is very likely that PBANK_1442300 and TGME49_212800 are genuine SPC24 functional orthologs (**Fig. 6C, Sequence File S1, Table S4**).

Discussion

Cellular proliferation in eukaryotes requires chromosome replication and precise segregation, followed by cell division, to ensure that daughter cells have identical copies of the genome. This happens through assembly of a spindle to which the centromeric region of chromosomes is attached through the kinetochore. Although the organisation of spindle microtubules, the molecular composition of kinetochores, and the modes of spindle pole separation vary extensively among eukaryotes (Akiyoshi and Gull, 2013; Drechsler and McAinsh, 2012; van Hooff et al., 2017), the microtubule-binding subunit NDC80 is conserved across most eukaryotes including apicomplexan parasites such as *Toxoplasma gondii* and *Plasmodium* spp. (Akiyoshi and Gull, 2013; Farrell and Gubbels, 2014; van Hooff et al., 2017). In addition,

since chromosomes only bear one kinetochore, the outer-kinetochore subunit NDC80 is an excellent tool to start probing the rather surprising chromosome dynamics during the different stages of the life cycle in *P. berghei*.

In this study we have sought to understand the assembly and dynamics of the mitotic machinery during the diverse modes of nuclear division in *Plasmodium* using the kinetochore-protein NDC80 as a marker for chromosome attachment to the mitotic spindle. For this we generated a transgenic parasite line to express endogenous C-terminal GFP-labelled NDC80, a protein which faithfully replicates kinetochore location and function during all diverse mitotic and meiotic stages of the life cycle. Live-cell imaging of the fluorescent protein, complemented with ultrastructural studies by electron microscopy, revealed a subcellular location of NDC80-GFP at discrete foci adjacent to nuclear DNA in all replicative stages of the *P. berghei* life cycle. The distribution and dynamic spatiotemporal profile corresponded to the replication of chromosomes during the atypical mitotic and meiotic processes of DNA replication in this organism. Non-replicating stages, including the intraerythrocytic ring stage, the extracellular mature merozoite, the non-activated female gametocyte and the motile male gamete show no evidence of kinetochore assembly as no NDC80-GFP expression was observed. Using a combination of GFP-pulldown and a sensitive homology detection workflow we identified all four components of the NDC80 complex throughout apicomplexans, including a likely SPC24 ortholog candidate in Plasmodiidae.

The subcellular localization data for NDC80-GFP revealed a discreet single focus adjacent to the haploid nuclear genome, which presumably contains the centromeres of all 14 chromosomes. Such clustering of centromeric regions has been demonstrated in yeast (Richmond et al., 2013), human cells (Solovei et al., 2004) and *Toxoplasma* (Farrell and Gubbels, 2014). It is thought to be important for genome integrity, but the exact reason for it is not well understood. In the stages of the life cycle where this NDC80 clustering is not detected, perhaps the 14 kinetochores are not fully assembled and therefore NDC80-GFP expression is not detectable (Hoeijmakers et al., 2012). This is consistent with the idea that clustering of centromeres in *Plasmodium falciparum* occurs only prior to the onset of chromosome segregation (Hoeijmakers et al., 2012). It is of interest that following fertilization there is a single focus despite the genome being diploid, this may well reflect the tight pairing of sister chromatids allowing recombination to occur at this stage.

Although our data showed a clustered location of NDC80, its actual role in chromosome clustering is not known. Previous studies on yeast and *Toxoplasma* showed no role of NDC80 in clustering and suggested a sole role in attachment to spindle MTs during chromosome

segregation. A single MT binds each kinetochore in budding yeast (Westermann et al., 2007); whereas in *Toxoplasma* a maximum of only 11 MTs were detected despite the fact that there are 13 chromosomes (Bunnik et al., 2019; Farrell and Gubbels, 2014; Swedlow et al., 2002). Within the closely related Coccidian parasites, a number of variations in the details of the process of asexual division has been described, relating to timing and number of genome and nuclear divisions (Ferguson et al., 2008) The coccidian parasite *Sarcocystis neurona*, which divides by endopolygeny forms a polyploid nucleus culminating in 64 haploid daughter cells (Farrell and Gubbels, 2014; Vaishnava et al., 2005). During this process, intranuclear spindle poles are retained throughout the cell cycle, which suggests constant attachment of chromosomes to spindle MTs via kinetochores to ensure genome integrity throughout *Sarcocystis* cell division (Farrell and Gubbels, 2014; Vaishnava et al., 2005). In contrast, *Plasmodium* (a hemosporidian), undergoes classical schizogony with a variable number of cycles of genome replication and nuclear division resulting a multinucleated cell (Arnot et al., 2011). This is similar to what is seen in the coccidian parasites *Eimeria* spp. and *Toxoplasma*, with daughter cell formation being associated with the final nuclear division. This fact is well demonstrated by the NDC80 localization in this study, which shows 1 or 2 NDC80-GFP foci per nucleus. The asynchronous nature of the division during these stages is shown by nuclei having either one or two NDC80-GFP foci (and intermediate forms) within the same cell. We made similar observations in our previous study of the SPB/MTOC/centriole plaque marker for centrin, CEN-4 (Roques et al., 2019). These studies also suggest that the kinetochore and centrosome/centriolar plaque duplicate before nuclear division starts; therefore the duplication of NDC80 and CEN-4 sets the stage for mitosis at each round of nuclear division in *Plasmodium*, as in *Toxoplasma* (Suvorova et al., 2015). In contrast, during male gametogenesis the genome size increases to 8N, and this corresponds to the formation of 8 distinct NDC80-GFP foci, before nuclear division, with asynchronous chromosome replication and segregation. Most notable is the presence of unique kinetochore bridges or rod-like structures during genome replication and chromosome segregation. It appears that two hemi-spindles associated with the kinetochores are joined together to form a mitotic spindle at the earliest stages of endoreduplication during male gametogenesis. This is then followed by kinetochore movement to opposite poles once all the duplicated chromosomes are segregated. The mitotic process of duplication proceeds in the absence of nuclear division (karyokinesis), which results in the very atypical kinetochore dynamics that are consistent with the live cell imaging data.

Similarly, during meiosis in ookinete development the genome size increases to 4N, represented by bridge-like kinetochore structures during replicative stages and at the end,

resulting in four distinct NDC80-GFP foci suggesting kinetochore clustering to facilitate chromosome segregation, although no nuclear division takes place.

During sporogony multiple lobes are formed and the intranuclear spindle may be formed during multiple nuclear divisions, as revealed by ultrastructural studies during sporozoite formation (Schrevel et al., 1977). Further ultrastructure analyses identified typical nuclear spindles with attached kinetochores, radiating from the nuclear poles located within an intact nuclear membrane during schizogony, male gametogenesis, ookinete development and sporogony. Our data are consistent with a previous report, in which kinetochore localization was revealed using ultrastructural studies of *P. berghei* sporogony, and the duplication of hemi-spindles during replication was suggested (Schrevel et al., 1977). Overall, based on all these results, consistent kinetochore clustering occurs within Apicomplexa.

NDC80 is a major constituent of kinetochores and is highly conserved among eukaryotes including *Plasmodium*. However, many of the molecular details of kinetochore architecture and function in *Plasmodium* remain to be explored. A recent comparative evolutionary analysis suggested a distinct kinetochore network in many eukaryotes including *P. falciparum* and other alveolates, with many of the highly conserved kinetochore complex proteins being absent in *Plasmodium* (van Hooff et al., 2017). Of 70 conserved kinetochore proteins only eleven were found to be encoded in the *P. falciparum* genome. These eleven proteins include NDC80, NUF2, CENP-C/-A/-E, SPC25 and others that are highly conserved across 90 eukaryotic species, but genes for many other conserved proteins like SPC24, MAD-1/-2 and MIS12 were found to be absent (van Hooff et al., 2017). Recent studies have shown the presence of unconventional kinetochore proteins in kinetoplastids (Akiyoshi and Gull, 2013; D'Archivio and Wickstead, 2017), suggesting that different kinetochore architectures are possible. Previous reports have shown the association of CENP-A and CENP-C with centromeres in *P. falciparum* (Verma and Surolia, 2013; Verma and Surolia, 2014). Using a combination of GFP-pulldown and a sensitive homology detection workflow we identified all four components of the NDC80 complex throughout apicomplexans, including a highly divergent SPC24 ortholog candidate in Plasmodiidae (PBANKA_1442300). Previous studies in *Toxoplasma gondii* only identified NDC80 and NUF2 (Farrell and Gubbels, 2014), but here we predict the presence of SPC25 (TGME49_232400) and a SPC24-like homolog (TGME49_212800) in this organism. We favour the interpretation of PBANKA_1442300 being a bona fide SPC24 ortholog for two reasons: (1) it was detected as a putative NDC80-GFP binding partner, and (2) our sequence analyses indicate it has a C-terminal RWD-like domain most similar to that of SPC24. It seems unlikely that the ancestor of hematozoa and coccidia would have duplicated the gene and then discarded the functional kinetochore SPC24 ortholog. Further proteomic and co-localisation

studies of SPC25 and SPC24 with each other as well as other kinetochore markers (for example, NUF2 and CENP-C) will be needed to confirm that the candidate SPC24 ortholog is a true kinetochore protein in *Plasmodium*. Our approach has illustrated that conspicuous absences of subunits of highly conserved and essential complexes, such as the obligate tetrameric NDC80 complex, are to be treated with caution, and additional sensitive homology searches using HMM-HMM comparison should be employed to more thoroughly test for homolog presences or absences. As such, patterns of extensive loss have been observed in eukaryotic parasites before, and we expect that systematic searches for conspicuous absences of subunits of specific complexes will yield a large number of highly divergent homologs in Apicomplexa.

NDC80 and NUF2 are the most conserved subunits of the NDC80 complex, whereas the SPC24-SPC25 dimer that interacts with the centromere is highly divergent both in sequence and in length. Strikingly, the candidate SPC24 orthologs in Plasmodiidae and Coccidia are 3 to 4 times longer than those of other eukaryotes (**Fig. 6C**), and their RWD domains are also relatively divergent. What this means exactly is unclear, but we speculate that the larger NDC80 complex present in some but not all Apicomplexa engages centromere-proximal kinetochore proteins in a different way than in other eukaryotes, and possibly through direct interaction with CENP-A and CENP-C. This notion is supported by the apparent absence of the canonical interaction partners of SPC24 and SPC25, namely CENP-T and the MIS12 complex (van Hooff et al., 2017), and the extended length of all NDC80 complex members within coccidia (~1.2 - 3 times longer). In Plasmodiidae only the SPC24 candidate ortholog is extended (~3 to 4 times longer), but in Coccidia, SPC25 is also twice as long as canonical SPC25s, potentially resulting in a larger and longer NDC80 complex. The extended length of SPC24 candidate orthologs results largely from N-terminal extensions of the coiled-coil region (Coccidia and Plasmodiidae), and/or large insertions into the loop of the RWD domain (Coccidia). We envision that these extensions may provide additional binding sites for novel interactors important in kinetochore clustering and/or the remarkable ‘bridge/rod’ phenotype we have observed in this study. The SPC24 N-terminal coiled-coil extension may also have additional interactions with the heterodimeric coiled-coils of the NDC80 and NUF2 subunits, providing extra rigidity to the tetrameric NDC80 superstructure.

In summary, this study demonstrates the dynamic expression and location of NDC80 during the different proliferative stages of the malaria parasite and reveals both the disassembly and reassembly, as well as clustering, of kinetochores. It also shows the asynchronous closed mitotic division of *Plasmodium* during schizogony and sporogony and provides novel insights into the chromosome segregation in male gametogenesis and in various stages of meiosis

during zygote differentiation to ookinetes. ChIP-seq and colocalisation studies clearly showed the centromeric location of NDC80 in *Plasmodium*. The protein pulldown and bioinformatics studies revealed that NDC80 has the full complement of four subunits, though SPC24 is highly divergent compared to other eukaryotes. This analysis of NDC80 will also facilitate future studies of cell division and comparative analyses of chromosome dynamics in evolutionarily divergent eukaryotic cells.

Material and Methods:

Ethical statement

All animal-related work performed at the University of Nottingham has undergone an ethical review process and been approved by the United Kingdom Home Office with the project license number 30/3248 and PDD2D5182. The work has been carried out in accordance with the United Kingdom ‘Animals (Scientific Procedures) Act 1986’ and was in compliance with ‘European Directive 86/609/EEC’ for the protection of animals used for experimental purposes. A combination of ketamine followed by antisedan was used for general anaesthesia and sodium pentobarbital was used for terminal anaesthesia. Proper care and efforts were made to minimise animal usage and suffering.

Six to eight-week-old female Tuck-Ordinary (TO) (Harlan) or CD1 outbred mice (Charles River) were used for all experiments.

Generation of transgenic parasites

The transgenic lines for NDC80 (PBANKA_1115700) were created using single homologous recombination as shown in **Fig. S1**. The oligonucleotides used to generate transgenic lines are provided in **Supplementary Table S1**. For GFP tagging, a 1153bp region of *Ndc80* without the stop codon was inserted upstream of the *gfp* sequence in the p277 plasmid vector using KpnI and Apal restriction sites as described previously (Tewari et al., 2010). The p277 vector contains the human *dhfr* cassette, conveying resistance to pyrimethamine. Before transfection, the sequence was linearised using EcoRV. The *P. berghei* ANKA line 2.34 was used for transfection by electroporation (Janse et al., 2006). Immediately, electroporated parasites were mixed with 100µl of reticulocyte-rich blood from a phenylhydrazine (6 mg/ml, Sigma) treated, naïve mouse and incubated at 37°C for 30 min before intraperitoneal injection.

Pyrimethamine (70 mg/L, Sigma) was supplied in the drinking water from 1-day post-infection (dpi) to 4-dpi. Infected mice were monitored for 15 days and drug selection was repeated after passage to a second mouse. Integration PCR and western blot were performed to confirm successful generation of the transgenic line. For integration PCR, primer 1 (IntT259) and primer 2 (ol492) were used to confirm integration of the GFP targeting construct. Primer 1 and primer 3 (T2592) were used as a control. We also generated a mCherry-tagged NDC80 transgenic parasite line as shown in the schematic provided in **Fig. S1**.

Western Blot

For western blotting, purified schizonts were lysed using lysis buffer (10 mM TrisHCl pH 7.5, 150 mM NaCl, 0.5 mM EDTA, 1% NP-40 and 1% Sarkosyl). The samples were boiled for 10 min after adding Laemmli sample buffer to the lysed cells. The sample was centrifuged at 13500 g for 5 min and electrophoresed on a 4–12% SDS-polyacrylamide gel. Subsequently, resolved proteins were transferred to nitrocellulose membrane (Amersham Biosciences) and immunoblotting was performed using the Western Breeze Chemiluminescence Anti-Rabbit kit (Invitrogen) and anti-GFP polyclonal antibody (Invitrogen) at a dilution of 1:1250, according to the manufacturer's instructions.

Localization of NDC80-GFP throughout the parasite life cycle

Live cell imaging of transgenic parasite lines was performed at different proliferative stages during the parasite life cycle (**Fig. 1**) as described previously (Roques et al., 2019; Saini et al., 2017) using a Zeiss AxioImager M2 microscope fitted with an AxioCam ICc1 digital camera (Carl Zeiss, Inc).

Blood stage schizogony

Infected mouse blood provided asexual blood and gametocyte stages of the *P. berghei* life cycle. Schizont culture (RPMI 1640 containing 25 mM HEPES, 1:10 (v/v) fetal bovine serum and penicillin/streptomycin 1:100) at different time points was used to analyse various stages of asexual development from ring to merozoite. The periods used for analysis and imaging were 0 to 1 h for ring stage parasites, 2 to 4 h for trophozoites, 6 to 8 h for early and mid-stage schizonts, 9 to 11 h for late segmented schizonts, and 18 to 24 h for mature schizonts and released merozoites in schizont culture medium.

Male gametocyte development

In vitro cultures were prepared to analyse non-activated gametocytes, activated gametocytes and male exflagellation. For *in vitro* exflagellation studies, gametocyte-infected blood was obtained from the tails of infected mice using a heparinised pipette tip. Gametocyte activation was performed by mixing 100 μ l of ookinete culture medium (RPMI 1640 containing 25 mM HEPES, 20% fetal bovine serum, 10 mM sodium bicarbonate, 50 μ M xanthurenic acid at pH 7.6) with gametocyte-infected blood. To study different time points during microgametogenesis, gametocytes were purified using NycoDenz gradient (48%) and monitored at different time points to study mitotic division (male gametogenesis, 0 to 15 min post-activation (mpa).

Ookinete development

To study ookinete development, gametocyte infected blood was incubated in ookinete medium for 24 hpa and various stages of zygote differentiation and ookinete development were monitored at different time points (0 min for nonactivated gametocytes, 30 min for activated gametocytes, 2 to 3 h for zygotes, 4 to 5 h for stage I, 5 to 6 h for stage II, 7 to 8 h for stage III, 8 to 10 h for stage IV, 11 to 14h for stage V, and 18 to 24 h for mature ookinetes post activation in ookinete medium).

Oocyst and sporozoite development

For mosquito transmission stages and bite back experiments, triplicate sets of 30 to 50 *Anopheles stephensi* mosquitoes were used. The mosquito guts were analysed on different days post-infection (dpi): 7 dpi, 12 dpi, 14 dpi and 21 dpi to check expression and localization of NDC80GFP during oocyst development and sporozoite formation.

Schizogony in liver stages

To study localization of NDC80-GFP in *P. berghei* liver stages, 100,000 HeLa cells were seeded in glass-bottomed imaging dishes. Salivary glands of female *A. stephensi* mosquitoes infected with NDC80-GFP parasites were isolated and sporozoites were released using a pestle to disrupt salivary gland cells. The released sporozoites were pipetted gently onto the HeLa cells and incubated at 37 °C in 5% CO₂ in air, in complete minimum Eagle's medium containing 2.5 μ g/ml amphotericin B. For initial infection, medium was changed at 3 h post-infection and thereafter once a day. To perform live cell imaging, Hoechst 33342 (Molecular Probes) was added (1 μ g/ml) and imaging was done at 55 h post-infection using a Leica TCS

SP8 confocal microscope with the HC PL APO 63x/1.40 oil objective and the Leica Application Suite X software.

Indirect immunofluorescence assay

IFA studies were performed using poly-L-lysine coated slides on which schizonts had been previously fixed in 2% paraformaldehyde (PFA) in microtubule stabilising buffer (MTSB:10 mM MES, 150 mM NaCl, 5 mM EGTA, 5 mM MgCl₂, 5 mM glucose) in 1X-PBS for 30 min at room temperature (RT) and smeared onto slides. The fixed cells were permeabilized using TBS containing 0.2% TritonX-100 for 5 min and washed three times with TBS before blocking. For blocking, 1 hour incubation was performed with TBS solution containing 3% BSA (w/v) and 10% goat serum (v/v). TBS containing 1% BSA and 1% goat serum was used to dilute the antibodies for the incubations. Anti-GFP rabbit antibody (Invitrogen) was used at 1:250 dilution, anti-alpha-tubulin mouse antibody (Sigma-Aldrich) was used at 1:1000 dilution, and anti-centrin mouse clone 20h5 antibody (Millipore) was used at 1:200 dilution; each was incubated for 1 hour at RT. Three washes were performed with TBS, then AlexaFluor 568 labelled anti-rabbit (red) and AlexaFluor 488 labelled anti-mouse (green) (Invitrogen) (1:1000 dilution) were used as secondary antibodies and incubated for 40 min at RT. A similar protocol was followed for gametocytes except the cells were fixed in 4% PFA in MTSB. Slides were mounted with Vectashield containing DAPI (blue) and sealed using nail polish. Images were captured as described for live imaging.

Super resolution microscopy

A small volume (3 µl) of schizont culture was mixed with Hoechst dye and pipetted onto 2 % agarose pads (5x5 mm squares) at room temperature. After 3 min these agarose pads were placed onto glass bottom dishes with the cells facing towards glass surface (MatTek, P35G-1.5-20-C). Cells were scanned with an inverted microscope using Zeiss C-Apochromat 63x/1.2 W Korr M27 water immersion objective on a Zeiss Elyra PS.1 microscope, using the structured illumination microscopy (SIM) technique. The correction collar of the objective was set to 0.17 for optimum contrast. The following settings were used in SIM mode: lasers, 405 nm: 20%, 488 nm: 50%; exposure times 100 ms (Hoechst) and 25 ms (GFP); three grid rotations, five phases. The band pass filters BP 420-480 + LP 750 and BP 495-550 + LP 750 were used for the blue and green channels, respectively. Multiple focal planes (Z stacks) were recorded with 0.2 µm step size; later post-processing, a Z correction was done digitally on the 3D rendered images to reduce the effect of spherical aberration (reducing the elongated view in Z; a process previously tested with fluorescent beads). Images were processed and all focal planes were digitally merged into a single plane (Maximum intensity projection). The images

recorded in multiple focal planes (Z-stack) were 3D rendered into virtual models and exported as images and movies (see supplementary material). Processing and export of images and videos were done by Zeiss Zen 2012 Black edition, Service Pack 5 and Zeiss Zen 2.1 Blue edition.

Chromatin Immunoprecipitation sequencing analysis (ChIP-seq)

For the ChIP-seq analysis, libraries were prepared from crosslinked cells (using 1% formaldehyde). The crosslinked parasite pellets were resuspended in 1 mL of nuclear extraction buffer (10 mM HEPES, 10 mM KCl, 0.1 mM EDTA, 0.1 mM EGTA, 1 mM DTT, 0.5 mM AEBSF, 1X protease inhibitor tablet), post 30 min incubation on ice, 0.25% Igepal-CA-630 was added and homogenized by passing through a 26G x ½ needle. The nuclear pellet extracted through 5000 rpm centrifugation, was resuspended in 130 µl of shearing buffer (0.1% SDS, 1 mM EDTA, 10 mM Tris-HCl pH 7.5, 1X protease inhibitor tablet), and transferred to a 130 µl Covaris sonication microtube. The sample was then sonicated using a Covaris S220 Ultrasonicator for 10 min for schizont samples and 6 min for gametocyte samples (Duty cycle: 5%, Intensity peak power: 140, Cycles per burst: 200, Bath temperature: 6°C). The sample was transferred to ChIP dilution buffer (30 mM Tris-HCl pH 8, 3 mM EDTA, 0.1% SDS, 30 mM NaCl, 1.8% Triton X-100, 1X protease inhibitor tablet, 1X phosphatase inhibitor tablet) and centrifuged for 10 min at 13,000 rpm at 4°C, retaining the supernatant. For each sample, 13 µl of protein A agarose/salmon sperm DNA beads were washed three times with 500 µl ChIP dilution buffer (without inhibitors) by centrifuging for 1 min at 1000 rpm at room temperature, then buffer was removed. For pre-clearing, the diluted chromatin samples were added to the beads and incubated for 1 hour at 4°C with rotation, then pelleted by centrifugation for 1 min at 1000 rpm. Supernatant was removed into a LoBind tube carefully so as not to remove any beads and 2 µg of anti-GFP antibody (ab290, anti-rabbit) were added to the sample and incubated overnight at 4°C with rotation. Per sample, 25 µl of protein A agarose/salmon sperm DNA beads were washed with ChIP dilution buffer (no inhibitors), blocked with 1 mg/mL BSA for 1 hour at 4°C, then washed three more times with buffer. 25 µl of washed and blocked beads were added to the sample and incubated for 1 hour at 4°C with continuous mixing to collect the antibody/protein complex. Beads were pelleted by centrifugation for 1 min at 1000 rpm at 4°C. The bead/antibody/protein complex was then washed with rotation using 1 mL of each buffers twice; low salt immune complex wash buffer (1% SDS, 1% Triton X-100, 2 mM EDTA, 20 mM Tris-HCl pH 8, 150 mM NaCl), high salt immune complex wash buffer (1% SDS, 1% Triton X-100, 2 mM EDTA, 20 mM Tris-HCl pH 8, 500 mM NaCl), high salt immune complex wash buffer (1% SDS, 1% Triton X-100, 2 mM EDTA, 20 mM Tris-HCl pH 8, 500 mM NaCl), TE wash buffer (10 mM Tris-HCl pH 8, 1 mM

EDTA) and eluted from antibody by adding 250 µl of freshly prepared elution buffer (1% SDS, 0.1 M sodium bicarbonate). We added 5 M NaCl to the elution and cross-linking was reversed by heating at 45°C overnight followed by addition of 15 µl of 20 mg/mL RNAase A with 30 min incubation at 37°C. After this, 10 µl 0.5 M EDTA, 20 µl 1 M Tris-HCl pH 7.5, and 2 µl 20 mg/mL proteinase K were added to the elution and incubated for 2 hours at 45°C. DNA was recovered by phenol/chloroform extraction and ethanol precipitation, using a phenol/chloroform/isoamyl alcohol (25:24:1) mixture twice and chloroform once, then adding 1/10 volume of 3 M sodium acetate pH 5.2, 2 volumes of 100% ethanol, and 1/1000 volume of 20 mg/mL glycogen. Precipitation was allowed to occur overnight at -20°C. Samples were centrifuged at 13,000 rpm for 30 min at 4°C, then washed with fresh 80% ethanol, and centrifuged again for 15 min with the same settings. Pellet was air-dried and resuspended in 50 µl nuclease-free water. DNA was purified using Agencourt AMPure XP beads. Libraries were prepared using the KAPA Library Preparation Kit (KAPA Biosystems), and were amplified for a total of 12 PCR cycles (15 s at 98°C, 30 s at 55°C, 30 s at 62°C) using the KAPA HiFi HotStart Ready Mix (KAPA Biosystems). Libraries were sequenced using the NovaSeq 6000 System (Illumina), producing 100-bp reads.

FastQC (<https://www.bioinformatics.babraham.ac.uk/projects/fastqc/>), was used to analyze raw read quality. Any adapter sequences were removed using Trimmomatic (<http://www.usadellab.org/cms/?page=trimmomatic>). Bases with Phred quality scores below 25 were trimmed using Sickle (<https://github.com/najoshi/sickle>). The resulting reads were mapped against the *P. berghei* ANKA genome (v36) using Bowtie2 (version 2.3.4.1) or HISAT2 (version 2-2.1.0), using default parameters. Reads with a mapping quality score of 10 or higher were retained using Samtools (<http://samtools.sourceforge.net/>), and PCR duplicates were removed by PicardTools MarkDuplicates (Broad Institute). For Chip-seq analysis, raw read counts were determined to obtain the read coverage per nucleotide. Genome browser tracks were generated and viewed using the Integrative Genomic Viewer (IGV) (Broad Institute). Proposed centromeric locations were obtained from Iwanaga and colleagues (Iwanaga et al., 2010).

Inhibitor studies

Gametocytes were purified as above and treated with 1 µM taxol (Paclitaxel, T7402; Sigma) at 1 min post activation (mpa) and then fixed with 4% Paraformaldehyde (PFA) at 8 mpa. DMSO was used as a control treatment. These fixed gametocytes were then examined on a Zeiss Axio Imager M2 microscope fitted with an AxioCam ICc1 digital camera (Carl Zeiss, Inc).

Electron microscopy

Samples for different mitotic stages of parasite development including schizonts (24 hours in culture), activated male gametocytes (8 min post-activation), infected mosquito guts (12 to 14 days post infection) and the meiotic stage from the mature ookinete (24 hours post-activation) were fixed in 4% glutaraldehyde in 0.1 M phosphate buffer and processed for electron microscopy as previously described (Ferguson et al., 2005). Briefly, samples were post-fixed in osmium tetroxide, treated in bloc with uranyl acetate, dehydrated and embedded in Spurr's epoxy resin. Thin sections were stained with uranyl acetate and lead citrate prior to examination in a JEOL1200EX electron microscope (Jeol UK Ltd).

Protein pulldown, Immunoprecipitation and Mass Spectrometry

Schizonts, following 8 hours *in vitro* culture, and male gametocytes 1 min post activation of NDC80-GFP parasites were used to prepare cell lysates. Purified parasite pellets were crosslinked using formaldehyde (10 min incubation with 1% formaldehyde in PBS), followed by 5 min incubation in 0.125M glycine solution and 3 washes with phosphate buffered saline (PBS, pH 7.5). Immunoprecipitation was performed using crosslinked protein lysate and a GFP-Trap®_A Kit (Chromotek) following the manufacturer's instructions. Proteins bound to the GFP-Trap®_A beads were digested using trypsin and the peptides were analysed by LC-MS/MS. Mascot (<http://www.matrixscience.com/>) and MaxQuant (<https://www.maxquant.org/>) search engines were used for mass spectrometry data analysis. The PlasmoDB database was used for protein annotation. Peptide and proteins having minimum threshold of 95% were used for further proteomic analysis. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository with the dataset identifier PXD017619 and 10.6019/PXD017619.

To assess covariance among all proteins identified by mass spectrometry, a principal component analysis (PCA) was performed on proteins having peptide spectrum matches in NDC80-GFP, but not control-GFP samples (schizont and gametocytes), and further excluding proteins that were annotated to be part of the ribosome (see **Fig. S5**). NA values were transformed into zero, indicating the absence of any peptide detection for a particular protein. PCA was done on $\ln(x)$ transformed unique peptide values using the 'prcomp' function, which is part of the R-package stats (v3.6.2). In a similar approach Spearman rank correlation was calculated for all proteins identified with both NDC80 and SPC25, using the 'corr' function, which is part of the R-package ggpibr (v0.2.5). Graphs were visualised using the R-package ggplot2.

Comparative genomics of the NDC80 complex in Apicomplexa

Database: To increase the sensitivity for detecting highly divergent members of the NDC80 complex (NDC80, NUF2, SPC25, SPC24) in apicomplexan parasites, we constructed a large sequence database of (new) genomes and (meta)transcriptomes (see **Supplementary Table 3** for sources). This database consisted of 43 Apicomplexa and 8 Apicomplexa-affiliated lineages, including newly sequenced gregarines and ‘apicomonada’ (Janouskovec et al., 2019; Janouskovec et al., 2015; Mathur et al., 2019), and the non-parasitic apicomplexan *Nephromyces* (Muñoz-Gómez et al., 2019). A set of 13 eukaryotes representative of a wider range of the eukaryotic tree of life were added for which the NDC80 complex presence/absence pattern was studied previously (van Hooff et al., 2017). For transcriptomes for which no gene predictions were available, ORFs were predicted using TransDecoder (Long Orfs algorithm: <https://github.com/TransDecoder/TransDecoder>).

Ortholog detection: To uncover an initial set of sequences orthologous to NDC80 complex subunits in our database we made use of previously established eukaryote-wide Hidden Markov Models (HMM) of the Calponin Homology domains of NDC80 and NUF2, and the RWD domains for SPC24 and SPC25 (Tromer et al., 2019; van Hooff et al., 2017). Although potentially informative, coiled-coil regions were avoided as they tend to have sequence similarities with (other) non-homologous coiled-coil proteins. For details on the strategy for finding orthologs, see previously established protocols (van Hooff et al., 2017). Briefly, HMM-guided hits were realigned using mafft (option:eins-i) (Katoh and Standley, 2013), modelled as HMM, and searched iteratively against the database until no new orthologs could be detected. Conspicuous absences were further inspected by iterative searches using sequences of closely related lineages, including lower stringency matches (higher E-values, lower bitscores) that had a similar length and coiled-coil topology. HMMs were modelled (hmmbuild) and iteratively searched (hmmsearch or jackhmmer: E-value<0.05, bitscore>25) using the HMMER package (v3.1b) (Johnson et al., 2010). SPC24 orthologues in Coccidia and Hematozoa were detected by comparing our custom-made eukaryote-wide HMMs with HMM profiles of automatically defined orthologous groups (OrthoFinder (Emms and Kelly, 2019)), using the secondary structure aware HMM-vs-HMM search algorithm HHsearch (Steinegger et al., 2019). Specifically, HMM profiles of the orthologous groups containing PBANKA_1442300 (*P. berghei*) and TGME49_212800 (*T. gondii*) were merged and the resulting HMM was searched against a dataset containing HMMs of the orthologous groups defined by OrthoFinder, a previously established set containing scop70, pdb70 and PfamA version 31.0 [7], and custom-made HMM of kinetochore proteins (Tromer et al., 2019; van Hooff et al., 2017).

Alignments were visualised and modified using Jalview (Waterhouse et al., 2009). Figure 6 was made using Inkscape (<https://inkscape.org/>).

Competing interests

The authors declare no competing interests.

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Author contribution

RT conceived and designed all experiments. RT, MZ, ED and DB performed the experiments; RRS performed liver stage imaging; RP and RM performed super-resolution imaging; DJPF performed electron microscopy; ET and RW performed the bioinformatics and phylogenetic analysis; SA and KLR performed the ChIP-seq analysis; MZ, RT and ARB performed pulldown experiments, RP, MZ, AAH, and RT analysed the data; MZ, RP DSG, ET, RW AAH and RT wrote the manuscript and all others contributed to it.

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Figures

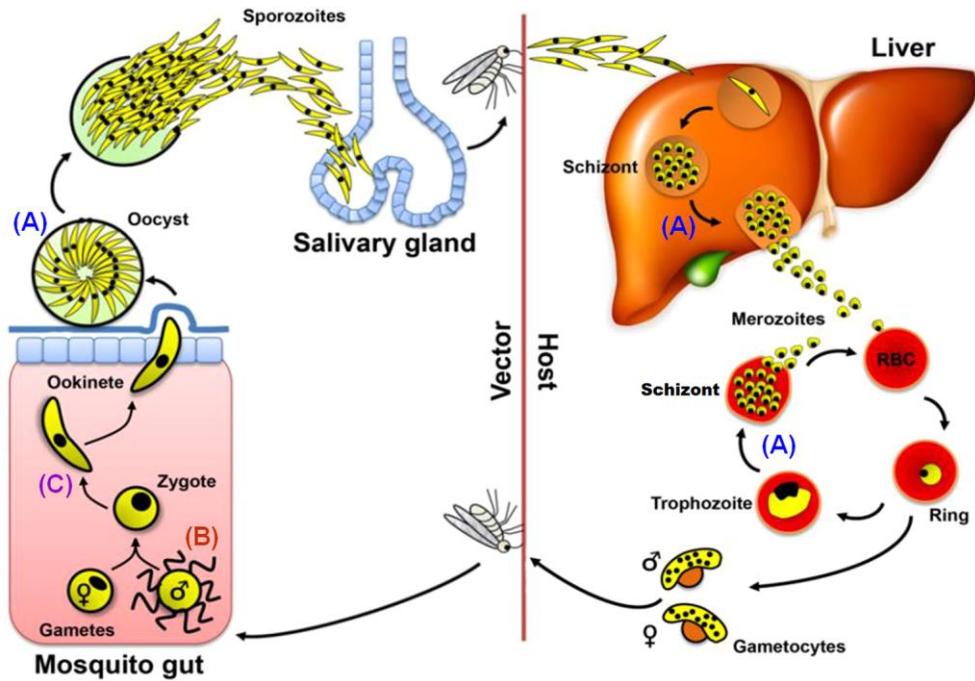


Fig. 1: Life cycle of rodent malaria parasite *Plasmodium berghei*. (A) represents endomitotic division: schizogony (schizont formation) in the liver and blood cells of the mammalian host and sporogony (sporozoite formation) in the mosquito gut. (B) and (C) represent atypical mitotic division by endoreduplication during male gametogenesis and meiotic division during the zygote to ookinete differentiation in the mosquito gut, respectively.

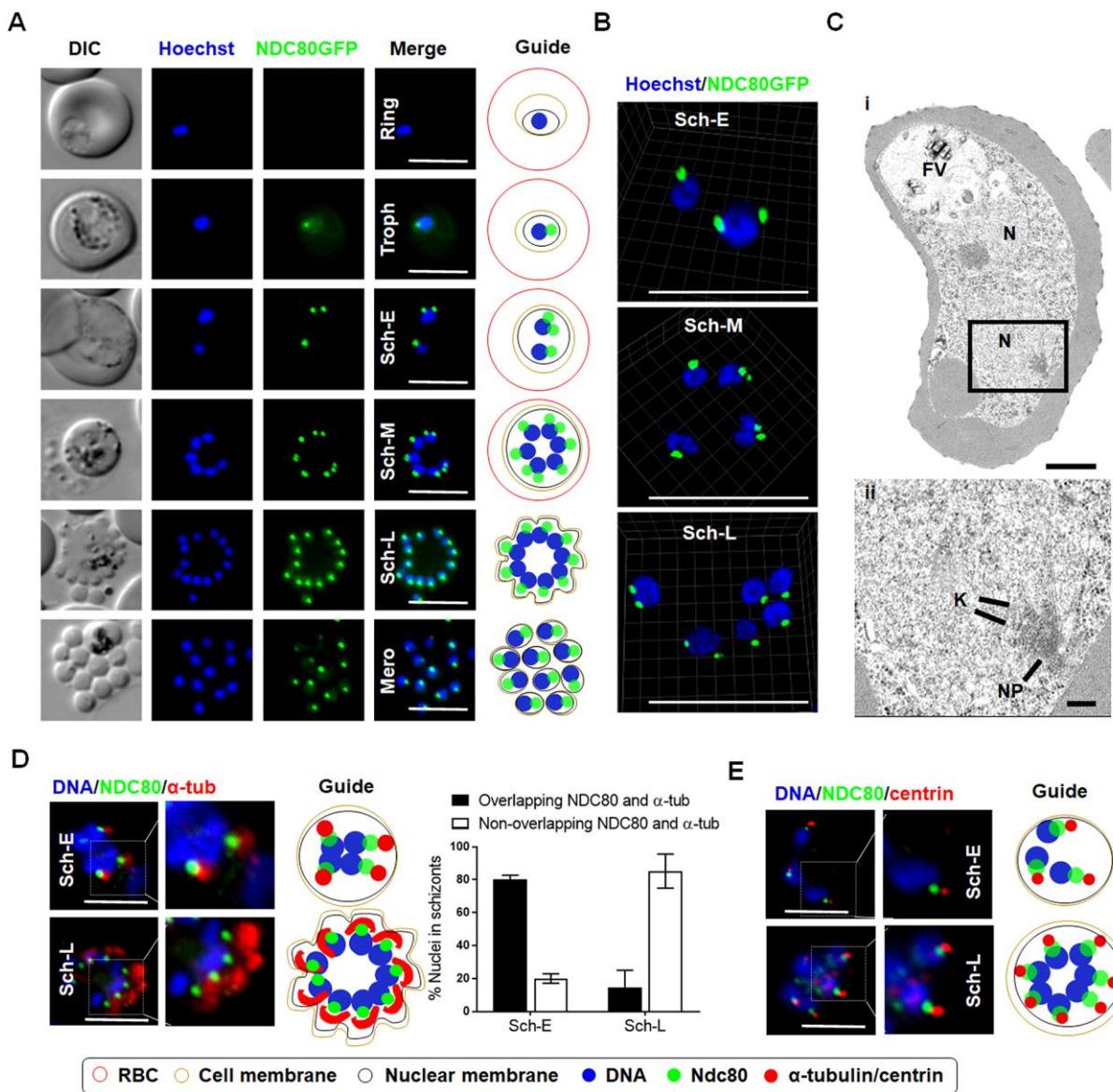


Fig. 2: NDC80-GFP localization during endomitotic cell division in schizogony. **(A)** Live cell imaging of NDC80-GFP expression and location during asexual blood stage (DIC: Differential interference contrast, Hoechst: DNA, NDC80-GFP: GFP, Merge: Hoechst and GFP fluorescence, Troph: Trophozoite, Sch-E: Early Schizont, Sch-M: Mid Schizont Sch-L: Late Schizont, Mero: Merozoite, 100x magnification, Scale bar = 5 µm.) and schematic guide depicting NDC80 localization during various developmental stages in the bloodstream of the parasite life cycle. **(B)** Live cell super-resolution 3D imaging for NDC80-GFP localization during asynchronous blood stage mitotic division. Scale bar = 5 µm. **(C)** Electron microscopy imaging of an early schizont showing kinetochore localization. **(i)** Section through an early schizont within a red blood cell, showing two nuclei (N) and the food vacuole (FV). Scale bar = 1 µm. **(ii)** Enlargement of the enclosed area showing part of the nucleus in which a nuclear

pole (NP), microtubules and attached kinetochores (K) can be seen. Scale bar = 100 nm. (**D**) Immunofluorescence fixed cell imaging and schematic guide of NDC80-GFP and co-localization with α -tubulin showing overlap during early schizont (**E**) Immunofluorescence fixed cell imaging and schematic guide of NDC80-GFP and co-localization with centrin (100x magnification). Scale bar = 5 μ m.

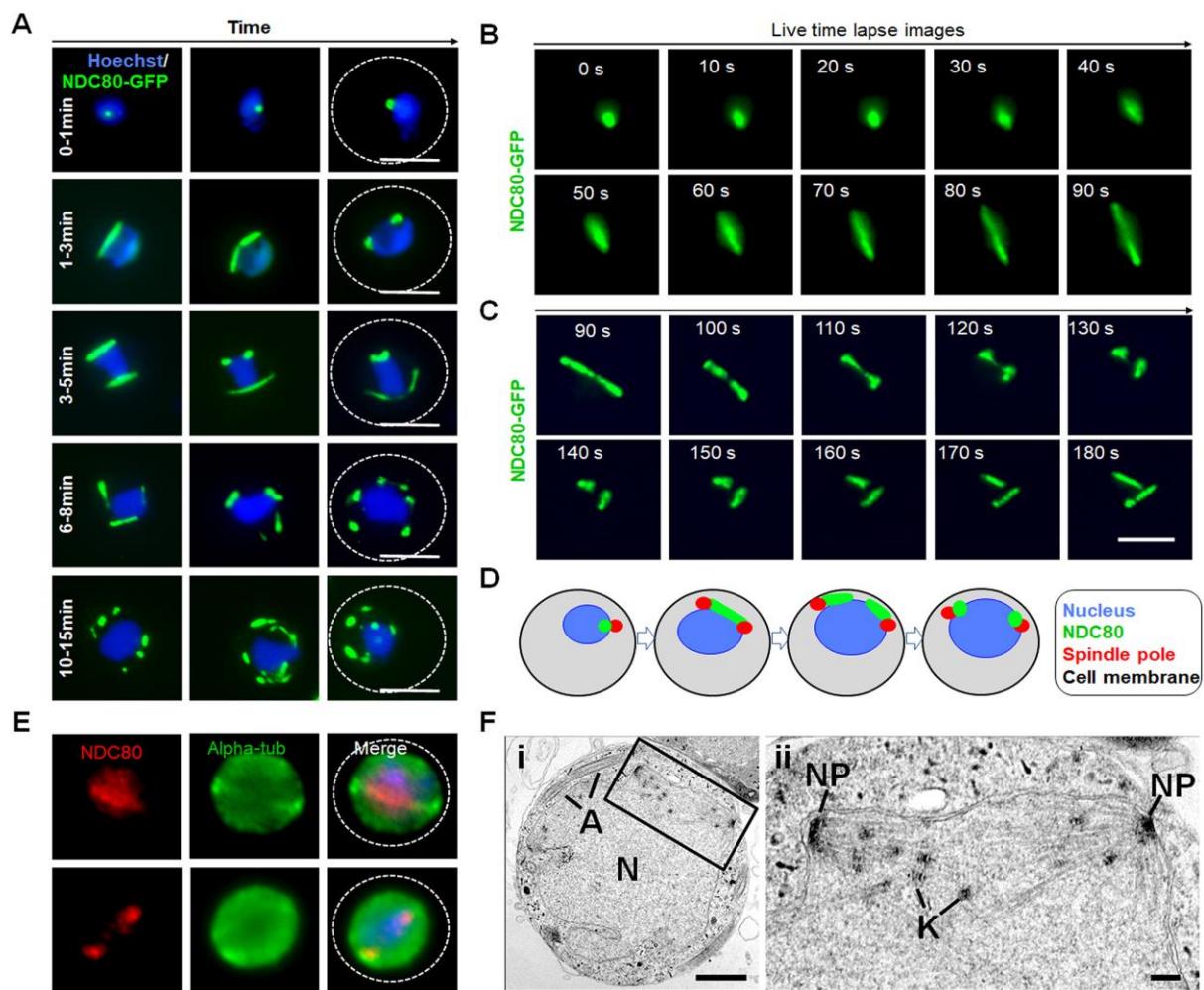


Fig. 3: Temporal dynamics of NDC80-GFP during male gametogenesis. **(A)** Live cell imaging for NDC80-GFP expression and location during endoreduplicative mitotic division in male gametogenesis (100x magnification). **(B-C)** Time-lapse screenshots for NDC80-GFP localization during male gametogenesis. Scale bar = 5 μ m. **(D)** Schematic representation showing dynamic localization of NDC80 during 1-3 min post activation of gametocyte (first round of nuclear division). **(E)** Indirect immunofluorescence assays showing co-localization of NDC80 (red) and α -tubulin (green) in male gametocytes activated for 1-3 min. **(F)** Electron microscopy on 8 min post-activation gametocytes for kinetochore localization. **(i)** Section through a mid-stage microgametocyte showing the large central nucleus (N) with axonemes (A) present in the peripheral cytoplasm. Scale bar = 1 μ m. **(ii)** Enlargement of the enclosed area showing the details of an intranuclear spindle with microtubules with attach kinetochores (K) radiating from the nuclear poles (NP). Scale bar = 100nm.

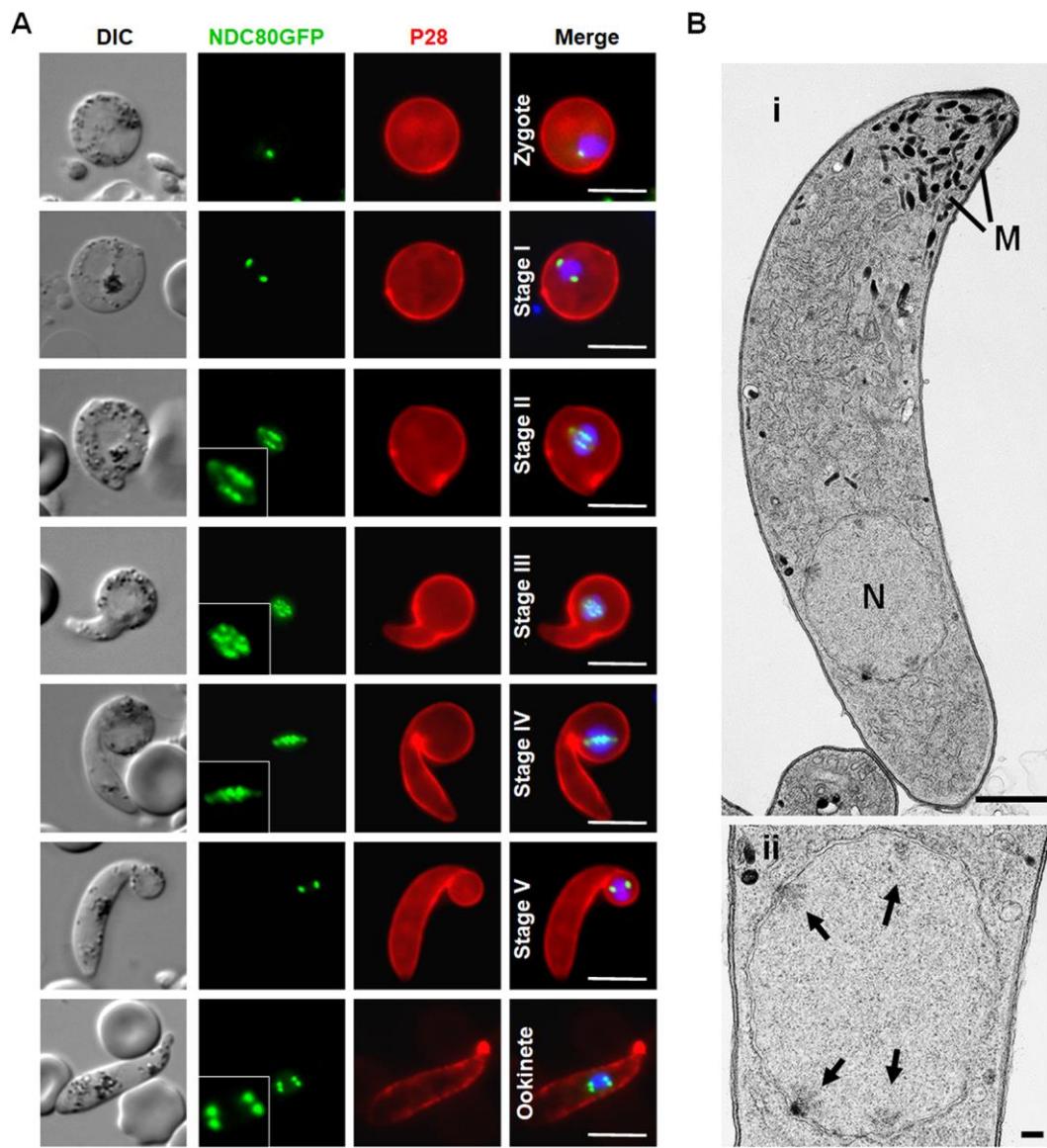


Fig. 4: Spatiotemporal profile of NDC80-GFP expression during meiotic stages in the ookinete. (A) Live cell imaging of NDC80-GFP localization during various stages of ookinete development from zygote to mature ookinete in the mosquito gut (100x magnification). Merge: Hoechst (blue, DNA), GFP (green) and P28 (red, cell surface marker of activated female gamete, zygote and ookinete stages). Scale bar = 5 µm. (B) Ultrastructural analysis of kinetochore localization in a mature ookinete. (i) Longitudinal section through a mature ookinete showing the apical complex with several micronemes (M) and the more posterior nucleus (N). Scale bar = 1 µm. (ii) Enlargement of the nucleus showing the location of the four nuclear poles (arrows). Scale bar = 100 nm.

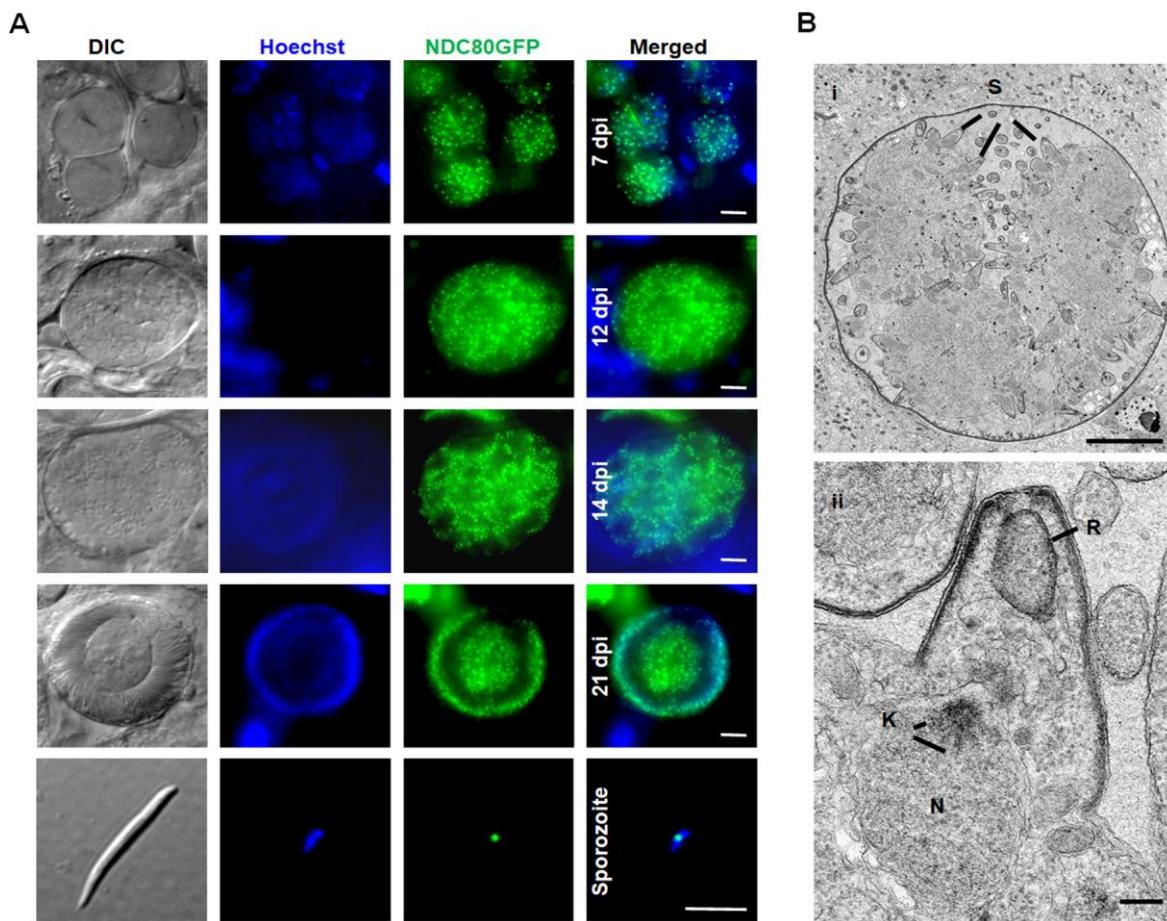


Fig. 5: NDC80-GFP localisation during oocyst development and sporozoite formation.

(A) Live cell imaging of NDC80-GFP in oocysts at 7, 12, 14 and 21 days post-infection (dpi) and a sporozoite. Panels: DIC (differential interference contrast), Hoechst (blue, DNA), NDC80-GFP (green, GFP), Merged: Hoechst (blue, DNA) and NDC80-GFP (green, GFP) Scale bar = 5 μ m. (B) Electron microscopy analysis of kinetochore location in an oocyst 12 day post-infection. (i) Central section through a mid-stage oocyst showing the early stages of sporozoite formation (S) at the surface of the oocyst cytoplasm. Scale bar = 10 μ m. (ii) Detail showing the early stage in sporozoite budding. Note the underlying nucleus (N) with the nuclear pole and attached kinetochores (K) directed toward the budding sporozoite. R – Rhopty anlagen. Scale bar = 100 nm.

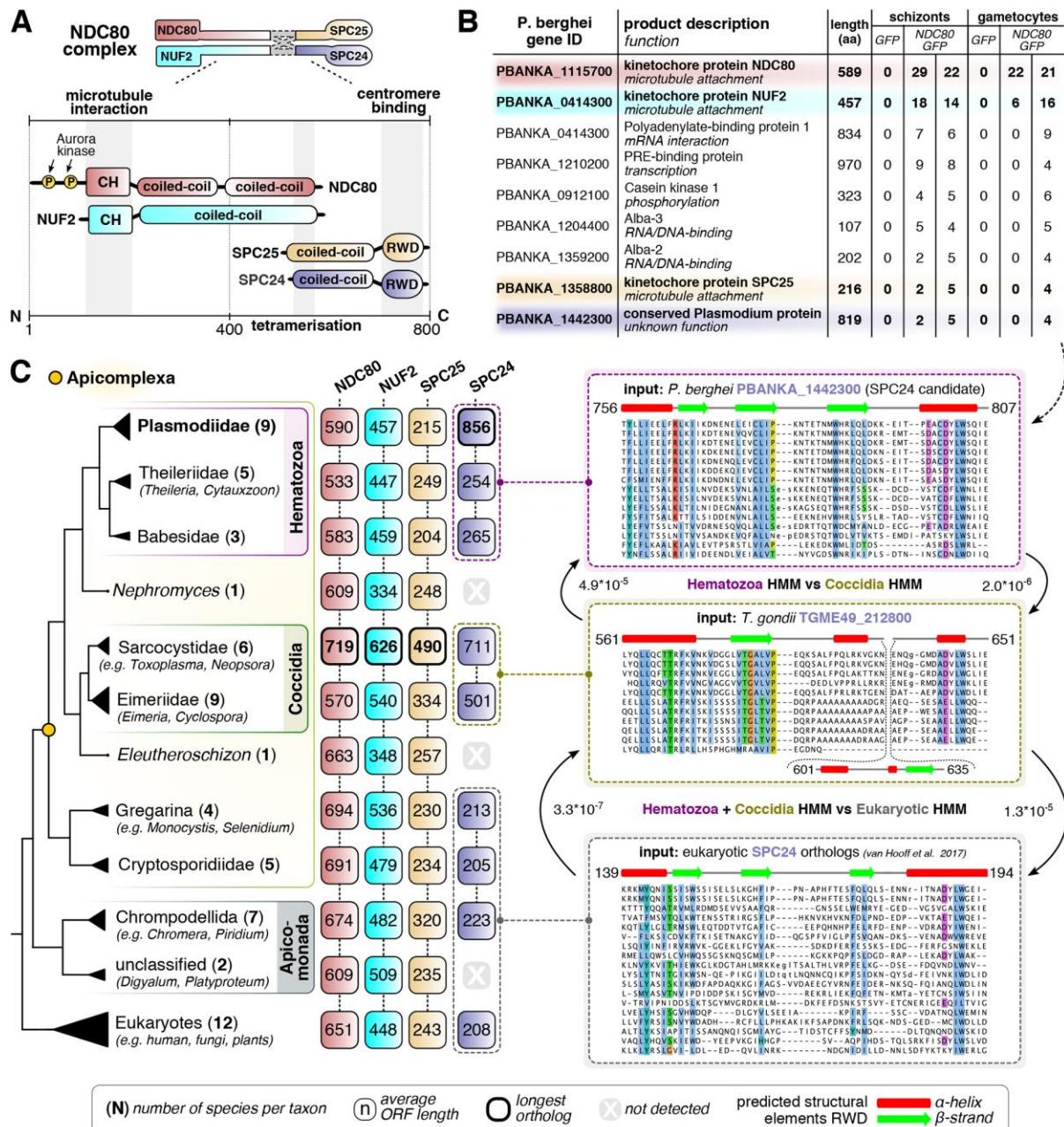


Fig. 6: The NDC80 complex is conserved in *Plasmodium spp.* and most Apicomplexa

(A) Domain composition of the four subunits of the NDC80 complex tetramer as commonly found in model eukaryotes. N and C denote the amino and carboxy termini of the proteins. Numbers indicate the length in amino acids. **(B)** List of candidate proteins identified by mass spectrometry analysis of anti-GFP immunoprecipitants from lysates of WT-GFP and NDC80-GFP schizonts (after 8 hr in culture) and gametocytes (activated for 1 min) (see Fig. S5). Numbers are the total number of peptides identified for each protein. We identify the previously unannotated PBANKA_1442300 gene as coding for a candidate SPC24 ortholog in *P. berghei* (see panel C). Colours correspond to panel A and indicate each of the four subunits of NDC80

complex. **(C)** On the left: presence/absence matrix of NDC80 complex subunits in a large set of (newly sequenced) apicomplexan parasites, various Apicomplexa-affiliated lineages (outgroup e.g. chromopodellids) and a subset of eukaryotes. (see **Table S4** for presence/absence table, and **Sequence File S1** for sequences of NDC80 complex orthologs). Numbers indicate the average length of the orthologs in each collapsed clade represented in the phylogenetic tree. Note the consistently longer orthologs in Sarcocystidae (including *T. gondii*) and the expanded SPC24 ortholog in Plasmodiidae. On the right: workflow for the discovery of SPC24 candidate orthologs in Hematozoa and Coccidia. Alignments represent RWD domains of putative SPC24 orthologs found using seed sequences (top: e.g. TGME49_212800). See **Fig. S6** for full alignments of RWD domains. The panels represent Hidden Markov Models (HMMs) of SPC24-like RWD domains, which were found to be significantly similar (see E-values) using HMM-vs-HMM search algorithm HHsearch (see methods).

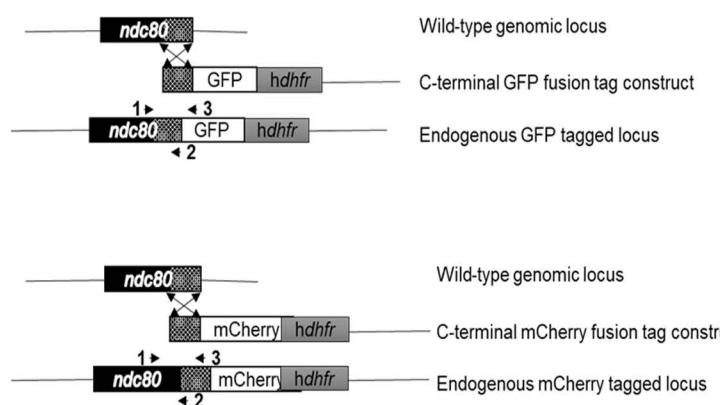
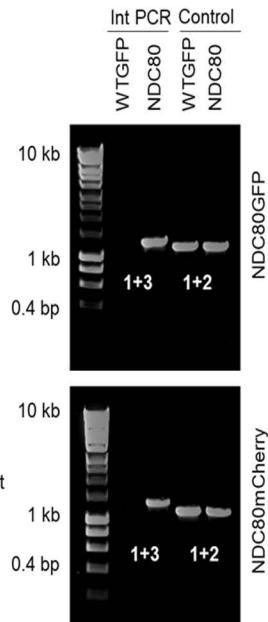
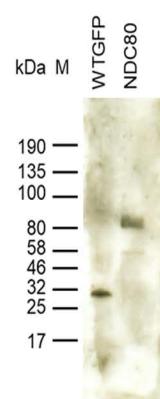
Fig. S1**A****B****C**

Fig. S1: Generation and genotype analysis of NDC80-GFP/mCherry parasite lines. (A) Schematic representation of the endogenous *Ndc80*, the GFP/mCherry-tagging construct and the recombined *Ndc80* locus following single homologous recombination. Arrows 1, 2 and 3 indicate the position of PCR primers used to confirm successful integration of the construct. **(B)** Diagnostic PCR of NDC80-GFP and WT-GFP parasites using primers IntT259 (*NDC80*, Arrow 1) and ol492 (for GFP line)/mCherry (for mCherry) (Arrow 3). IntT259 and T2592 (*NDC80*, Arrow 2) primers were used as control. Integration of the NDC80 tagging construct gives a band of 1269bp (GFP line) and 1335bp (mCherry line) for NDC80 parasite line. For controls, both WT and NDC80 tagged constructs gave a band size of 1153 bp. **(C)** Western blot of NDC80-GFP (96 kDa) and WT-GFP (29kDa) protein to illustrate the presence of intact NDC80-GFP in schizont stage extracts.

Fig. S2

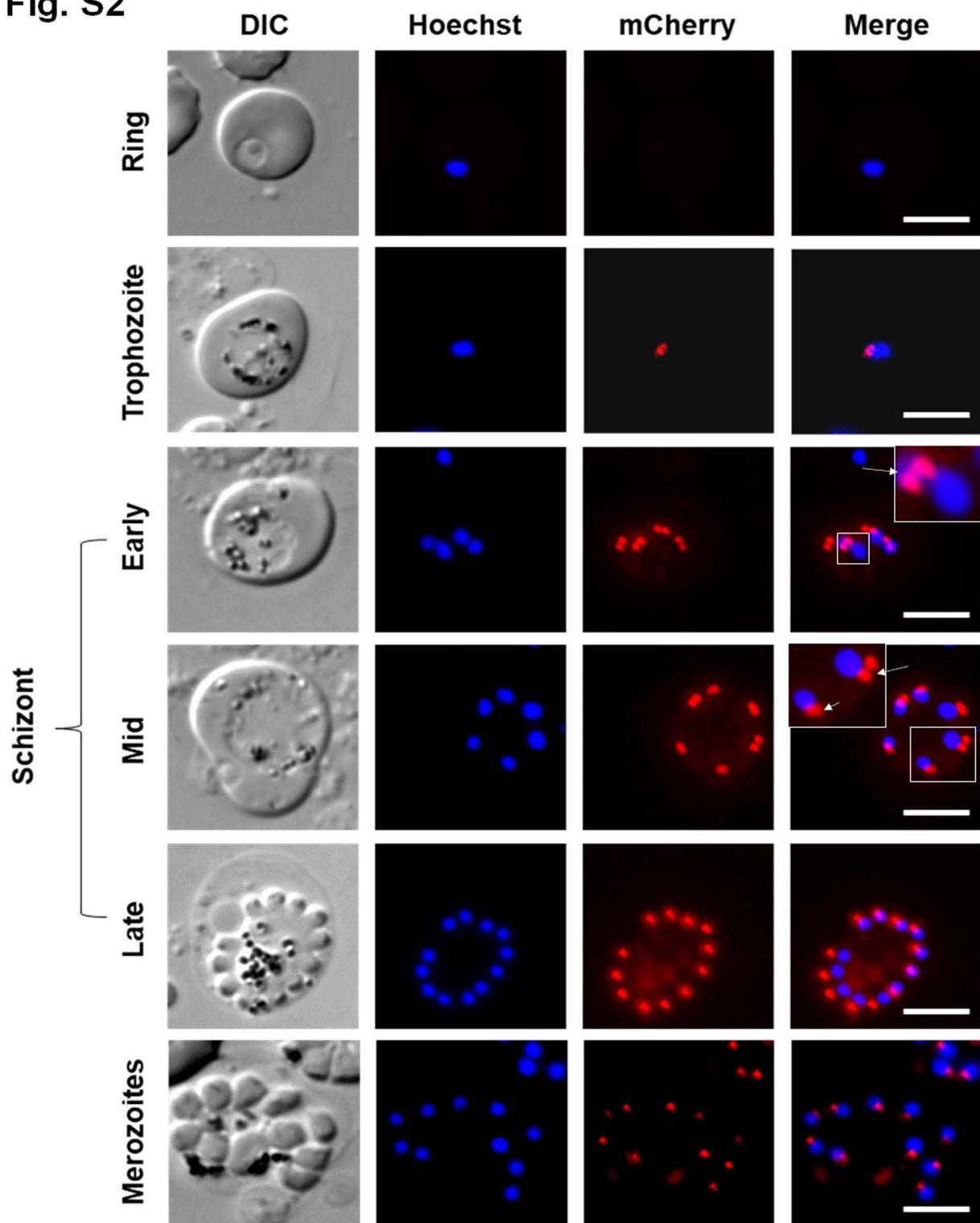


Fig. S2: Expression and location of NDC80-mCherry in asexual blood stages. Fluorescence was detected by live cell imaging. DIC: Differential interference contrast; Hoechst: blue, DNA; mCherry: red, NDC80-mCherry; Merge: Hoechst and mCherry fluorescence. Arrow shows doublets and arrow head shows singlet of Ndc80-mCherry. Scale bar = 5μm.

Fig. S3

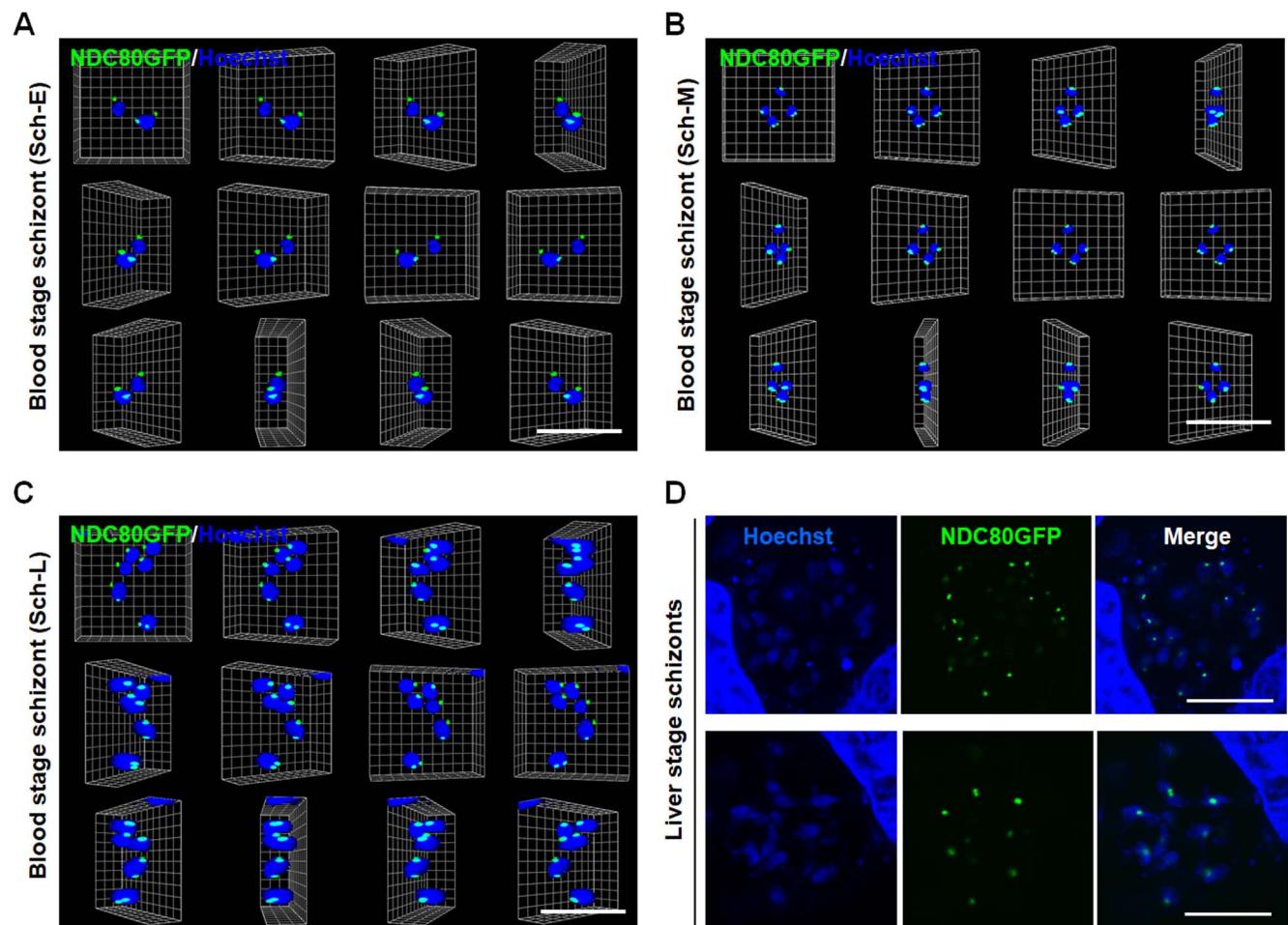


Fig. S3: Localization of *Plasmodium* NDC80-GFP during schizogony in blood and liver stages.

Different views of three-dimensional super-resolution NDC80-GFP images during **(A)** early stage schizont, **(B)** middle stage schizont, **(C)** late stage schizont during blood stage schizogony ; Scale of the grid is 0.5 μm , **(D)** Expression of the protein was detected in liver schizonts by live cell imaging. Merge = DAPI and GFP. Scale bar = 5 μm .

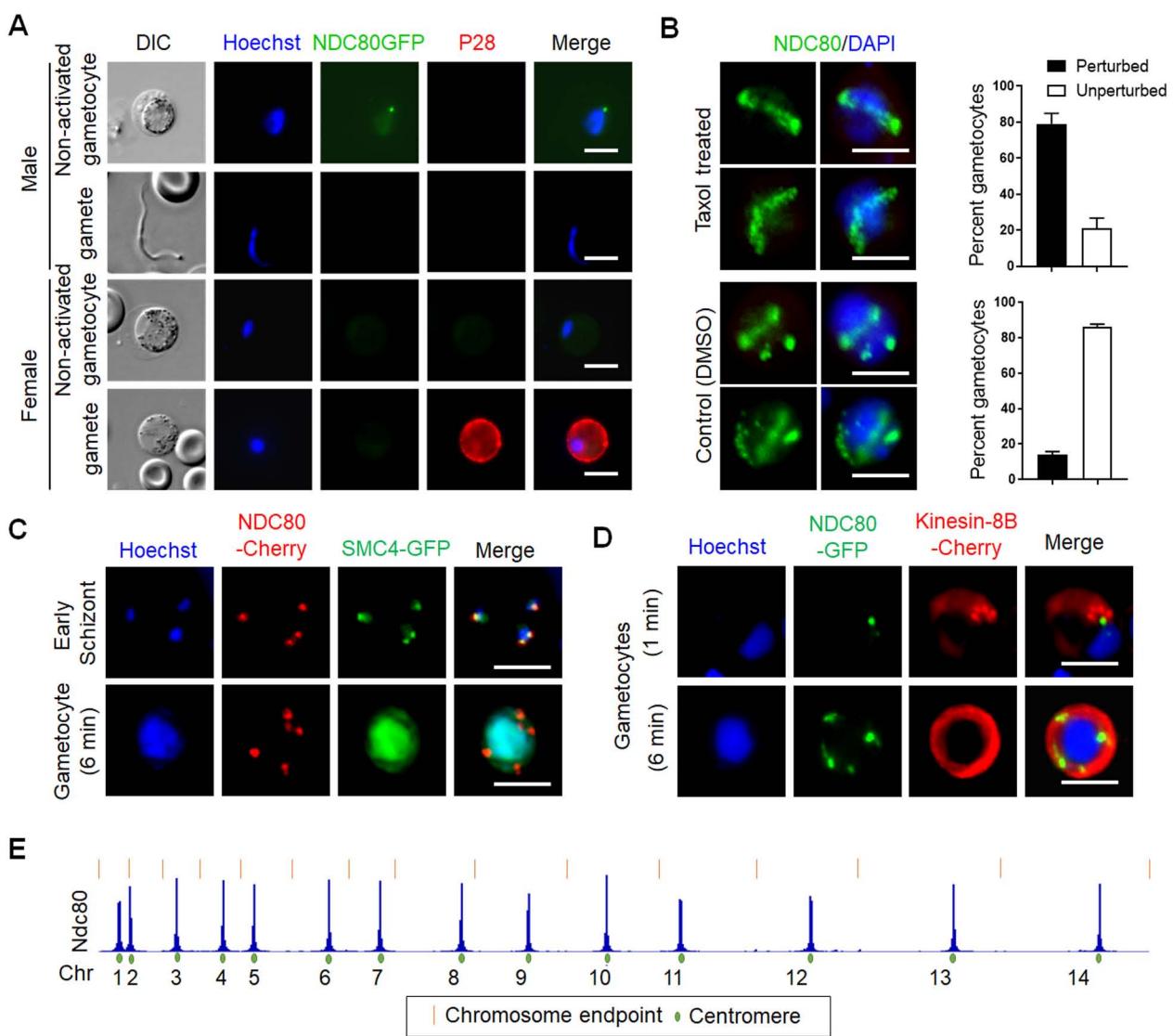
Fig. S4

Fig. S4: NDC80 is not expressed in female gametocytes and gametes but shows centromeric location in male gametocytes. **(A)** Non-activated and activated gametocytes and gametes were examined by live cell imaging (100x magnification). DIC: differential interference contrast; Hoechst (blue, DNA); NDC80-GFP (green, GFP); P28 (red, cell surface marker of activated female gamete); Merge: Hoechst (blue, DNA), GFP (green) and P28 (red). **(B)** Male gametocytes treated with taxol (tubulin depolymerisation inhibitor) at 1 min post activation showing location of NDC80. DMSO was used as control. **(C)** The location of NDC80 (red) in relation to SMC4 (green), a kinetochore marker, in schizont and male gametocyte. **(D)** The location of NDC80 (green) in relation to kinesin-8B (red), an axonemal marker, in male gametocytes at 1 min- and 6 min- post activation. Scale bar = 5 μ m. **(E)** Centromeric localization confirm by ChIP-seq analysis of NDC80-GFP profiles for all 14 chromosomes in gametocyte stage. Signals are plotted on a normalized read per million (RPM) basis.

Fig. S5

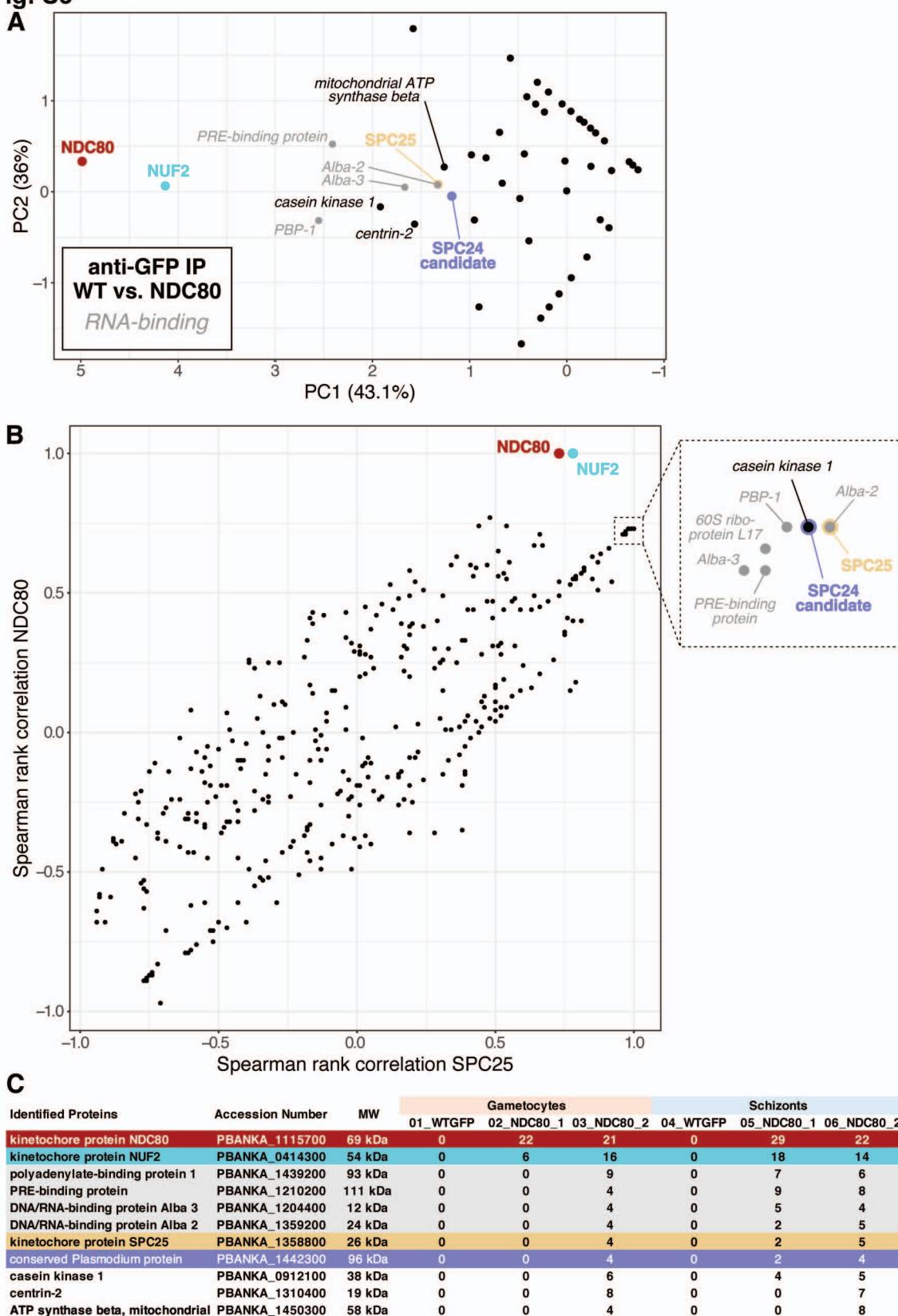
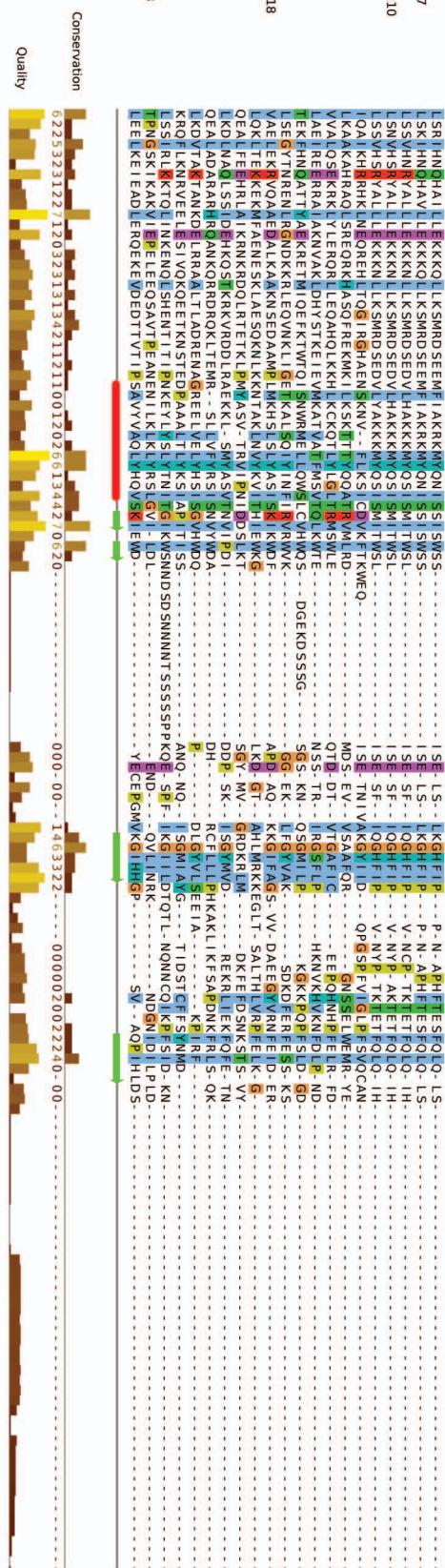


Fig. S5. Covariance, and correlation analysis of proteins detected in GFP pulldown experiments (WT vs. NDC80). **(A)** principal component analysis of natural log-transformed peptide counts of identified proteins in all NDC80-GFP pulldown experiments (schizonts and gametocytes) reported in this study (**Table S2**), excluding ribosomal proteins (often found as background hits). Subunits of the NDC80 complex are specifically highlighted, and colours correspond to those in Figure 6. Percentages indicate the variance explained by each component (PC1: 36%, PC2: 43.1, adding up to a total of 79.1%). Candidate proteins that show similar behaviour in different pulldown experiments are specifically annotated. Grey represents candidates that function in transcription and/or are known to interact with (m)RNA. **(B)** Plot of Spearman rank correlation of NDC80 (y-axis) and SPC25 (x-axis) with all proteins identified across control and NDC80-GFP pulldown experiments. Candidates were selected based on $R>0.7$ for both SPC25 and NDC80. NDC80 complex subunits are specifically highlighted, and colours are similar to panel B. **(C)** Table showing peptide count values in both control and NDC80-GFP pulldown experiments for candidates selected from the analyses in panel A and B.

Fig. S6

<i>Hammonia_jammarum</i>	<i>Neoscyphus_cannini</i> [543-581]
<i>Sarcocystis_neumanni</i> [543-580]	<i>Besiostoma_pessotti</i> [386-731]
<i>Cystoisospora suis</i> [645-802]	<i>Cystoisospora_necatrix</i> [382-548]
<i>Eimeria_acervulina</i> [221-383]	<i>Eimeria_brunetti</i> [383-560]
<i>Eimeria_falcirostris</i> [27-441]	<i>Eimeria_maxima</i> [306-549]
<i>Eimeria_praecox</i> [354-484]	<i>Eimeria_praecox</i> [354-484]
<i>Eimeria_tenella</i> [380-536]	<i>Eimeria_tenella</i> [380-536]



KKAQY DOK-
LKAQY DOK-
UNIQY DOK-
LKKAQY DOK-
LKKKEY DOK-
LKKAY DOK-
KLKAY DOK-
KLKAY DOK-
KLKAY DOK-
KLKAY DOK-
RANKKIDE GLSKFSTI SPDE KIRHL SPFLSALK L-
KIKKCO IDQ KMDSYNHR STONKURRLL FLSALK L-
RGEH KOTTE VEKOHLA RD VROGL EFLVSS TTVS-
EE RHEE KELLSIS ATIE KESADAEEROLLE EFTS S-
TSKOK EKOK FE LOON-1 KIEHMF EFK AYVE
VEEKVKLNLLID DE LANE LINNNNKKR R-
VLLV

DNE	NE	LE	CLI	P	K-	NTEINMMARLQD	KK
DNE	NE	LE	CLI	P	K-	NTETINMMARLQD	KK
DNE	NO	LE	CLIP	-	K-	NTETDMWRLQD	KS
DKE	NE	LEVCLP	K-NTKTMWMLQD	KR			
DTE	NE	LEVCLP	K-NTKTMWMLQD	KR			
DDE	KO	LEVCLP	K-NNTNMWRLQD	KN			
DND	NE	LEVCLP	K-NNTNMWRLQD	KN			
DTE	NE	LEVCLP	K-NNTNMWMLQD	KR			
DNE	NE	LEVCLP	K-NNTNMWMLQD	KR			
DNE	NO	LEVCLP	K-NNTNMWMLQD	KR			
VDE	K-S	NWLALSL	SK-KENETIWRFLSS	K-			
IDQEN	VNLALSL	-EKEKNERVNLALSL					
IDQEN	GNG	ANLALSL	SK-AGSEQIWRFLSS	LR			
VDE	K-S	NWLALSL	SK-KENETIWRFLSS	K-			
RNE	SO	VNLALSL	SK-DRTTITMDYAN	LD			
AND	KE	VNLALSL	PE-DRSTOMDVTVK-T	TS			
VTR	SR	STLVAP	LEKEDKMMI	ID	S		
EEN	DL	WVEALV	NYVSD	SUNTRIP	LS		

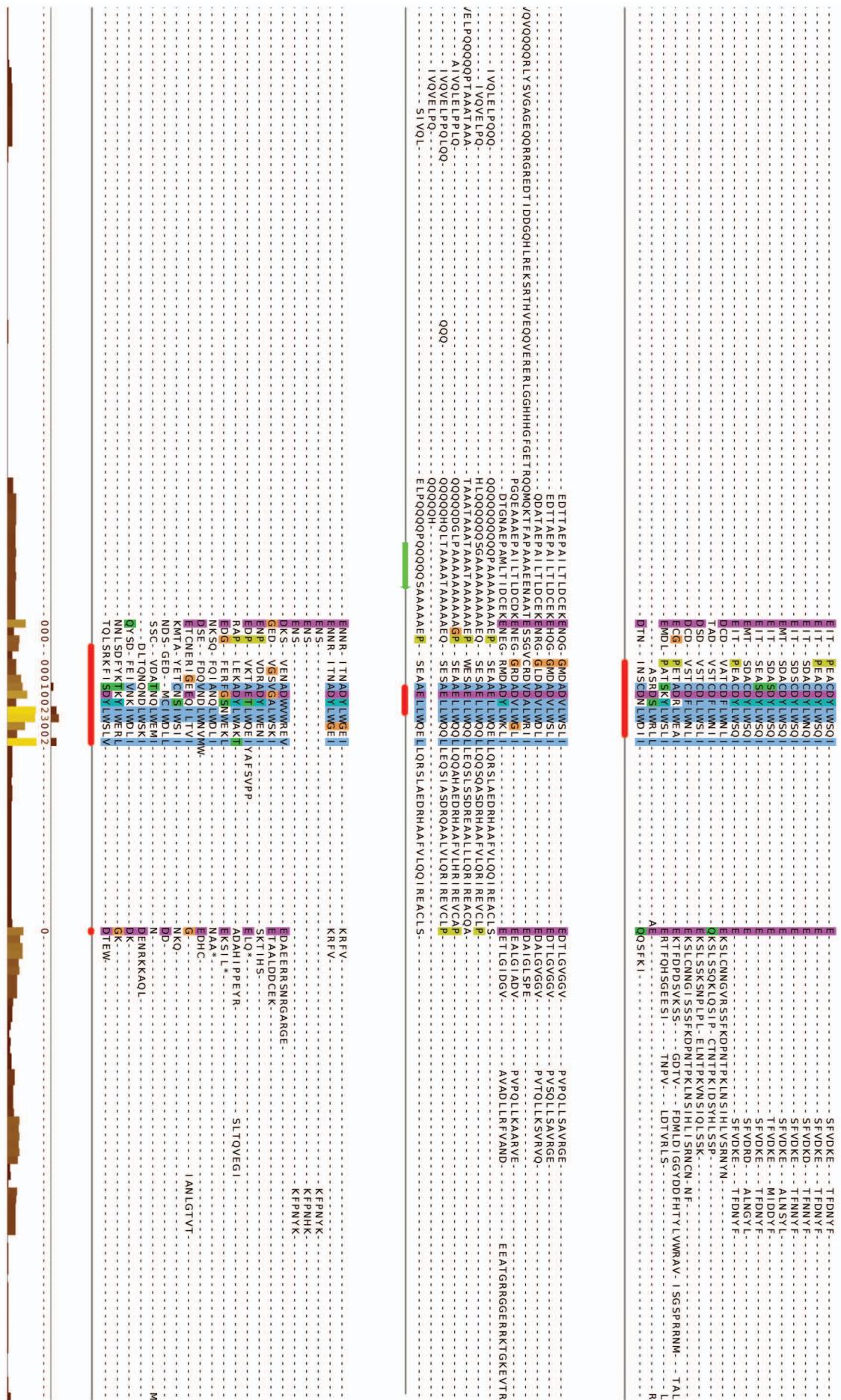
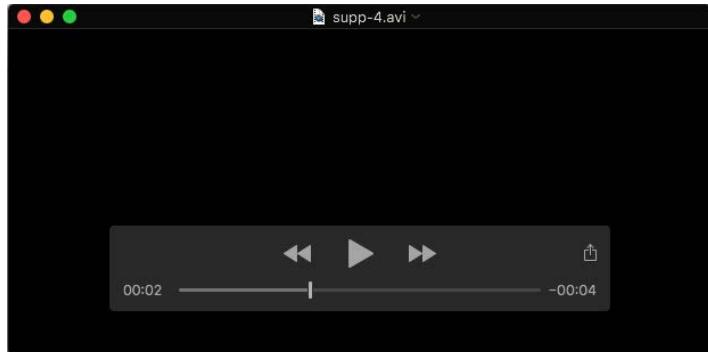
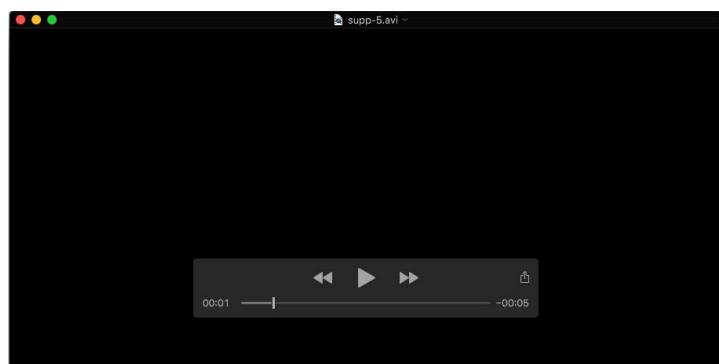


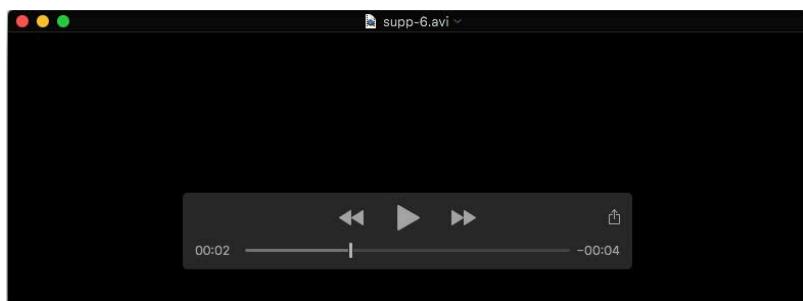
Fig. S6 Full-length alignment of RWD domains of different SPC24 candidate orthologs shown in Fig. 6C. RWD domain alignment of SPC24 candidate orthologs from our sequence database. Secondary structure of the three groups of orthologs (see **Fig. 6C**) is based on the HHsearch algorithm. Colours and conservation metric are based on the Clustal scheme as implemented in Jalview (Waterhouse et al., 2009).



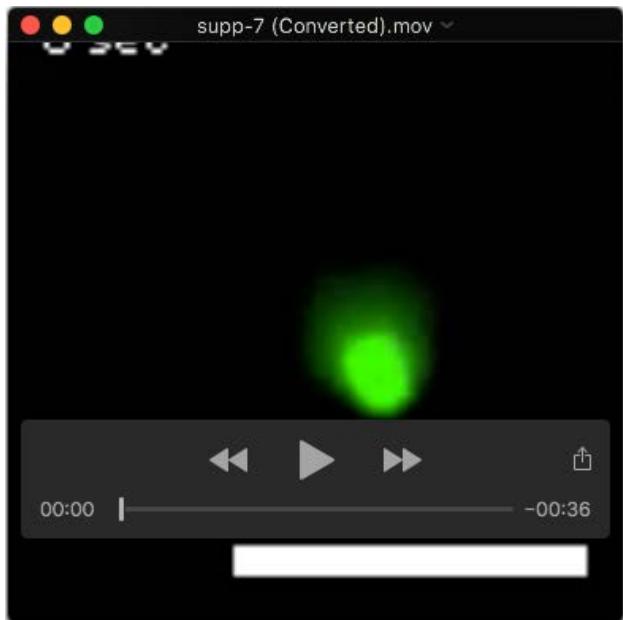
Movie 1. Three-dimensional visualization (3D rendered SIM Structured Illumination) of NDC80-GFP with respect to DAPI-stained nuclear DNA during early (SV1), middle (SV2) and late (SV3) stages of schizogony showing asynchronous division. Scale of the grid is 0.5 μm . See also Fig S3A, B and C.



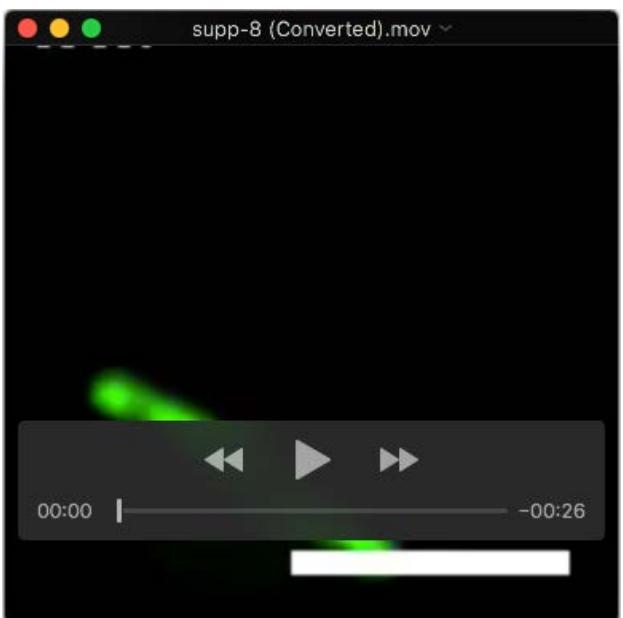
Movie 2. Three-dimensional visualization (3D rendered SIM Structured Illumination) of NDC80-GFP with respect to DAPI-stained nuclear DNA during early (SV1), middle (SV2) and late (SV3) stages of schizogony showing asynchronous division. Scale of the grid is 0.5 μm . See also Fig S3A, B and C.



Movie 3. Three-dimensional visualization (3D rendered SIM Structured Illumination) of NDC80-GFP with respect to DAPI-stained nuclear DNA during early (SV1), middle (SV2) and late (SV3) stages of schizogony showing asynchronous division. Scale of the grid is 0.5 μm . See also Fig S3A, B and C.



Movie 4. Time lapse video showing dynamics of NDC80-GFP during gametogenesis over 0 to 90 seconds after activation. Scale bar = 5 μ m. See also Fig. 2B.



Movie 5. Time lapse video showing dynamics of NDC80-GFP during gametogenesis over 90 to 180 seconds after activation. Scale bar = 5 μ m. See also Fig. 2C.

Table S1: Primers used in this study.

Name	Sequence (5' to 3')	Notes
T2591	CCCC <u>GGTACCGCGAATTCTAATAATAGGTTAAC</u> T	KpnI site <u>underlined</u>
T2592	CCCC <u>GGGCCCTTCAGTTACACTTGATGTAAATTTTATATAATT</u> C	Apal site <u>underlined</u>
IntT259	GATCTGGAAGAACATCAATCAGAAAGAC	
ol492	ACGCTGAACTTGTGGCCG	For GFP line
mCherry	TTCAGCTTGGCGGTCTGGGT	For mCherry line

Table S2: List of main protein hits identified by mass spectrometry in the NDC80-GFP immunoprecipitation experiments. The two last columns represent Spearman rank correlation values of a particular gene product with NDC80 and SPC25 (see also **Fig. S5**). Colour coding corresponds with **Fig. 6** and **Fig. S5**. (separate excel sheet)

[Click here to Download Table S2](#)

Table S3: Sources of genomes and transcripts of the sequence database used for sequence similarity searches including hyperlinks and/or reference to papers. (separate excel sheet).

[Click here to Download Table S3](#)

Table S4: Presence/absence table of the NDC80 complex (NDC80, NUF2, SPC25, SPC24). Absences are denoted by '-' and presences by the length of the ortholog. (separate excel sheet)

[Click here to Download Table S4](#)

Supplementary Sequences:

Sequence File S1: (candidate) orthologs of (including length per sequence in the FASTA headers): *PbNDC80* (PBANKA_1115700), *PbNUF2* (PBANKA_0414300), *PbSPC25* (PBANKA_1358800), *PbSPC24* (PBANKA_1442300).

Supplementary Sequence File 1 (File S1)

>SPC25_Plasmodium_berghei (PBANKA_1358800) [216]
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EFSNIQISDPKKYRASITLHDGMYEAVETIPRINKFEDYVNGLNRLPFTTFCCLLRKSFKELQ
>SPC25_Plasmodium_chabaudi_chabaudi [216]
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>SPC25_Plasmodium_gallinaceum [216]
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>SPC25_Plasmodium_vivax [216]
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>SPC25_Theileria_annulata [259]
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>SPC25_Theileria_equi [273]
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>SPC25_Theileria_orientalis [231]
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>SPC25_Theileria_parva [219]

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>SPC25_Cytauxzoon_felis [264]

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KKLLQLEVKVLGLKFINEDGSLKMIFTNLSITDPHKKCSFTIKLTNDRTGINCDPPIKQFDSDLICDLNNRLEFGL
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>SPC25_Nephromyces [248]

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>SPC25_Toxoplasma_gondii_ME49 (TGME49_232430)[432]

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>SPC25_Eimeria_mitis [244]

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>SPC25_Eimeria_necatrix [287]

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HALLAGS

>SPC25_Eimeria_acervulina [355]

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>SPC25_Eimeria_falciformis [343]

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>SPC25_Eimeria_maxima [256]

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>SPC25_Eimeria_praecox [221]

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>SPC25_Eimeria_tenella [287]

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>SPC25_Cyclospora_cayetanensis [572]

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ADCRERASRGTRQRAEFAKFIDRQMAELEELATESARLQAEVKELNTKTEQNLQEVEALLEEDKASCPQF
MVVSQRQLHARKEWRSRVDAYKFTHGIRIHQQRGKPFTRICFLGLYKELAQLECHLDVSLEDQSCQEPTG
SHKGGLSGAASDLIAVSEADIQCIIHDPKNASPPYAPAFLVSDCMYLDLEFALALVRGSTLISSLDSMLSFSRLC
LHLILAKPLLPLCPRNPHLAAAAPVSSATAAHDCIAIPLFWDVPVMYSLARVLFDLFFCDSVVGAWPEAP
LSELCALYAEGRVSFTALVSHSRLIFKAQLTRMSHAQKLVALHEQQQRQAEEKQEEAHKLQHQIHTGPL
SDD

>SPC25_Eleutheroschizon_duboscqi [257]

MAAADVSAADHVAATVLARTPPTPVDQLAPRCQGLYDALKQDTTELLLDGDQTRLTGTQQKLLKAIAQIRE
VKLATQTNLTKASELSERTRENKRTVKQLQLEEKELDRQLGELPTALSALELKKEKARKEKELSDIRSQYMVVK
KQGVHQEDQWADECKSYQRAFGISYHLEDNLHKIFFNLSPDEGKDLECVFEFRLLEDRKGTGAPGSLKLS
DTRCDPPVSDFRELLADFNANMSTFPQLLLLRLDAFHAGLGRG

>SPC25_Cryptosporidium_andersoni [257]

MDFFLKKKGIFRISIDKLNSRILYLTFIIMDSHVLKSNILNEEVTNLENKVQEWCNNINNSIANMGDFENN
NINLNFIREAKVRIDGLQKRIGDSKELFEIRRVMESEQEQYQKHLNTNLQNMLEVKQKLEMRIAEDRNRLEG
EKTAYENDLHIAKMQSSEREKGCRYSDNLGLYINKVNGNNRFTFIYIDANNPLREFYFDLYYNEKTSSYNG
VHCEPYIHEFQDLTNKLNNRSICLRKFISNMRQAFKDSL

>SPC25_Cryptosporidium_hominis [228]

MEAHSIISNEISEEIQDLKNKLLEWTSGINGMLELGVEKLNDQVSRDINMIKDSSIRIDGLEKRIKDTRELSEIK
KVVIEKEEFKKCFNTNTIALSEKLKRLEKQLQEDRQRFEQEKLVYEADLKVAQQSLEREKSCRIYSENLG
LNIGKVNGNNRFTFTCIDENFPEREFFFDLFYNEKAQLYEGIGCNPQVSDFKPNIELLNSKSITFRQFICKMR
KSFQQLIK

>SPC25_Cryptosporidium_meleagridis [228]

MEAHSIISNEISEEIQDMKNKLLEWTNGINGMLELGVEKLNDQVSRDINMIKDSSIRIDGLEKRIKDTQELSEIK
KVVIEEEEFKKCFNTNTIALSEKLRRLERKQLQEDRQRFEQEKLVYEADLKVAQQSLEREKSCRIYSENLG
LNIGKVNGNNRFTFTCIDENFPEREFFFDLFYNEKAQLYEGIGCNPQVSDFKSNVELLNSKSITFRQFICKMR
KSFQQLVK

>SPC25_Cryptosporidium_muris [227]

MDSHVLKSNILNEEVTNLENKVQEWCNNINNSIANMGDFENNINLNFIREAKVRIDGLQKRIGDSKELF
EIRRVMESEQEQYQKHLNTNLQNMLEVKQKLEMRIAEDRNRLEGEKTAYENDLHIAKMQSSEREKSCRIYS
DNLGLYINKVNGNNRFTFIYIDANNPLREFYFDLYYNEKTSSYNGVHCEPYIHEFQDLTNKLNNRSICLRKFIS
NMRQAFKDSL

>SPC25_Cryptosporidium_parvum [228]

MEAHSIISNEISEEIQDLKNKLLEWTSGINGMLELGVEKLNDQVSRDINMIKDSSIRIDGLEKRIKDTQELSEIK
KVVIEKEEFKKCFNTNTIALSEKLKRLEKQLQEDRQRFEQEKLVYEADLKVAQQSLEREKSCRIYSENLG
LNIGKVNGNNRFTFTCIDENFPEREFFFDLFYNEKAQLYEGICCNPQVSDFEPNIELLNSKSITFRQFICKMR
KSFQQLIK

>SPC25_Monocystis_agilis [208]

MGSLRVLQRACKTLSQKGARDAVETVTICQETFSVSSHMSEIVRVQIEETKRAVEQMCFINESMKEHHEKV
EKQLKTELETLSKYEGKLESEKDRMAESETKLSVCREQHEASQDQLIKERHRLQLFLGLVTGMEDAGQVV
CLKFIDPNPQREFKFTIRRNNIICVPQIAAVEEISNNLKNCRITLTEAVCLMRQAFRQSV

>SPC25_Lecudina_tuzetae [226]

MATISTKPDDLQLRIQKQEQRFARFENEILQCANAHAEKWCEVQEEGLIVHQVEELCRRKKQDEEALAE
VLQCFKVEGEVEEWTKISSNKAQVEALSAQLAEEEEAIARIQQEIDKTVGDDAQLDRERQEIEEYERCLGI
AIHRTGGATKFIFTRLDRSSPQREFSFLLKLLGNAFKVFGTQPEIEGVKDVEEINQNGGRLTLSAMRRR
FVEAVAEGCV

>SPC25_Pterospora_schizosoma [256]

GINAEYMGGIRFVTTQYTAMNQHVHTNSIEEESRIIFQTLTKEIEDIHNSIDTWNNVTTKREQYAKLWLEIQ
KSGLEVLFKKNSKKIKEDQKTLLEVLKCHKVVEAEVALLKEMPDEERLQLLEQFNSLSNKIERLQME
NAKIHSECESIEHQTEKMLFYKRFPGIEIRRPQQQTVFIFTNIDPYNPSREFMFSLKISSENNLTVVDCEPK
VGTLKSLTDELNSNGLEIFVYNLRTKFKQFALQQHTI

>SPC25_Selenidium_pygospionis [229]

MQTEQWCDEVKQEIGILYQQFSKTEVCKKTLFEEKNAFNEEVEERKQAIDSLKKRREELLKKELMEKTM
KREEDEQAEIKELVKALERQVAELEEEARKKEEEAABLGEERRKLCVRQEEIQHEKNRSSAAAKTELKFYR
DYMGLAIEPIKENQIKFVFTMLDKNNLCREFWVSDITDTIRVLECEPPIKDIDALVAATDRENVISFLKRARA
CFVLSLEEKTEQ

>SPC25_Vitrella_brassicaformis [334]

MAPQDGAFIRQLLASQQQLKDALQEEVEVAQKVRDQCLKIAEVGKQRCQDSKTTKQGVLGIEIQKEVFAYR
TKLQEYQAKEGLFQQKEADFQQQLSDVLSKLERLPEDVRNAETETGHMKAKVERAKKVVYEREKAKMVER
HAEYEVLISSFQRKLGLKLSRVADSVKGHVRVIFTNLHPADDKEYAFELYLKDDETEYKVLLCDPPIADLP

ALVTALNAKQIEFGAFVCIIRTRFKAIASTQPHELQQPEATEEEQEQQEKEKPEQESPPAAAAAGGGGGAA
MNLPRFRIPENGDESPPMHQGGLLSRLTGGLLRRSTAAAKRGDGEQDDSV
>SPC25_Chromera_veila [421]
MSLGATVLHLKQENWFFFKAERHEQICEALSRDYSTCAPLFDKLSGACDSAAGQIAKQQEAENDRRDT
VVKMREALKRLDEQLEGQPKDHREAEVFKEAESRLDMENKALKLQEERIRAQHQTVAHLKRTEFSGL
RIEKSRAVRYVFSHLSERLPELECHMDLRLEVQTDQSQKETRSYSIVSCSPPISWSPATSKASTDLSRLL
SDLAKKVVDLACLTAVVRQRFRDFVRREDEEAEKERNKQPPATQLSNRSALRPPRHSTVKITGRPSQVD
GSASTPTPSPPVASSPQGTLGMCRASVQAQAAAAPLPGCSSSTPEKSRGAGGTPEEAESRTPVPPVPPR
VRVASAKTWTSSPVSAAERPPGTERESETVLLHGDIEEGRTEGLDGAPSSGGPARPPLSTLMEMDDY
>SPC25_Colpoda_aspera [235]
MKGMDLIKQKSQLLNQTTIVFGKNSITKLNQKSLQEESTLKKFETGFLQCQEVLLENANKTNEALMASAKEA
RSELDVQKDIVGFEEDYIEAMKENEIRRLEDNKTRYRDLLANLNEELEKERQEHNQACAHSAHFEALTAI
DQYRNNLIGISFEKTEDKQLVIFFEAVSRQPGYDLCRIDLKLDDEDDGSFIVTRTEPEAPAIAEFAEELSEDRN
LRFLVKSKEFKVRAEQ
>SPC25_Colpodella_angusta [288]
MTELNSKRLQQLVERSDMMVDGVLSSNDQFIAQLQKRREALERQVEILEPLSVTCAEARSNQRQIQDEIQ
TSEPERNRNLNEAQLRLEVENLPKLIAAQEKIFAEARKKLDNVSNHTMMLEKYRREERMLNDELNVYET
QAGIRIVRDHGSLKYIFNIDQNEYYMLMSMEICADGRKIYRLLDCRPSINIADVISYLNEGKLKIGGCLAI
FKQAAGIRVNNTSLSALHSSQYRMNDIFHGNDNSQFNKQLQQQQHHHQHSNFPSGGIARGSILKP
VEMMP
>SPC25_Colpodella_sp [288]
MTELNSKRLQQLVERSDMMVDGVLSSNDQFIAQLQKRREALERQVEILEPLSVTCAEARSNQRQIQDEIQ
TSEPERNRNLNEAQLRLEVENLPKLIAAQEKIFAEARKKLDNVSNHTMMLEKYRREERMLNDELNVYET
QAGIRIVRDHGSLKYIFNIDQNEYYMLMSMEICADGRKIYRLLDCRPSINIADVISYLNEGKLKIGGCLAI
FKQAAGIRVNNTSLSALHSSQYRMNDIFHGNDNSQFNKQLQQQQHHHQHSNFPSGGIARGSILKP
VEMMP
>SPC25_Piridium_sociabile [356]
MNTKWLPVFALLISPREGGTVLSDRKRCGVSFVPHALRRWRRFGSSRMHDAAPAVISNRSANKFLHVV
YRVDLEKTVEFYTKVLGMTVLRRDDPVGRYTNFLGYGTQKNNKHFAIELTYNYGVREYRQAKLGFLG
LGLVRENLTETGVAVAHHGSVLKDVADEITPSQFPDEPAEQYRIWMATIEDPNGYPIELVEKSQADSM
HRVRLHVSTMKSIEFYTQVMGMLLKRSMPLPKWPAISSWLGYGIVGDDFAVPILELVYEYSTEKLDVG
GYGHЛАISTPDLKTSLDNIRDKGKIVKEVTVPKGITKIAACLDPDGYKIVLVDAQDFDKELACEKIKTEIA
>SPC25_Digyalum_oweni [241]
IRIVVCDKIKDQTRIRSRALAVMDALRRVASQIEDECNRISDRLLTICEKKLQQAQLNKENMEKEI
QIHARAQ RDMVAQLSEIEVQNVKTKDMLKHQSQETESRRTELRLQERLASLSIDD
KEDQISKAP
EEDNSEAVLKALH QEE
LTYTRLFGLRLGQASSGGTKVFT
TSINPLDPEEEFSFILVPGKRM
EAVHF
SHRIDNERAA
ILDSELNREP
RCKSRLAWFIAQIRKEFKLCEV
>SPC25_Platyptreum_vivax [229]
MTSSEYIKSLRRAYMHIDEECTRINDRVNIITQKLENITLHKH
HNKIIENLSGEQSDIVNQISDIEVQQLKLN
DLFSYRLEHDAKRNELKVLQERLASLVL
YDKKTN
KENKEIVDN
KEDSSM
LENMQMIE
FYERIFGIRL
GQA VSGG
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SLLNE
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>SPC25_Perkinsus_marinus [215]
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>SPC25_Arabipdosis_thaliana [315]

MEQISNIAGGDTTKE¹MASLG²LICEKDIHEQRLKIDSFIASP³FRRSMNSLVERAQATAQSQVELMNLKADLR
EAEDELVKVLAVKTRKEARQMGIRDSISATQSRIEV⁴RRNLQLQSKSKDDSVRIISQQQLQALSKSKDNAGKV
TEDKADIHEAISWYNHALGFHVEAGHG⁵VKFTFTNIDAKRPTREFSFTVHYGNDIYTLLSDLQLDYINEMVQ
ELNKTNDLFRFVR⁶LMREQFLKSTLSELPTHSGQLQQETSAISASAPAISFSTD⁷NMSTPENKR⁸SKVQVNRR
QKRGSESP⁹LAPVSTSATRRSSRFKGKK

>SPC25_Giardia_intestinalis [240]

MEDEQKSAAERLVSLADTLTISLNTFVTKNLDAISNMGSTFISFVDETLHLLKKSKDDYEERLKQELEVEKLS
TSASEEEQKLNAQLARARTQLDTLKQQYSIMQEEYRKALADFEERRVAFEALPATQKAHV¹⁰KEDLEWRLR
NYE¹¹MLRMKIEQRDENSIVIFWGLNP¹²ADESQQYSFRLITREDGEIII¹³EPTIEIANLDLFLSDAKITGNIP¹⁴LLIRR
IRLSFLQLAECE¹⁵DSES¹⁶VTQD

>SPC25_Trichomonas_vaginalis [212]

MENDSINESLKHFLDAIAESKD¹KLNK²FVTQQTDLMKKRQMINNLKTD³MTRNERNLKDEQRLVMENIKRIKN
QNENISVQNK⁴KQQIEIKLNNNDIQTIPK⁵NFEDYQV⁶KIRLLETTANH⁷I⁸GKEKEKADEFYSSYQ⁹TDSL¹⁰LYEALYGI
KISISNGKTKFVFLPQSYEVELDCIQNNF¹¹VLSNSTLP¹²SSIATTILVDEFNN¹³SRDLFSFLNSIRQQI

>SPC25_Naegleria_gruberi [236]

MFQE¹VIEIHDRVAQKTNQFEQNF²KGFINSQINDNNK³RHD⁴FEQNCQNFQSEFQSL⁵TRVERGINERQHVEE
IRKRYCEDEGELDQR⁶ITDMK⁷VRADEQPK⁸I⁹E¹⁰SKQ¹¹KLLQLK¹²SEHEEKSNMLNR¹³VESDFVDHATKKFIEKEKY
RTLLGMTFEARQDG¹⁴DIAYLLISFSI¹⁵QNMQALGLTSCQIALRIDGG¹⁶LYS¹⁷VAGCQPAI¹⁸QYDDL¹⁹IASLNDNKNFGS
FIILVRNRF²⁰KNL²¹INQHKPRK

>SPC25_Dictyostelium²² discoideum [324]

MEQQQLK¹TKNGQK²TKPKKK³TS⁴TI⁵VQ⁶QDN⁷NNMENSGYNSQNDSEFD⁸P⁹SSSSL¹⁰SSSSQQIVNNNQ
PLITNITTP¹¹NNNNNNNNNNNNNNQILED¹²IYSDEYCENQM¹³KCMEGFI¹⁴KKW¹⁵KDEYIGSI¹⁶KGN¹⁷VEHHQ¹⁸I¹⁹ELNNYTF
QLSELKQERLFLQQYKES²⁰EQRTIELQ²¹IIQLQ²²RINNEIESIEKNRH²³Q²⁴L²⁵PIDLESKKL²⁶KY²⁷QEETQR²⁸IRL²⁹TS³⁰TI³¹K³²SIE
DQT³³IKQLLD³⁴LD³⁵YPL³⁶TLFK³⁷NYL³⁸GLEFQ³⁹KLS⁴⁰NGD⁴¹GL⁴²KL⁴³IFT⁴⁴K⁴⁵DRNN⁴⁶H⁴⁷N⁴⁸REFT⁴⁹ISITVDNL⁵⁰T⁵¹DQY⁵²IL⁵³V⁵⁴NCNP⁵⁵MIS⁵⁶DL
DQS⁵⁷IK⁵⁸KL⁵⁹NETG⁶⁰NFSFFV⁶¹K⁶²S⁶³I⁶⁴R⁶⁵K⁶⁶Q⁶⁷F⁶⁸V⁶⁹N⁷⁰K⁷¹T⁷²I⁷³H⁷⁴K⁷⁵

>SPC25_Saccharomyces_cerevisiae [221]

MASIDAFSDLERRMDGFQKDVAQVL¹RQQN²HARQQLQQFQAEMRQLHNQHQHL³IDE⁴LQRLATQRT⁵ALQQ
QIHAQQATNTTREQWRSYHERE⁶SELSRRQ⁷STLAQSREL⁸DS⁹LLQ¹⁰Q¹¹R¹²G¹³K¹⁴E¹⁵C¹⁶V¹⁷Q¹⁸L¹⁹R²⁰A²¹Q²²S²³G²⁴N²⁵D²⁶A²⁷E²⁸V²⁹LY³⁰FLVV³¹ARD³²M³³L³⁴ASL³⁵

>SPC25_Homo_sapiens [224]

MVEDELALFDKSINEFWNK¹FKSTDTSCQMAGLRDTYKDS²IK³FAE⁴KLS⁵V⁶K⁷KEE⁸ER⁹M¹⁰V¹¹EM¹²F¹³EY¹⁴Q¹⁵NQ¹⁶IS¹⁷RQ¹⁸NKLIQ¹⁹EKKD²⁰NLLK²¹LIAEV²²KG²³KK²⁴QE²⁵LEV²⁶L²⁷T²⁸AN²⁹I³⁰Q³¹D³²L³³K³⁴EE³⁵Y³⁶R³⁷K³⁸K³⁹E⁴⁰T⁴¹ST⁴²A⁴³E⁴⁴R⁴⁵L⁴⁶K⁴⁷R⁴⁸L⁴⁹Q⁵⁰K⁵¹S⁵²A⁵³D⁵⁴L⁵⁵Y⁵⁶K⁵⁷D⁵⁸R⁵⁹L⁶⁰G⁶¹L⁶²G⁶³L⁶⁴Y⁶⁵Y⁶⁶Y⁶⁷Y⁶⁸Y⁶⁹Y⁷⁰Y⁷¹Y⁷²Y⁷³Y⁷⁴Y⁷⁵Y⁷⁶Y⁷⁷Y⁷⁸Y⁷⁹Y⁸⁰Y⁸¹Y⁸²Y⁸³Y⁸⁴Y⁸⁵Y⁸⁶Y⁸⁷Y⁸⁸Y⁸⁹Y⁹⁰Y⁹¹Y⁹²Y⁹³Y⁹⁴Y⁹⁵Y⁹⁶Y⁹⁷Y⁹⁸Y⁹⁹Y¹⁰⁰Y¹⁰¹Y¹⁰²Y¹⁰³Y¹⁰⁴Y¹⁰⁵Y¹⁰⁶Y¹⁰⁷Y¹⁰⁸Y¹⁰⁹Y¹¹⁰Y¹¹¹Y¹¹²Y¹¹³Y¹¹⁴Y¹¹⁵Y¹¹⁶Y¹¹⁷Y¹¹⁸Y¹¹⁹Y¹²⁰Y¹²¹Y¹²²Y¹²³Y¹²⁴Y¹²⁵Y¹²⁶Y¹²⁷Y¹²⁸Y¹²⁹Y¹³⁰Y¹³¹Y¹³²Y¹³³Y¹³⁴Y¹³⁵Y¹³⁶Y¹³⁷Y¹³⁸Y¹³⁹Y¹⁴⁰Y¹⁴¹Y¹⁴²Y¹⁴³Y¹⁴⁴Y¹⁴⁵Y¹⁴⁶Y¹⁴⁷Y¹⁴⁸Y¹⁴⁹Y¹⁵⁰Y¹⁵¹Y¹⁵²Y¹⁵³Y¹⁵⁴Y¹⁵⁵Y¹⁵⁶Y¹⁵⁷Y¹⁵⁸Y¹⁵⁹Y¹⁶⁰Y¹⁶¹Y¹⁶²Y¹⁶³Y¹⁶⁴Y¹⁶⁵Y¹⁶⁶Y¹⁶⁷Y¹⁶⁸Y¹⁶⁹Y¹⁷⁰Y¹⁷¹Y¹⁷²Y¹⁷³Y¹⁷⁴Y¹⁷⁵Y¹⁷⁶Y¹⁷⁷Y¹⁷⁸Y¹⁷⁹Y¹⁸⁰Y¹⁸¹Y¹⁸²Y¹⁸³Y¹⁸⁴Y¹⁸⁵Y¹⁸⁶Y¹⁸⁷Y¹⁸⁸Y¹⁸⁹Y¹⁹⁰Y¹⁹¹Y¹⁹²Y¹⁹³Y¹⁹⁴Y¹⁹⁵Y¹⁹⁶Y¹⁹⁷Y¹⁹⁸Y¹⁹⁹Y²⁰⁰Y²⁰¹Y²⁰²Y²⁰³Y²⁰⁴Y²⁰⁵Y²⁰⁶Y²⁰⁷Y²⁰⁸Y²⁰⁹Y²¹⁰Y²¹¹Y²¹²Y²¹³Y²¹⁴Y²¹⁵Y²¹⁶Y²¹⁷Y²¹⁸Y²¹⁹Y²²⁰Y²²¹Y²²²Y²²³Y²²⁴Y²²⁵Y²²⁶Y²²⁷Y²²⁸Y²²⁹Y²³⁰Y²³¹Y²³²Y²³³Y²³⁴Y²³⁵Y²³⁶Y²³⁷Y²³⁸Y²³⁹Y²⁴⁰Y²⁴¹Y²⁴²Y²⁴³Y²⁴⁴Y²⁴⁵Y²⁴⁶Y²⁴⁷Y²⁴⁸Y²⁴⁹Y²⁵⁰Y²⁵¹Y²⁵²Y²⁵³Y²⁵⁴Y²⁵⁵Y²⁵⁶Y²⁵⁷Y²⁵⁸Y²⁵⁹Y²⁶⁰Y²⁶¹Y²⁶²Y²⁶³Y²⁶⁴Y²⁶⁵Y²⁶⁶Y²⁶⁷Y²⁶⁸Y²⁶⁹Y²⁷⁰Y²⁷¹Y²⁷²Y²⁷³Y²⁷⁴Y²⁷⁵Y²⁷⁶Y²⁷⁷Y²⁷⁸Y²⁷⁹Y²⁸⁰Y²⁸¹Y²⁸²Y²⁸³Y²⁸⁴Y²⁸⁵Y²⁸⁶Y²⁸⁷Y²⁸⁸Y²⁸⁹Y²⁹⁰Y²⁹¹Y²⁹²Y²⁹³Y²⁹⁴Y²⁹⁵Y²⁹⁶Y²⁹⁷Y²⁹⁸Y²⁹⁹Y³⁰⁰Y³⁰¹Y³⁰²Y³⁰³Y³⁰⁴Y³⁰⁵Y³⁰⁶Y³⁰⁷Y³⁰⁸Y³⁰⁹Y³¹⁰Y³¹¹Y³¹²Y³¹³Y³¹⁴Y³¹⁵Y³¹⁶Y³¹⁷Y³¹⁸Y³¹⁹Y³²⁰Y³²¹Y³²²Y³²³Y³²⁴Y³²⁵Y³²⁶Y³²⁷Y³²⁸Y³²⁹Y³³⁰Y³³¹Y³³²Y³³³Y³³⁴Y³³⁵Y³³⁶Y³³⁷Y³³⁸Y³³⁹Y³⁴⁰Y³⁴¹Y³⁴²Y³⁴³Y³⁴⁴Y³⁴⁵Y³⁴⁶Y³⁴⁷Y³⁴⁸Y³⁴⁹Y³⁵⁰Y³⁵¹Y³⁵²Y³⁵³Y³⁵⁴Y³⁵⁵Y³⁵⁶Y³⁵⁷Y³⁵⁸Y³⁵⁹Y³⁶⁰Y³⁶¹Y³⁶²Y³⁶³Y³⁶⁴Y³⁶⁵Y³⁶⁶Y³⁶⁷Y³⁶⁸Y³⁶⁹Y³⁷⁰Y³⁷¹Y³⁷²Y³⁷³Y³⁷⁴Y³⁷⁵Y³⁷⁶Y³⁷⁷Y³⁷⁸Y³⁷⁹Y³⁸⁰Y³⁸¹Y³⁸²Y³⁸³Y³⁸⁴Y³⁸⁵Y³⁸⁶Y³⁸⁷Y³⁸⁸Y³⁸⁹Y³⁹⁰Y³⁹¹Y³⁹²Y³⁹³Y³⁹⁴Y³⁹⁵Y³⁹⁶Y³⁹⁷Y³⁹⁸Y³⁹⁹Y⁴⁰⁰Y⁴⁰¹Y⁴⁰²Y⁴⁰³Y⁴⁰⁴Y⁴⁰⁵Y⁴⁰⁶Y⁴⁰⁷Y⁴⁰⁸Y⁴⁰⁹Y⁴¹⁰Y⁴¹¹Y⁴¹²Y⁴¹³Y⁴¹⁴Y⁴¹⁵Y⁴¹⁶Y⁴¹⁷Y⁴¹⁸Y⁴¹⁹Y⁴²⁰Y⁴²¹Y⁴²²Y⁴²³Y⁴²⁴Y⁴²⁵Y⁴²⁶Y⁴²⁷Y⁴²⁸Y⁴²⁹Y⁴³⁰Y⁴³¹Y⁴³²Y⁴³³Y⁴³⁴Y⁴³⁵Y⁴³⁶Y⁴³⁷Y⁴³⁸Y⁴³⁹Y⁴⁴⁰Y⁴⁴¹Y⁴⁴²Y⁴⁴³Y⁴⁴⁴Y⁴⁴⁵Y⁴⁴⁶Y⁴⁴⁷Y⁴⁴⁸Y⁴⁴⁹Y⁴⁵⁰Y⁴⁵¹Y⁴⁵²Y⁴⁵³Y⁴⁵⁴Y⁴⁵⁵Y⁴⁵⁶Y⁴⁵⁷Y⁴⁵⁸Y⁴⁵⁹Y⁴⁶⁰Y⁴⁶¹Y⁴⁶²Y⁴⁶³Y⁴⁶⁴Y⁴⁶⁵Y⁴⁶⁶Y⁴⁶⁷Y⁴⁶⁸Y⁴⁶⁹Y⁴⁷⁰Y⁴⁷¹Y⁴⁷²Y⁴⁷³Y⁴⁷⁴Y⁴⁷⁵Y⁴⁷⁶Y⁴⁷⁷Y⁴⁷⁸Y⁴⁷⁹Y⁴⁸⁰Y⁴⁸¹Y⁴⁸²Y⁴⁸³Y⁴⁸⁴Y⁴⁸⁵Y⁴⁸⁶Y⁴⁸⁷Y⁴⁸⁸Y⁴⁸⁹Y⁴⁹⁰Y⁴⁹¹Y⁴⁹²Y⁴⁹³Y⁴⁹⁴Y⁴⁹⁵Y⁴⁹⁶Y⁴⁹⁷Y⁴⁹⁸Y⁴⁹⁹Y⁵⁰⁰Y⁵⁰¹Y⁵⁰²Y⁵⁰³Y⁵⁰⁴Y⁵⁰⁵Y⁵⁰⁶Y⁵⁰⁷Y⁵⁰⁸Y⁵⁰⁹Y⁵¹⁰Y⁵¹¹Y⁵¹²Y⁵¹³Y⁵¹⁴Y⁵¹⁵Y⁵¹⁶Y⁵¹⁷Y⁵¹⁸Y⁵¹⁹Y⁵²⁰Y⁵²¹Y⁵²²Y⁵²³Y⁵²⁴Y⁵²⁵Y⁵²⁶Y⁵²⁷Y⁵²⁸Y⁵²⁹Y⁵³⁰Y⁵³¹Y⁵³²Y⁵³³Y⁵³⁴Y⁵³⁵Y⁵³⁶Y⁵³⁷Y⁵³⁸Y⁵³⁹Y⁵⁴⁰Y⁵⁴¹Y⁵⁴²Y⁵⁴³Y⁵⁴⁴Y⁵⁴⁵Y⁵⁴⁶Y⁵⁴⁷Y⁵⁴⁸Y⁵⁴⁹Y⁵⁵⁰Y⁵⁵¹Y⁵⁵²Y⁵⁵³Y⁵⁵⁴Y⁵⁵⁵Y⁵⁵⁶Y⁵⁵⁷Y⁵⁵⁸Y⁵⁵⁹Y⁵⁶⁰Y⁵⁶¹Y⁵⁶²Y⁵⁶³Y⁵⁶⁴Y⁵⁶⁵Y⁵⁶⁶Y⁵⁶⁷Y⁵⁶⁸Y⁵⁶⁹Y⁵⁷⁰Y⁵⁷¹Y⁵⁷²Y⁵⁷³Y⁵⁷⁴Y⁵⁷⁵Y⁵⁷⁶Y⁵⁷⁷Y⁵⁷⁸Y⁵⁷⁹Y⁵⁸⁰Y⁵⁸¹Y⁵⁸²Y⁵⁸³Y⁵⁸⁴Y⁵⁸⁵Y⁵⁸⁶Y⁵⁸⁷Y⁵⁸⁸Y⁵⁸⁹Y⁵⁹⁰Y⁵⁹¹Y⁵⁹²Y⁵⁹³Y⁵⁹⁴Y⁵⁹⁵Y⁵⁹⁶Y⁵⁹⁷Y⁵⁹⁸Y⁵⁹⁹Y⁶⁰⁰Y⁶⁰¹Y⁶⁰²Y⁶⁰³Y⁶⁰⁴Y⁶⁰⁵Y⁶⁰⁶Y⁶⁰⁷Y⁶⁰⁸Y⁶⁰⁹Y⁶¹⁰Y⁶¹¹Y⁶¹²Y⁶¹³Y⁶¹⁴Y⁶¹⁵Y⁶¹⁶Y⁶¹⁷Y⁶¹⁸Y⁶¹⁹Y⁶²⁰Y⁶²¹Y⁶²²Y⁶²³Y⁶²⁴Y⁶²⁵Y⁶²⁶Y⁶²⁷Y⁶²⁸Y⁶²⁹Y⁶³⁰Y⁶³¹Y⁶³²Y⁶³³Y⁶³⁴Y⁶³⁵Y⁶³⁶Y⁶³⁷Y⁶³⁸Y⁶³⁹Y⁶⁴⁰Y⁶⁴¹Y⁶⁴²Y⁶⁴³Y⁶⁴⁴Y⁶⁴⁵Y⁶⁴⁶Y⁶⁴⁷Y⁶⁴⁸Y⁶⁴⁹Y⁶⁵⁰Y⁶⁵¹Y⁶⁵²Y⁶⁵³Y⁶⁵⁴Y⁶⁵⁵Y⁶⁵⁶Y⁶⁵⁷Y⁶⁵⁸Y⁶⁵⁹Y⁶⁶⁰Y⁶⁶¹Y⁶⁶²Y⁶⁶³Y⁶⁶⁴Y⁶⁶⁵Y⁶⁶⁶Y⁶⁶⁷Y⁶⁶⁸Y⁶⁶⁹Y⁶⁷⁰Y⁶⁷¹Y⁶⁷²Y⁶⁷³Y⁶⁷⁴Y⁶⁷⁵Y⁶⁷⁶Y⁶⁷⁷Y⁶⁷⁸Y⁶⁷⁹Y⁶⁸⁰Y⁶⁸¹Y⁶⁸²Y⁶⁸³Y⁶⁸⁴Y⁶⁸⁵Y⁶⁸⁶Y⁶⁸⁷Y⁶⁸⁸Y⁶⁸⁹Y⁶⁹⁰Y⁶⁹¹Y⁶⁹²Y⁶⁹³Y⁶⁹⁴Y⁶⁹⁵Y⁶⁹⁶Y⁶⁹⁷Y⁶⁹⁸Y⁶⁹⁹Y⁷⁰⁰Y⁷⁰¹Y⁷⁰²Y⁷⁰³Y⁷⁰⁴Y⁷⁰⁵Y⁷⁰⁶Y⁷⁰⁷Y⁷⁰⁸Y⁷⁰⁹Y⁷¹⁰Y⁷¹¹Y⁷¹²Y⁷¹³Y⁷¹⁴Y⁷¹⁵Y⁷¹⁶Y⁷¹⁷Y⁷¹⁸Y⁷¹⁹Y⁷²⁰Y⁷²¹Y⁷²²Y⁷²³Y⁷²⁴Y⁷²⁵Y⁷²⁶Y⁷²⁷Y⁷²⁸Y⁷²⁹Y⁷³⁰Y⁷³¹Y⁷³²Y⁷³³Y⁷³⁴Y⁷³⁵Y⁷³⁶Y⁷³⁷Y⁷³⁸Y⁷³⁹Y⁷⁴⁰Y⁷⁴¹Y⁷⁴²Y⁷⁴³Y⁷⁴⁴Y⁷⁴⁵Y⁷⁴⁶Y⁷⁴⁷Y⁷⁴⁸Y⁷⁴⁹Y⁷⁵⁰Y⁷⁵¹Y⁷⁵²Y⁷⁵³Y⁷⁵⁴Y⁷⁵⁵Y⁷⁵⁶Y⁷⁵⁷Y⁷⁵⁸Y⁷⁵⁹Y⁷⁶⁰Y⁷⁶¹Y⁷⁶²Y⁷⁶³Y⁷⁶⁴Y⁷⁶⁵Y⁷⁶⁶Y⁷⁶⁷Y⁷⁶⁸Y⁷⁶⁹Y⁷⁷⁰Y⁷⁷¹Y⁷⁷²Y⁷⁷³Y⁷⁷⁴Y⁷⁷⁵Y⁷⁷⁶Y⁷⁷⁷Y⁷⁷⁸Y⁷⁷⁹Y⁷⁸⁰Y⁷⁸¹Y⁷⁸²Y⁷⁸³Y⁷⁸⁴Y⁷⁸⁵Y⁷⁸⁶Y⁷⁸⁷Y⁷⁸⁸Y⁷⁸⁹Y⁷⁹⁰Y⁷⁹¹Y⁷⁹²Y⁷⁹³Y⁷⁹⁴Y⁷⁹⁵Y⁷⁹⁶Y⁷⁹⁷Y⁷⁹⁸Y⁷⁹⁹Y⁸⁰⁰Y⁸⁰¹Y⁸⁰²Y⁸⁰³Y⁸⁰⁴Y⁸⁰⁵Y⁸⁰⁶Y⁸⁰⁷Y⁸⁰⁸Y⁸⁰⁹Y⁸¹⁰Y⁸¹¹Y⁸¹²Y⁸¹³Y⁸¹⁴Y⁸¹⁵Y⁸¹⁶Y⁸¹⁷Y⁸¹⁸Y⁸¹⁹Y⁸²⁰Y⁸²¹Y⁸²²Y⁸²³Y⁸²⁴Y⁸²⁵Y⁸²⁶Y⁸²⁷Y⁸²⁸Y⁸²⁹Y⁸³⁰Y⁸³¹Y⁸³²Y⁸³³Y⁸³⁴Y⁸³⁵Y⁸³⁶Y⁸³⁷Y⁸³⁸Y⁸³⁹Y⁸⁴⁰Y⁸⁴¹Y⁸⁴²Y⁸⁴³Y⁸⁴⁴Y⁸⁴⁵Y⁸⁴⁶Y⁸⁴⁷Y⁸⁴⁸Y⁸⁴⁹Y⁸⁵⁰Y⁸⁵¹Y⁸⁵²Y⁸⁵³Y⁸⁵⁴Y⁸⁵⁵Y⁸⁵⁶Y⁸⁵⁷Y⁸⁵⁸Y⁸⁵⁹Y⁸⁶⁰Y⁸⁶¹Y⁸⁶²Y⁸⁶³Y⁸⁶⁴Y⁸⁶⁵Y⁸⁶⁶Y⁸⁶⁷Y⁸⁶⁸Y⁸⁶⁹Y⁸⁷⁰Y⁸⁷

AQYKIDKLQEYTDAIKLKTMAISKKYKRTYLLIEELFRLKIICKDNENELEICLIPKNTETNMWHRLQLDKKEITPE
ACDYLWSQIESFVDKETFDNYF

>SPC24_Plasmmodium_falciparum3D7 [1115]

MNPLYTKSKFKTTEFANNENKGVRINKYKTNVNNTYSKNSETSQSTQLTHKNGIMKNSNIKNKGTINYSFK
YPPKSNLKNMKSNLKGDTKNISNTYPNKIAFPHLNKGSGYNEYKNEEKNDLMNTRRSIMVDNLRKPLAS
SSNRLISQMKKDQLFNGKNGEDPNVNDDEPLTNKYKNKTHNNNSNNNKISSNNKLHNNNSNSSNMKKS
NIYYKPQDNNKNKFPNKSYNVSKIFYSNQYDKDCAMSSEQFGTYSSPVKRKYDAMDNSRNMDKDCKNE
MSYGKKKFVGRRNYYLDKMNDEEYSNCDFGDMNMSYNVKDTTQRENNDNRNDNRNDNRNDRND
RNDNRNDNRNDNRNDNRNDNRNDNWNDKCNDNKISNNINNLKVNVNKDGDLFHSQNINKKEIN
NNLPFNKRDDEDIILNRSPRVSMSVRNDTFNSNSLGRNYTYERLKHSNNNNMNNVNIGYNNVFGFNN
KEESALPTDTYNDKNDMVNNNMSVKKKMMMTNPYLEKGKLLNGNMNIYKNEKITKSYKPTPENNENNN
NNNNNNNNNNNISSSSSKVLYSHHSSMVRNHNNVPHYISDTNSEKIVMNHIDNINNKDVLNTYGDYLS
EENNNKNRRSSIFNVNTNINRNNLKKKDITNVSNVSNSMSILSNSRSKKFSCLVKNDLESEKGRGRSSVLLS
SEKKGGYGINNMMINSEGMDFKNVSNAINFYSTNDMDMGYNNDNNNDNNNDNNNDKNNNGEDENN
NGEDKHNIDEDKHNGEDKDLNKSRYNIDKNQYSNDNYYDESYFNNYVESDFQRRQMMEKGGTYFFNNIY
NNDMNNMTHKNEHLHEHLHEHLHKSILKKYNNIDNYKKNDSEYLYFQNMIRNITKYNNEENDMNNLYENV
MNLNDYCLNFTSENLEYINEILNLSEVNNNIQDIQQILYNKKVVQDKRHMVNKEYELTCQKLKECESIFCDS
NDEHGEKKLVFYEIDEKKNLLKNTYEVKMLLNKVQYKIDKLQEYTNALKLKTVMMSKRYKKTFLIEELFR
LKIKDNEQLEVCLIPKNTETDMWHRFQLDKSEITSACDYLWNQIESFVDKDTFNNYF

>SPC24_Plasmodium_gallinaceum [818]

MNQVSVKPKFKAEFVNNEKGVRFMNKTKNQNQNTISKNTVQNL禄PKNTYAKNSNLKLRTNNNTFNHP
PKLTFKNSQNAPHKNFQAPSTKVHNELRYEEKTLFNTRNTHDDSKKNFFNKSTMQNKNEEKYNDRKNV
KKSFDNNSLRKNDNNTRLTCAFNTNSTNNISNNPAYNIVYNIQNKEIENNKNNDVNNSENFIPTYSSPTKR
KYETFENINKNDEKDNEKMTQNKRFQAQKNYNIDNNFNICCNKFNNNNILNNTNENMVSYSKSNDNLFN
RNNLINKCSDKNIFYNKNKNETLNSSPYIVNVIKYDNNFSNSLGRNYTNYERLINNNKLINNHNYELNHKHY
FKEKNAKFNLLENNKNKNNTFMEQFNNHKNENHNNIYLDPIAENDNKNLNYLTETNSEKIMNNTSNENN
NDNFNSCVFDRLEENFNSYMKRKIDKSDDITIKDIEKFDTLNNSSNKNISNNSKFSNLLKNNLETNKIKCIL
NNEKIMLRNNIIETNNADLKNLSNSQAFSEASISNNMENYKEKLTHLYNGMNRMSKSLNQNELRDTTNKKI
NIDTYKSNDCEYIYFQNIINNIDKCNNVEKIDTLCENVMNINDYSLNFNYENECDINDILNLSNNVSDNIQNIQQ
IINYKKVVQDKRNIVNEEYENMCQKLKDSESIFCDTNEETGEKKLKVFYEIDEKKNILKNKYEMKKMLLKKA
QYKIDKLQEYTEAIKLKTLAISKRYKKTFLIEELFRLKIIKDKENELEVCLIPKNTETDMWHRLQLDKNEITSDS
CDYLWSQIESFVDKETFNNYF

>SPC24_Plasmmodium_knowlesi [896]

MNQVSAKPKLKTTFVGGESRGVRFMNRAGHLSAVTKNVANVSSAANPQFTHKTTFAKNSHLKLRTNNP
VFKQPPKSSLKSQRSSLKVSHVEFAKYSNKKAHHNGDDNGDMGDGYDNDNAGSNDCLNDDARDEMNDKMHNMQMD
EMHHQMNDDEMHHQMNDDEMHDGVHDDGGREAKPGGTKVPVSNARNATNAEGRKSYLNTPFLRKYRIDKYGESGKT
TIGSGKNNYNRGNPKLGQKKVLSQPQHYEREEDGAGENEWGRKAEDAIYSYS
SPMKRKYDTMEVKHRGAATRRGKGSDSDNSGGDIREASHMESEKARRDTNIQNCKRYASESLPHDSGNL
HYDNNLNPVNISNGTNEKINAYGSMMLYSNEKLLGSNNTGLGAHPSLNSSGFHGINQENGVLRSYQPLST
TGGNTAGNATDVNEGEGNVKTLLGRSYAKFDRLRKGEAMLAAPAQKYQQQTQQQQTQQQQTQQQQG
QHLNNPLQYLSKDPRRTHRRSRHTNYITDTNSEKMMNNSSSHNELFNSCTLDRLSEEKNSYANKRRD
GGMDTQVGDGMDGGDLGINRISSVRNTASNGSQKFSRLLRNDFPMGSHQLGVLRSDKGVLKNRHVES
STVEFRNMSTSQQYSEDLPLGVKGGSVLSMRGLDKQEGGDSEDYYRNVVENITNCSEENDMNILYKN
IVNLNDSVLKMDKENECLDEIICKTNGISETIRTIQHVVLTRKKVIQDKRNLINEEYELISQKLKECESIFCDSNE
GGNGEKKLVFYEIDEKNTLKGKYEMKKMLKKAEYKIEKLQEYTDAVKLKTAISERYKKTFLIEELFRLKI
IKDTENEVQVCLIPKNTKTNMWHLQLDKREMTSDACDYLWSQIESFVDKEALNSYL

>SPC24 Plasmodium malariae [1064]

MNQVSAKSKFKTNEFINENKNVRFMNKAKSNLNGVSKTINGQTTQPSHKQIYIKNSNLKQRTYNNILKPP
FKSSLKSNQNQNGSHKIPPVSFPAHNELGNGGKKVSLYDTRNLTNNNFKPHFNKLIAQNHSDNRFDEQKN
VRRNLNEDIHLEKDATGNAVSSNVNTIGNKYSILYNSPYNCEGGENTKNVSGNNVEGIMYSYTSPVKRKY
NAIEGVNKNYDNENKDLKRSDKKFDQSINYDFDNSKSYHSMNNVSSNTKENSNVHNSKLNFEKPTNGSG
DDDSNGDANGSSNADDNYNYKYNNGNINTDGNMNSSGNLISSGKIISSGKIISSGKIISSGKIIINGGN
IIGSGNIIGSGKIIIGSGKIIIGSGNIINSGSGKHSGDGKESAKGPSANIFFGSNKESTILNRSLPTMYATSGVNV
NTTFNNNMNKNFNFASTLGRNNTNYERLKNSDNCLNGTNYEGNYKYHYMEKEICTNMEELKKKEKKKKD
VNLVRNKLGSTNANTNFNESSNYINRNLNSSLYYLNLGTRNEQNDERHLQYITDTNSEKILNNNSNKNDM
FNTSTFEVLSEENFNAHVKRKTYKKGINDSHHVEGGGGVTTKHSSSNRKKNISNNSLFTNLLKEDAVNQQ
VNSVFLSNEKGVLKNSLIESNNQDLKMLSNSYHDSEVNDIGSTGGRSISKSATKSGSRNGSRNGSRNGSR
NGSRNGSRNGSRNGSRNGSRNGSRNGSRNGSGNGSRSGSRSGSRSGSRNDDRRGNKPNGHKK
QGERTMEGIYYSKYDEINSISRGVQEMDLKKTRGMRIHNMDAYKNRDNNFLHFQNIVHNITECSNSEKD
DILYKNVMILNNYSLNFNSENDCIGEIIIRNSVNSNIQNIQQAIYNNKKNVQDKRNIVNEEYETISQKLRECES

>SPC24_Plasmodium_ovale_curtisi [901]
MNMNHSSAKSKFKSTDLVSNEKNVRFMNKSNNLNGVSRNTTNIQNYHIGQKNTFIKNPNSRWPKNLNF
KHPPKANLKSQNQNVNNIPQVGLTKSSHNELRNGGGKKAFPPWNTKNLSHGDTKKQYFNKLSIQNNNEDK
YKEQKNUVYKLNDDTNLKNDNPQGSNSNNPTRLHNSSYSSVNDQLENSKNDYGNNAEDMIYTHSSPI
KRKYEVLDNINKNDERENKDIKTYNKKFAQRNYNFENNLEYNKTFCSMNNISDSINENSSNNMLHSN
EKLLSNNNISHKGNSNTLFGSNHDDVVFNRLHQARGTVHATNTAGGKSGTIVSAGDLFNSSSLGRNYSNY
NRLQSSDNVLSNSNYESGYKYFYGERGAKTNLEVQTNVTANEIAGKIPIAIASGSKNMVDNENRYFSENVH
NNYRSENHNNASSYFSNCVMRNKGHRKTSYITDTNSEKMINSSNNKDMLNSSIIDRLSEENFNSYMKRKTE
NKCSIGNSNNLEEGSVFPNNGNRNAVAKSSKNCNSLMTHDEENYQGNEALESSKKMDLKNSLIIESNNMD
LKNMSTSLPHSEGKKMPSNNGEQVHMKNHRDMHDTNIDNSYNARGEDEAKEQEEEEVVREKESSYFDEY
GEVEILPKNITHNDHWKNVVLKRHNMDVFKNNDNDLYFKNIIHNITKCSSGENDINMLCKNVMNLNDYSLN
FDTETEYINEIANISNSVNNNIRTIQQAIYNNKKIVQNKRVVNEEYEIIQSQKLKDCEIFCDTDEETGGKKLV
FYEIDEKKNILRKKYELKKMQLKKAQYKIDKLQEYTNVIKMKTMAIAEKYKKCFSMIENFFRLKIICKNDNELE
VCLIPKNTETNMWHRLQLDKNEITSEASDYLWSQIESFVDKETFDNYF
>SPC24_Plasmodium_vivax [923]
MNQVSAKSKFKTTEFLGGESRGVRFMNRAKSHLNAVSKNAPVSSAANPQFTHKTSFLKNSHLKVRTNNP
VLKQPPKSSLKSNQRSSLRVSHVEFARHSHKKGHNGGDGGDSDGNGDDNGDGYDDDAGVKDGL
NDDVHGDVHGDDGRETIQGGRKVPPPAGSATNPEERKSYMNKFGARNHGRDKYGESGRATTIGGGNN
QSSGNPKLGQSRVLSQPHHYEREAGAGDNEWGLSAQGAIYSYGSPMKRKYDAMENVHRGDAARGGK
GSDNDGDGDENGGGDRDSNGGGDPHSDRERDRERDRARRDTNAQSKRYGPRGHPEGGNSHYGN
SYSGVNSSGTNEKVSAYGSMMMLSNEKLLGSNNTGLGAHPSLNSSGLHGMNQEDALLRSYQPLSTSG
GNAGGNNAPGAANMPGGDQGEDNARATPLGRSYAKFDRLRRGQPLLARPPPQQQLKSQLSQPSQQSV
SAHRLNSSLYHPSKNPARPDRRSRHTSYINDTNEQMANHATGSHNELFNSTFDRLSEENFNFCGGKRT
EGGGERRGDERRVESYGEGSACNGAERAASANRSAPPHVQVTPNVQATPNPRVASNGSHKFSRLLRND
GMTNHQQESNQVQLRNVSTPMKYSEGRPFAVGFSSTMGPVGSLKGAPPNEARNGRGESELPAAATRLD
SAEGDSEDDYYQRVVRNISNCSEEKDMALLENVENLNNYVLNMERENEHLNEILQTNNSETVRCIQQ
VVLTRKKVIQDRRNVINEEYELISQKLKECESFFCDSNDGGKGEKKLVFYEIDEKKNTLKGKYEMKKMLK
KAEYKIEKLQEYTDALKMTLAISERYKKTFLIEELFRLKIICKDTENELEVCLIPKNTTNMWHKLQLDKREM
TSDACDYLWSQIESFVDRDALNGYL
>SPC24_Plasmodium_yoelii [822]
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NVMKNTKNNIKNNRNSKFYNSENNEVNNNKDDNINNPESILYTSTPIKRKYEAVNNTNQNTENDNNKMN
TLNKKYAQRNYYFDNNLSTYYNNKINNENSNSNNINENVNNYNNFYSNEKLFNNNNTLDVSYITCKTGSNID
TANEVNINNNNNSIKSHSLARNYDKQKNSDNLNNNSNCEPNYKCYLEQNGQHNLDVHINKLYTNDNPK
NGNCNSLPYYLNANILRSKEIRESKYLADTHSEKMMNSTNNNDVFNACTIDRLSEENFNSYVKRKSHNKNT
FNGINNLNDITTNHMTNNNNNNNNRNESSNNSKKFHNLNMNNEDSQLKNIINNQDASLKNSIIGSNIVDFK
NISMAPISEGNLLSHKSENGSVKRRSSSHNNMNANSVHFQDDSSYFNKYRDMNNMQQYIKKNDNRNK
KIMNRYNIDTYINNDNDLYFRNIINNITKYNSGESDIDDLCKTVMNINDYSLNFDGENEGINEIINITNNVNNNI
INIQQAIYNNKKIVQDKRNVINEEYEIIQSQKLKDCEIFCDTDETGEKLLKVFYEIDEKKNILKNKYQVKKMLL
KKAQYKIDKLQEYTDALKMTAISKKYKRTYLLIEELFRLKIICKDNENELEICLIPKNTTNMWHRLQLDKKEIT
PEACDYLWSQIESFVDKETFDNYF
>SPC24_Theileria_annulata [272]
MALTHHGVLDTIKEKLDKLSKLDHDFLDPDQNSLLSTFNDIFNLTSEFIKNTSKEDSLISEFVNYRNDVFNQ
YNDKLQNSIQLTNGILFYQKLEQKKEFEEQANKNAKESLKSQLNFSDFGEFRILMSRQEIMDKLNEWKHICD
LEQIEIKKLKCQIDILEDKISTNKFNSNQISYYNKKLYELLTSALKISILNVDEKSNVNLAILSESKKENEQTWHR
FSSSKDCDVATCDFLWNLIEKSLCNNGVRSSFKDPNTPKLNSIHLVSRNYN
>SPC24_Theileria_equi [261]
MNSDKNNEAIRKILGKLDRLSELEVDFLNYNANQSIISPFDDLIETCSSYIEDTEIENGLEEFLTQKKRHAEL
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ALKCKLDELEGKLSFEKTTSPDSEIYKRLFYSFITSALKTILSIDDQENVNLAILSESEKNEHVWHRLSYSLR
TADVSTCDFLWNIIQKSLSSQKLQSQIPCTNTPKIDSYHLSSP
>SPC24_Theileria_orientalis [262]
MNSAPNDLLGTIKHNLDVLSNVDVDFLNSTQNSNFLSSLYNTIDAISQFISDTSHEDVLLDEFVNYRKELSDV
FNKKVEYAKSQCKELERKLEVETRINNDAINRLKSHQTDYSPGDIEALISRQEIIDRLNQWKHSCELESIDIKN
LKCRIIDILQEKMDSVNHNSTQHKERNRLLYEFLSSALKTILNIDENGANLAILSESAGSEQTWHRFSSSK
DSDVSTCDLLWSLIEKSLSSKSNNPLPLENTPKVNSIQLSSK
>SPC24_Theileria_parva [262]

MTLTHYGVLDTIKEKLDRLSKFDHDFLDQNSLLSTFNDIFNLTSFIKNTSKEDSLVNEFVNYSRSDVLNQYNNKLQNSINLTNELEKQKDHEEQANRSAKERLKSQQLNFSGMSRQEIVDKLNEWKHICDLEQIEIKKLKCQIDILEDKISTNKFNSNQISYYNKLYYELLTSALKISILNVDEKSNVNLAILSESKKENEQTWHRFSSKDCDVSTCDFLWNLIEKSLCNNGISSSFKDPNTPKLNSIHLISRNCNNF
>SPC24_Babesia_bigemina [293]
MPDVAVLLEALSTLRDVEGDDGNSELLSRVSEAAEALAAYVADDSAEAALFSDFENLHESLRQKRDAELRDANASVDKTKQELSEAEATHTRLTEELRSLKAKTEAAIGECVRPFGVIGAGTDGNALAAIHEAQQMDELQLQLRTVEERLALDIRGLEHKLQTLDEVEKQHLAIRDDIVKRDKLLYEFVTSSLNITVVDRNESQVQLALLSEEDRTTQTWDCMYANLDECQPETADRWEAIEKTFDPDSVKSSGDTVFDMLDIGGYDDFTYLWRAVISGSPRRNMTAL
>SPC24_Babesia_bovis [255]
MPDVEAQLEALSELRFDIKSLLDNFTLKSRLSTSVDALCAFIRDDIQEWSLVNEFERVHQFLCQVFDELRAANASVDGLKKELEEAEAEHRKLTAELDEIESSYNNVDESSMLQAQQASSLIEEAFAFSYAYEESILDEESSRHQEKLSSDIATIEKASDADAERDRLLYEFLTSSLNITVVSANDKEVQFALLNEPEDRSTQTWDLVTVKTSEMDLPATSKYLWSLIERTFQHSGEESITNPVLDTVRLSL
>SPC24_Babesia_microti [225]
MDGFEFDVLDEISHSFEQLALEKIDSATFDDNVTKLKELTSKATTVLSDIKNEVSLINASKEYNQHVLSKIDVVERENERKEYNELLVLKDERQRYNELKSRLIELENKYKSSCPDHTLMTKVLREQMQWQTALESTLLETSKLKHEYKLAQKQKIFELQQENIIKIEHSRMFYEFLKAALKIAVLEVTPSRSTLVIAPLEKEDKWMLIDTDSASRDSLWRLLAER
>SPC24_Cytauxzoon_felis [233]
MALTNVKLKLKNLSQLDLDLVNGRNDLLSIFDDVVRETVDYINDKSIEDGLVNEFINTHEKLKKIFSKKINNISESHNNLNRQLESEKNNYENTLDEFKKWKSNFNSNDENVLKMLKGQDDLDKLNLRSSFISEECQVEELKVKLNLLNDELALNELNNNNLKRRYRLYYNFLSSALRIIVIDEENDLNVEIALVTNYVGDSWNRIKIPLSDTNINSCDNLWDIIQQSFKI
>SPC24_Toxoplasma_gondii_ME49 (TGME49_212800)[671]
MVLVRSPRWGPapeAAGSSPLREPRVSPCERNDAVSPFPLSSKSRLRLTERQLPPSAECPFSFSSSPLPRLPSLNADVGIPHPRLSSLDAREGEEGRQADESRGRAAEEGEEEKRAASAGDSRGQAGNLSRSAIRSSRPKMGTSVGSSTTLIPLPSRASSTLRHASSPSRPVLARLSSSSPVGSGRGAQLAAPATAAREDAGHSDLSPSSVAAFSPSKRLRRTEDQTHSFLSQPSEPALPALHSLTGSEETERAGFKSMSSTASRSSPLKNTNAIFLMASDSRQEAGAPAASAEVSARGDMHGVSPSERISVAEERISPLSSRCSSRAPASANEVPQSAALCADKHSRNEKLRELENLQHVLPAS GSLREYVQQAGEEARWRERGDSGPFSLPKTAEEFRDQWRPMVDA TVKKLCDMTDEEKQVTPYIMLEGRMLRAVRRASASLANRRTKAVAFNSEVRDLFSREQRLYDESEREFEVQRAQDQAGGQEKEKRLKDTQRKKEQVEVYRQQVHKMANLSFTRQKKLRLFEALESAKKESSSKEKSLQALYQLLQCTTRFKVNKVDGLV TGALVPEQKSALFPQLRKVGKNAQQIHKGEGDEEDMLEDTTAEPAILTLDCEKENQGGMDADVLWSLIEDTLGVGGVPVPQLLSAVRGE
>SPC24_Hammondia_hammondi [670]
MVLVRSPRRGPapeAAGSSPLREPRVSPCERNDAVSPFPLSSKSRLRLTERQLPPSAECPFSFSSSPLPRLPSLNADVGIPHPRLSSLDAREGEEERQADERRGRAAEERE EEEGEEPVAESAGDSRGQAGNLSRSA SVICSARPKGPGPSVGSSTTLIPLPSRASSTLHHASSPSHSLAGLSSSSPVGSGRGAQLA VPATAAREDAGHTDVS PSSVAAFSPSKRIRRTEEHIIHSFLSQPSEPALPALHSLTGSEETERAGFEATSSMGSRSSPLKNNAMFLMGSDSRQEASAASAEAFARGDAHGASSNERISAAEERVSPPSSRCSRHAPASADEVPQSAALSADKHSRNEKLRELENLQHVLPPTGSLREYVQQASEEARWRERGDSGFSLPKTAEEFRDQWRPMVDA TVKKLCDMTDEEKQVTRYIMLEGRMLRAVRRASASLANRRTKAVAFNSEIRD LFSREQRLYDESEREFEVQRAQDQAGGQEKEKRLKDTQRKKEQVEVYRQQVHKMANLSFTRQKKLRLFEALESAKKESSSKEKSLQALYQLLQCTTRFKVNKVDGLV TGALVPEQKSALFPQLRKVGKNAQQIHKGEGDEEDVLEDTTAEPAILTLDCEKENQGGMDADVLWSLIEDTLGVGGVPVPQLLSAVRGE
>SPC24_Neospora_caninum [681]
MVLVRSPRRGSIPGQASSSSPLRLPRLPSSEQNEAASPLPASSETSGRLAADIEPPSADCPSFSFSSSPLP RPSLNAVGSGIPPLPHLSSLDRRREGRRTDERRGRADEERQEGEGT LAERV D STGAPRFLASRASSSPSSRIPILGANTSDIAPTLIPLPSRSSVCRNASSPSPPVRAGLASSSPVGGGGREAQLAVPSTASPKSLADQAASAARDHEGHSDLSPSSLA FSPSKRPRRGEEHAPSFLSQPSETSHSTWNSSLAGSQSLPTYTNAPRG GFSPTGSRSSPLKIHTVLPNSREPTSTFAAHFAGGDSARPADFSNQRISAMDAPKNSPLSSRYARRAPDGANEVSSCATSFSDGKRAERIRELENLQHVLPAS GALREYVQQAGEEAR GKERGDRGPFSLPKSADEFRDQWKPLVDATVKKLCMTDEEKQVTRYIMLEGRMLRAVRRASASLANRRTKAVAFNSETRDLFLREQQLYEE SEREFEVVQRVAQDQAGGQEMLEKRLKDTQRKKQQVEVYRQQVHKKMATLSFTRQKKLRLFEALEAAKK ESSSKEKSLQALYQLLQCTTRFKVNKVDGLV TGALVPEQKSALFPQLRKVTGENARQIKNRQGEDV LQDA TAEPAILTLDCEKENRGGLADVLWDLIEDALGVGGVPVTQLLKS VRVQ
>SPC24_Sarcocystis_neurona [1038]

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SHAAADAAAAGPAIRGEEGGSARRGPSLRHFSRAPATTGETCTNSKNESLQPQPQHATAAVSNTSVPKR
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CDIREEEKQVTQFIMLEGRALRVLRHSKALQKKRTEVAAVHATAAQQEAEQQQAHESEKHFDIVHRQAE
DAAGGQAHLLERRLKDQTQKKERLLQLRRQLYEEMFACALLRQKKAELAKQLVSLKNTSAARETRRNALYQ
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GHSRSRQRNALQMQLPQQRGIRVEENAATESSALPVKFSAGLPMDGSMAARDRETTPASSDTDPDVFL
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>SPC24_Besnoitia_besnoiti [731]

MVLVRSPRRGPAASRVSPSSSPSRIPRPFACATHSSRASEQGSPHASTSTRAPETTGTRLEFELTSSA
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GRDASSAAPSSLSSRRSLRPEAPLFGAPESLPPDASIGRSSGSPSAAATASPSRTAAADLAATASSREGVD
SSLAASSPSKRLRKEDFVYSFLSGLRGREAETSSPRSAPRPSVGDDQLGTAVSESSGSRRDFEPPSRDPL
ASPPPPVSAAGLPQGALPPREAPRSVADADALATEAAPPSSLRLSLRAPREGAESEAASASLRRRNVD
GREAEKRRVEELEGLAQMLPASEALRDYMQQAAEGARGSRDRDDARPRFAVPKSADEFRAQWRPLVDA
TVKKLCDMTDEEKQAPFPCGVSSGVPDEEPARRQGRASRLFRWAKAFRSAPLFSRIRSPLPVFSSSQVT
RYIMLEGRMLRAAHRAAASLASRTKAVAFNSETREVLEEEQLYEASEREFEVVQRVAQDQAGGQEKE
KRLKDTQRKREQAEVYRHQVHKKMAALSFTRQKKLRLFEALEAAKKESFSKERSLQAVYQLLQFTTRFKV
NKVEGGLVTGALVPEQQSALFPQLAKTTKNAKRAIQNGLRAAGEDEDLPGQEAAAEPAILTLCDFKENEGG
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>SPC24_Cystoisospora_suis [802]

MLVELTMVLVRSPRRGPVLPPRSASPGASPSFPSSDRFLFSDDFQESNISGEGQASSAQDRDRQLSSVKP
TRRSQELFSAADSSCSGSSSPRSPSRSQLGRPVSATLRRPPYSRVLFTSTSGVPERTGSRPNSTEAECER
EPEKLMVEQGREGLAAGPGEKDRKTRGGATVESRYAGDTCASPRLEPAGRSARVQESELPEVCDVYGP
APVRPSVSVSNAQMGRVSDGALAGTAALSTAGQVAGCSFSSDPSSVRTPARATRENSLPLSSPESAMP
ASAFIPSARGEKSQAPPQEQQVQRSTVAASHAFPSSSSHTTKDHGGLCFEEAACLLDRGRLGAMDAAVI
PSAPLSLLEESPSKLLRGDFQAFQISGKTQRESEASPLAQAPSGGTPEVRTIQALNNSPRFQSEGFSF
RVSPQRHLCTEAGVKEKEDAGDLFRTSGMVGVQQETNVFHASRSVGSPSRVGSSPCAQVENLARRLP
ASRALQEYVKQALGGGAEGGKKGAVPRTAQGFLEEWSSLVATVRRLCDMTEEEKQVTRYIMLEGRM
LRAVRHASSSLTRRRAAVSARHSEAEQELNQQRASYEESEREFDVVQRHAETAAGGQEKLERRLKDTQR
KREKVEAYRHRMHEKMSALSFARQKKIRLAEALAASKESNTQEKVYQGIHQQLRQVTRFVVNGVAGGVV
TGVLVPDEDLVPPRLLRKRRVGVMPLSHSEQENVDTGNAEPAMLTIDCEKENEGRMDADYLWKLIEETLGI
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>SPC24_Eimeria_necatrix [548]

MVLVRSPKRPVPSGSNEAQQQQQQQHPNKIPAAASYVRSRTASAAAATAATAAAAAARNRAGNAST
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AAAAAASAASDASEPQQQQEPLKRLSAQDELLQPYLQLQQQQQQDEEQQQQQQEEQQEEYQQQQQ
QQEEEQQQQQEEQQQQQEEQEHQQLLQQQLQQQQEESEDEVVDGVPRSAAFRQQWQQQLQQ
LQQQLGCMEEQQQLQQTVSLEQRSAAIIRSAALLQQQQQQQQQQQQRAQQQHEASESR
FVSIHTEAAAAGGTELLQRRLLQQTQEKKQQLQQQQQQKHELLFALKTEYKARQLQQQLQQLQQQQQ
QLDELHQQUELLSLATRFKITKISSSITGLTVPDQRPAAAAAAAADRAAGLNAAAAAAADRSSLEGDLA
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>SPC24_Eimeria_acervulina [383]

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AIRKAVLLLQQQRDSRQQELQQQQQQQQAAQQEHDSENRFVSIQSEAAAAGGAEMLQRRRLQQTQHV
KQQLQQQQQQKHSLLFSLSKTEYKARQLQQLQQLQQHQHQQEIDRLREEELLSLATRFRVSKISSSITGL
TVPDQRPAAAAAAAADGRTGMAAAAAAADSGLLAAEAEGASIVQVELPQHLQQQQQSGAAAAAA
AAEQSESAELLWQQLLQQSQASDRHAAFLQRIREVCLP

>SPC24_Eimeria_brunetti [560]

MVLVRSPKRSPAPPVTIDSQQQQQQQQQHPNKIPAASSLVRNRTASAAAAATAAAAAATAAAAAAPA
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>SPC24_Eimeria_falciformis [441]

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AEDDVNGVPRSAAFQQQWKHQLQQLQQHLGCMSEQQQQLQQAMLLENRCQAIRKTALLQKQQQQ
RKQLQLQQQQQQSQQQHTASESRFTVQTEAAAAAGGEELLQRRLQQTHVKKQQLHRQQQQKHALL
YTISKMEYKARQLQQQLMKLKQQQRDIETVREEETLLSLATRFRITKISSNSITGLTVPDQRPAAAAAAAAS
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WQQQLLQQHAEDRHAASFVLHIREVCAP

>SPC24_Eimeria_maxima [555]

MVLVRSPKRSPVPTGII DTNKEQQQQQQQQQQQQQH PKN KIPAASSL VRNRTASAAAAAATAAAT AAAAP
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ATTTAATAAAAATAAAAATAAAAFAAEKSV DVSLPQQETLKRLSAHDEPLQQQQQQQQQQY G
EQEENDQQQQQQQQEEEMEEYVPGR LPRSAGAFLRH WQQQQQLQQQLGCMDTEQQQLQQTISLE
KRSEEAIRKAALLLQQQR EHRQQLQ QQQQQQQEAQQQHEESES RFVSIQTEAAAAGGEELLQ RRLQQ
THIKKQQLQQQQQHQKHDLL YSLSKTEIKAQQLQQLLQQLKQQQQQQIETLREQELLLS LATTFKISKISSSI
TGLTVPDHRPAAAAAAATATAAGTGRDSSAAAAADND SNCLLEGQAECASIVQVELPPQL
QQQQQQQQQQHQQLTAAAATATAAAA EQSEASLWQQLLEQSIASDRQAALVLQRIREVCLP

>SPC24_Eimeria_praecox [484]

MVLVRSPKRPVPSGSGSSNE

MVEVRQPKKIVVQDQGQNEIQQQQQQQQPPIKRIIAVQGTVKRTAQA//VVVVVAVVVAALAKRKAQRAST
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AASDASEPQQQQEPLKRLSAQDELLQPYLQLQQQQQQEEEQQQQQQQEEQKQEEEHQEQQQQQEEEQ
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QQQLGCMDSQQQLQQTVALEQRSTAIRSAALLQQQQQRQQQLQLQQQQQQRAQQEHEASESRF
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>SPC24_Chromera_veila [207]

MLKIPQIPKETFQEEADELLAAWEENFQTVAHQHGIVEASIIQTQGAIAKVEESQESRGKASESQREIVRLEE
SERSKKAKLAFHAKEIEELRALTEKFHNQATTYAEKRETMIQEFKWTQISNWRMELLQWSLCVHWQSDF
EKDSSSGSGSKNQSGMILPKGKKPQPFSLDGDRAPLEKADYLWAKTADAHIPPEYRSLTQVEGI

>SPC24_Colpoda_aspera [195]

MNFGFGFDSQTIQDVLKDTKFEGRYVNAFLSKREEIQQLMNFRIEKLRKDIENVLQRNAQLAYALDDGTEES
IITERAEAFTEEYLRINAЕINALHKEAKDLAELSEGTYNRENLRGNDKKRLEQVNKLIGETKALSQIYINFIRV
RWVKGGKEKLFGYVAKSDKDFERFESSKSEDGFERFGSNWEKLEKSIL*

>SPC24_Cryptosporidium_andersoni [207]

MEIMSAKRSISASNESFRDINYMIQEVVNEFSDFKSEDETSLKLTAEINKLDKYISSVYDILKDSVENIRKNEE
AYKKELEDTNFENSISFNTKDSSEEKCYQLSKYCEEIELELSKIHNQHAVLLEKKKQLLKSMDSEEMFIAK
RKMYQNISSISWSSISELSLKGHFIPPNAHFTESFQLQLSENNRITNADYLWGEIKRFV

>SPC24_Cryptosporidium_hominis [93]

MSSKCEEIEKELSSVHNRYALLLEKKKNLLKSMRDSEDVLHAKKKMYQSISMITWSLISESFIQGHFIPVNCP
TKTETFQLQIHENSKFPNYK

>SPC24 Cryptosporidium meleagridis

MFESIQDRNGVGRGVRKMSISPQFNENIKE

MLIDDIKLIENSYLEELENGENICNNSGNEIYNCKEDNKRSALMSSKCEEIEKELSNSVHSRYALLLEKKKNLLK
SMRDSEDVLHAKKMYQSISMITWSLISESFIQGHFIPVNYPAKTETFQLQIHENSKFPNHK
>SPC24_Cryptosporidium_muris [207]

>SFC24_Cryptosporidium_muris [207]

MEIMSAKRSISASNESFRDINYMIQEVVNEFSDFKSEDETSLKLTAEINKLDKYISNVYDILKDSVENIRKNEE
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RKMYQNISSISWSSISELSKGHFIPPNAPHFTESFQLQLSENNRITNADYLWGEIKRFV
>SPC24_Cryptosporidium_parvum [211]
MFESIQDRNRVGRGVRKMSISPQFNENIKEIILTQEIAQIYEHLGGETDVLSQLIQNEIAELENYIKSTYDKLQ
MLIDDIKSIENSYLEELGENICNNSGNEIYNNFKEDNKRSALMSSKCEEIEKELSSVHSRYALLLEKKKNLL
KSMDSEDVLYAKKKMYQSISMITWSLISESFIQGHFIPVNYPTKTETFQLQIHENSKFPNYK
>SPC24_Monocystis_agilis [213]
VSDVDEKVVEEIEENEKKMEKDVKDRSEQIGSKKENVVREFNERARDELDDIAGQRSDELQELKRIQESLDEE
KAKEAENEELQKETEETKQQQLKIVTERTAEIGPEIQAQAIKRRHKLNEQREHLTQGIRGHAENSKNVFLKSICD
VKFTKWEQISETNIVAKGYIIDQPGSPFVIGLPFSVQCANDKSVENADWVWREVEDAEERRSNRGARGE
>SPC24_Lecudina_tuzetae [219]
RRVHGKSTPVCHYIFENLEMFDTSVVERFTEEIENRLLAHNSGPLHDIEKSCATFLEIVQTGDNTLATSLADR
SKELRTAESRLESLKKEEVKFREDVTREKQGTGRALKSSFEHYQREVEETARLKAAKAHRAQLSREQRKH
ASQFREKMKILSKTTTYQQATRVMRDMDSEVVSAAFQRGNSELWEMRYEGEDVGSGALWSKIETA
ALDDCEK
>SPC24_Pterospora_schizosoma [211]
VSTQSTWGDVVMIEPEFDVLKVKNLVESANESLSKCDYAANFDKIQNQEALLDDIIHSSDKALSKNIEKQHT
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LKKHLKCKQTLGLTRMSWLEQTDDTVTGAFCICEEPQHNPFLRFDENPVDRADYIWENISKTIHS
>SPC24_Selenidium [234]
MPRPTLAWSNSTLPEKTESAIAELRGLVEAVEQNLADFKDINQSTRALERSEKAVTQWIRDIDREFANRL
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RRAIAKNVAKLDHYSTKEIVMKATVATFMSVTQLKWTENSSTRIRGSFLPHKNVKHVKNFDLPNEDPVK
TAETLWQEIYAFSVPPELQ*
>SPC24_Aurantiochytrium_limacinum [218]
MEDLLAFARSSGGDGDEGSPWEMGSILAEVATFAGEQKELEALAQAQELSksVRDTYKDTETEASEAVS
ACEDEASAARARAFAAAQTPTSPEIEALERERMALLKRLDEASTAEETTMAMVAELEKRVQAAEDALKAA
KNSEDAAMPLMKHSLSLYASISKIKWDFAPDAQKKGIFAGSVVDAEEGYVRNFEIDERNKSQFQIANQLWD
LINAA*
>SPC24_Bigelowiella_natans [205]
MSLISDPVTALLRDTRQEFQTQGKDEEIIYRALESIQDSEVVMRHTEVNVKQLIEDLTSSVQLLEHKNGQDW
EEKKAELEERESTVSSDIAAMTKMDDLTLNRQGEEQYLQKLTEKKEKMFAENESKLAESQKNLKKNTAKL
NVYKVITHIEWKGLKDGT AHLMRKKEGLTSALTHLVRPFELKGDSEFDQVNDLWNVMWEDHC
>SPC24_Arabidopsis_thaliana_1 [212]
MRDQSRNFEMVISWGDELIHVLDLDRKGFDVLVQTLEQLRAIPFSCDEDFKEIHESLQDLQKKLDVCKEKTD
EANSEIADEEEIERLQKELDEELELECKLKEELRFIADELKDLNSQEALFEEHRLAIKRNKRDQLRTETKLP
YASVTRVIPNIDDSLKTSGYMGRDKRLMDKFEDSNKSTSVDTCNERIGEEQILTVIGIANLGT
>SPC24_Arabidopsis_thaliana_2 [201]
MGNASENFIEDLMSYGDDLINLLNVKNGFDIISQSSEQFKALNFACHEDFNQIQGSIEDCKTLYACKKKTE
EAYSDIAAEDEIERLQKELDEEMEREFLKDELRLVADELKDLNAQLSSIDEHKQSTKRKVRDDLRAEKKLS
MYASVTNVIPDIDPSKISGYMVDRKRLIEKFQFETNKMTAYETCNSIWSIINKQ
>SPC24_Giardia_intestinalis_1 [196]
MTQLEINDMVSVACAEMTRESLILQKTDLPTAFAELIDMFLIRRRIISKHDSTLKTTSNHNVIVAHLSAPTAP
DDTLLKSLEDELKSLQASLDETAEGIKTLQKEQEALADARARHRAEQANKQRRDRQLTEMRSLLVFYRSI
SNVYWDADHRCFLLLPHKAKLIKFSAPDNKFRLSQNDSGEDMCIWDLDD
>SPC24_Giardia_intestinalis_2 [184]
MVGPFDSLADALIRQMNLQEDMRHNGEYSLLLRIIRGQTSLDRVAKNQEALSVARKAGAILTVSRPKPQ
TDPGFIQRTQKALEETQDAIQETQRNIERLEAQLKDVTAKTANKDELRRAALTADRENAGREELVELYHSI
SGVHWDPQDGLGVYLSEEIAKPIRFSSCVDATNQLWEMINM
>SPC24_Trichomonas_vaginalis [177]
MFGGEVSPTRLCEKIKDKKVEDAHYNLQDENTKKTKELOMQKIQSLREEEKLKNSDKISSNSQKLEIE
KLKQAKEELIKKRQFLKNRVEELESIVQKQEETKNSTEDPAAALTLYKSIAPITISSANQNQISGMIAYGTDST
CFFSYNMDDLTQNQNDLWSKIDENRKKAQL
>SPC24_Dictyostelium_discoideum [281]
MIENKQNKQNKQNNKDNKNVNFIKMIEDLKEDKIKRYLDNDVENLKRAQNLYQDITEKMAQQDEFRNELDI
WKNESIQFEQSIPTPIDEKDHSKINFLETRKNQLIEKNNQFEEEIFIENELSSIRLKKTQLINKENQLSHENTI
TIPNKEYLYSLYTNTGIKWSNNDNN
NDSNNNNNTSSSSPPKQESPFIKGII LDQTLNQNNCQIKPFSIDKNQYSNDFEIVNKIWDLIDK
>SPC24_Saccharomyces_cerevisiae [213]

MSQKDNLNDNPVEFLKEVRESFDIQQDVTAMKRIRHDLVIKEESEARISKEHSKVSESNKKLNAERINVAK
LEGDLEYTNEESNEFGSKDELVKLLKLDLGLERNIVSLRSELDEKMKLKDSEIISTPNSKIKAKVIEPELE
EQSAVTPEANENILKLKYRSLGVILDLENDQVLINRKNDGNIDILPLDNNLSDFYTKYIWERLGK
>SPC24_Homo_sapiens [197]
MAAFRDIEEVSQGLLSLLGANRAEAQQRRLLGRHEQVVERLLETQDGAEKQLREILTMEKEVAQSLLNAKE
QVHQGGVELQQLEAGLQEAGEEDTRLKASLLYLTEREELKEIEADLERQEKEVDEDTTVTIPSAVYVAQLY
HQVSKIEWDYCEPGMVKGIIHGPSVAQPIHLDSTQLSRKFISDYLWSLVDTEW
>NUF2_Plasmodium_berghei (PBANKA_0414300)[457]
MNTMNDPVPKLPFDEIRNEMNKYGVVTPSTLKHPTTEDVQGVYSICIKYILNKDINNIRIEEFTGDLKSSMPS
IDGIQILPNEGKNHLQAIGNLRFIRHCEKINKILNMDNTLSYIFKPTSGHITKLINAFMHFMRYREQIYNENDAT
IKQIEERKNESNLDNELKSIQSELQVLLSKHEEVRTSILNEKNIKRDYEEEIIENQNSLNSQQSILISLKSTKD
RIVNETNELIFQFSRYSRQKKEDLEDQIVPSPEKLQQYNDLKDLLYEHMSHCETSKKKNEDIKNKINIADLCIK
KLVNLLTILTSHINETLKVHIDKKNKLKDLGTLNLKSLKEENENLTKKKREHENILNETENNLFQEKNKWDEKIQ
DEKKNTIIVEENVKQIRESIDGITTNTQEIKEINNIVNHIQDTVNIYNKNFAIIADLIENTKKSQKILKNKIQNNIQ
NCIKTHL
>NUF2_Plasmodium_yoelii [457]
MSTMNDTPKLPFDEIRNEMNKYGVAITPSTLKHPTTEDVQGVYSICIKYILNKDINNIRIEEFTGDLKSSMPSI
DGIQILPNEGKNHLQAIGNLRFIRHCEKINKILNMDNTLSYIFKPTSGHITKLINAFMHFMRYREQIYNENDATI
KQIEERKNESNLDNELKSIKSELQVLLSKHEEVRTSILNEKNIKRDYEEEIIENQNSLNSQQSILISLKSTKDRI
VNETNELIFQFSRYSRQKKEDLEDQIVPSPEKLQQYNDLKDLLYEHMSHCETSKKKNEDIKNKINIADLCIKK
LVNLLTILTSHINETLKVHIDKKNKLKDLGTLNLKSLKEENENLKKKKREHENENILNETENHFLQEKNKWDEKIQD
EKKNTIIVEENVKQIRESIDDITTNTQEIKEINNIVNHIQDTVNTYNTYNKNFAIIADLIENTKNSQKILTNKIQNNIQ
CIKTHL
>NUF2_Plasmodium_vivax [454]
MSGGGGEVVPRLSFEEMRGEMSKYGVETQGTLKNPTTEDMQGVYSMCIKHILNKDINNIRIEEFTGDLKSS
MPSIDGIQILPNEGKNHLQAIGNLRFIRHCEQVNKILCVENTLSYLFKPVSSHMARLISAFVHFTKYKEQIYVD
NDMKIRRIEEGKSEDSLGAELKAVKNELQLSQENYEQVKNSVLSEKNKKRDYEEEIIENQNMLNAQQSTII
SLRAAKDKIVNETNEIIFQFSRFRQKKEDLEDQIVPSPEKLQEQYQELKNLLEHVSYFEKDKKKNEEIKN
ISDLCLKKLVDLVTILTGHFNETIKLHIGKKEELKGLEKHLNLKSEKEHTMKRKQQEKILLETEQYFAQQKD
KWNAAVQAEENAVVEEKASQLVQMDELKRQADREAREIDSIVKLIQETLNNYRRNFALIDDLTARTRSS
HALLAAQVRGQAAGRIG
>NUF2_Plasmodium_ovale_curtisi [457]
MNVTNDPVPRLGLEEMKSEMNKYGVEITQGTLKNPTTEDVQGVYSICIKYILNKDINNIRIEEFTGDLKSSMP
SIDGIQILPNEGKNHLQAIGNLRFIRHCEKVNKILNMDNTLGYLFKPVSSHVKLINAFIHFMKYKENIYSENDA
KIKNIQEKKNEDTTLENELKSVENEQQLLTKHDEIKNSIMSEKNKKRNFEIIIENQNLNSQQTILISLKSTK
DKIVNETNELIFQFSRFRQKKEDLEDQIVPSPEKLQQYNEELKDLLFEHLSYLETDKKKNEEIKN
KLIKVELLTTLSHVNETIKIHIDKKNEIKNLEKNIKSLKEKDYLTNKEKEQDNILNDTEQYFLKEKKWNEKID
EEEKNKIHVQKNVKLIYEQVDNINMQTNHELQEIKNIMKHTQDTVNMYSKHFIALAELTENTRNSQRTLASKV
QNNVHACVQTRM
>NUF2_Plasmodium_malariae [457]
MNGGNDLVPRLGFEEEMKSEMNKYGVEIAQNTLKNPTTEDVQGVYSICIKYILNKDINNIRIEEFTGDLKSSMP
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KKIQEKKIEDSTLDNELKSVQNELQILLSKHEEIRNSILKEKNTKRNYEEEIIENQNLNSQQSVIISLKSTKD
VNETNELIFQFSRRLRQKKEDLEDQIVPSPEKLQQYNEELKDLLFEHLSYFEADKKKNEEIKN
KINVADLCIKKL
V DLLTTLTSHIDETIKFHINNKSELKSLEKNIKSLEAEKQNLTNKKQEQQQLLINTEQSFVQEQNWKWNQKIEEE
QKNYVLVEKNVTQIYENIDSVNICKREVQEIKNHICQDTINSYNQNFAITDLIENTKKSKTILTEKIQNNVQKF
IKTHI
>NUF2_Plasmodium_knowlesi [456]
MSGGGGEVVPRLSFEEMRSEMSKYGVDTQGTLKNPTTEDMQGVYSMCIKHILNKDINNIRIEEFTGDLKSS
MPSIDGIQILPNEGKNHLQAIGNLRFIRHCEQINRILCVENTLSYLFKPVSSHITRKLINAFIHFMKYKEEYND
FKIRKIEEGKSEDLALDTELKAVRNELQSLDNYEQIKNSVLTEKNKKRDYEEEIIENQNLNAQQSTIISLR
DKIVNETNELIFQFSRFRQKKEDLEDQIVPSPEKLQEQYHELNLLLEHVSYYESDKKKN
NEEIKN
KINVADL
CLKKLVELLTCLTGHLNDTIKHIEKKDQLKDEKDLKTLKTDKDNLTLLKKDQE
KILRDTEQYYAEQKDKWN
AKVEGEEKNVVVEKKVNQMYEKIDE
LNREADREAQEIDSIVKLIQDTLDNSRNFAIIDDLTERTRN
SHALL
AGKVRTLVLPCAGKP
>NUF2_Plasmodium_gallinaceum [457]
MNIGNEIVPKYGFEEEMKNEISKYGVETQGTLKNPTTEDVQGVYSICIKYILNKDINNIRIEEFTGDLKSSMPSI
DGIQILPNEGKNHLQAIGNLRFIRHCEKINKILNLDNILSYIFKPVSSHITKLISAFIHFMKYKEEYND
QEKKNEDNILESELKTVDS
ELNMLLGRHDEVKTSIINEKSTKRNYEEEIIENQNLNAQQSIIISLKSTKD
KIINE
TNELIFQFSRRLRQKKEDLEDQIVPSPEKLQQYNEELKDLLSEHMSYF
ESDKKKN
EEIKSK
INVADLSIKKL
V D

LTILTNTVENTIKQHIDKKNELKNLEKNLKSIEKENLENKKEEQDNILSNTEQCFFEEKNKWNKKIHEEEKK
ITQVEKVNVKTIYENIDNINFKTNREIKEINNIVKHIHTTINSYSRNCDIITDIENTRNSQKILSDKVQNNIQKYIKA
NV

>NUF2_Plasmodium_falciparum3D7 [461]

MNKMASTHNEIIPRLGFEEMRNEMNKYGVEINQSTLKNPSTEDIQGIYSLCIKYILNKDIQNIRIEEYTGDLKS
SLPTVDGLQILPNEGKNHLQAIGNLRFLRHCEKINKILNLDNILSYIFKPVGSHMTKLINAFIHFMKYRDQLYNE
NGEKIKSIQEKKNEYDVLENEYDALENELNKLLLKHEDIRNNIINEKNIKRNYEEDIINKQNLLNSQQSLIISLS
TKDKIVNETNELIFQYSRYRQKKEDLEDQIVPSPEKLQKYNEELKDHYEHIAQFEDDRKKNEDIKNKINIADI
CIKKLVDLLTALNEHIEHTIKLHIEKKNNLQTIEKQYKSLTNEKQNFITKNTEQDKIIKETKEFLQQEQTKWNQK
IKQEQQHNTILIQQKVKDIYQNVDDLNKTNREINQINNIKHIQDIINHYNKNILLITELIQNTKNSHSILTHKVLNNI
QKDISANM

>NUF2_Plasmodium_chabaudi_chabaudi [454]

MNDTVPKLPFDEIRNEMHKYGVAITPATLKHPPTTEDVQGVYSICIKYILNKDINNIRIEEFTGDLKSSMPSIDGI
QILPNEGKNHLQAIGNLRFFRHCEKINKILNMENTLSYIFKPTSGHITKLINAFMHFMRYREQIYNENDTTIKQI
EERKTESSILDNELKSIHSELQVLLSKHEEVRTSILNEKNIKRDYEEEIIENQNSLNSQQSILSLKSTKDRIVNE
TNELIFQFSRYRQKKEDLEDQIVPSPEKLQQYNDDELKDLLYEHMSHCETSKKNEDIKNKINVADLCIKKLVN
LLTTLTSHINETLKVHIDKKNRLLKDLGTNLKSLKEENDNLTKKIEQENILNETENNLFQEKNKWNEKIQDEKK
NVIIVDENVKQIYENINDTTKTNQEIQEINNIVHIQDTINTYNKNFAIIADIENTKNSQKVLTNKIQTNIQNCIK
THL

>NUF2_Babesia_microti [437]

MDELVDALKLGAAQVSERIFKNPTKDETGLYCLAIQEVFGRAITQVRYEEAMGLKSNPKSDLFETFGFTIE
SNSYKSIGDLRFYRYCQRLHSALGLEPINPTTISNPTSADIKKFGKALLAFLKFRDNVLQLLGTDLDTLIQSLD
VFKFALINDSNQAVRELSEFSRCNEKNVHVLNMQQEKKCLETKLTLINEEFTMFKERLRTVNTRINELQESIE
QILLQTVAARQTNQDLQDQVVEDPDKFKSDLMQYNSQIDSGNLELDHLKKSEEKLHEKIANFQNSKSRIDTT
INLILSHLDNVYPPYSELISITSATADSQAIDDDLKRINLEINDCHETLKCCKLDQDNMQRNMSLLENKLADNT
KRLSMMDETGRERIESVHKRIKEKEEACLELEREIKKLEILKLAKDNTNFNNCIQDFVTLLHPYEKLLRYY

>NUF2_Babesia_bovis [477]

MDLTEEPQILYELFRKVRVDEVVEDLKELGVDVREETLKNPTPEAACFYGLAIQVVFGKTRTDIRPEDVYS
HIHVYDSGVEGLNITSNDNLEFLKRGIGNLRFWRYCQKLHETLGLQKIERYEIFNPTPESFHRFISAFVVYRFF
REALKSLFEDSVARLNFCVEQDAQLDESINKVRRELQNLQRLKDENLTDNIESSVEERAILEHSLLQAKNVFN
DSKDEKAKLDGEIEHIKLSINEVQLKKTCSRHLNETLSEQVVEPDALYNQNDLEAEDSVATATLNNLEER
FEDVIGRIKELEESSKFLVDMKTKLSQHVDEVLKPLLENNSTSQQNRNETLREQISQQKCKLEAKLSLTN
TQQQYEDKMNKAKELADAIEIKQAHKFAEEIERDVQGIKNTTAEQNDELSKLSNELVNRERYMIATFASIEEY
FSKVKEAAEAYKSSMDAVTERLHSTVTKSHPYGAEFIMDESTS

>NUF2_Babesia_bigemina [463]

MDLTDETRVIDELFRKVRFDEILEDLKVLGVDVREQTLKNPTTEEALGLYGLAIQVIFGKTRTDIRPEDVYSQI
HVYSSGVEGLEITSDNLDLFLKRGVGVLRFWRYCQKLHETLGLPDIERYELFNPTPESFHRFISAFVVYRFR
EALKSLFEDAIARLNFCAEQEEQLEDNIGKVQQELQNFKRLKDENADDIVSAGDREHLEGLLQAKSVFNE
HKEEKAKLDADINRVKQTINDVQLKKTARHRNEMLCEQIVSEPDLSHAQKDELEAQEVVQNATLKNLTEE
FDEAKRRFRRLLEDCHYVEGVKAKMARHLDEVLKPLAENLSASRTLKARNETLQQQISRQKEQRDSARAE
LTAKQQLNEEKRLLQEAADNELKAARKFSEDTHNEVVEMQQAISQEKKKVESLKHDLAEREERVTAVFSA
AQASWKVVTDAADSYAQRMDEIMDNVEKVIQEE

>NUF2_Theileria_parva [459]

MSDSSSQLVLIHDLFKSVRPDEILEDLKQFGTVVSASAFRNPKPEESLGLYSLAIQVVFGKTIHDIKPEELSGKL
YDTKTEPGVDFSTENVDTYKNAIGNLRLWRSQKLHKVGLGEIERYELFNPTAKSFNELISAFVVYLRFR
QALYSLYESSIQKLDNLAELDLRLDEDLKNLVRENDKFKNMVLENVVEHESLLKLNELENRMIVANEETN
AKETKKKLEIEKEKINSVLNDTMLSJKSKARFTFDDLSQHFSTNIETASKIFNELNAEYASQSQTLEALVSDVE
RLNTIHASLVESCTFELLKSNLEEHYNEVIIPHVKSLELSKRLOSSQTESLKLKIKQLTRTRDSILKDIHQKTL
QADRLKRLSDELQAKVDEAHRFAQESSNSESGLRAKISDMRDLISQMNSRAQDVRKHSQELFSQLSNMIN
TLYDSIESYNHKINNLNNSLKF

>NUF2_Theileria_orientalis [459]

MGDSSDPSIFYELFKKVRPDEILEDLKQFGVDVNPEAFTNPKSKESSLGLYSLAIQVVFGKTS HDIRPEEIYRQ
LIQPPNELLGVDFSTENVDFKNGIGNLRLWRYCQRLHSVGLKQIQRYELFNPTPKSFYDLISSFVVYLRFR
EAIYSLYEVSVSNLDLFLAEKELRLSEDIMNLIKENQKFKGVTADTAEHQLLLSNKRELEAKLLSANEEFTRF
KSRKKELDLEMGKIKNLLNDTMLSJKSKARYTFDDLSEHVTNIETASKLYTELKAEDANQRQRQLDSLSSKAE
DLRAKYSKLLEASSVFDRLKVNLSDHYRDVVVPHVDSLEFLKKASVQISRLNDRIAELTRRRDSLLSDLERH
KKVHEEKLRKLGEESQTRIDQAKKFASESKSGNELVLRKITDVKLMIEHNEKISETRANSFDLFSKIDERLE
SLKESAFLYNEKIQLLNKRLAEA

>NUF2_Theileria_equi [458]

MDRSDDPIIIDELFRKVRPEEISDDLKELGVIDPKAFKNPTADESLGIYSVAIQVIYGKTVDDIRPEERLGLFS
DSVSIIDGVDIGPNDIFLKCGIGNLRYWRYCQRLHQTLGLEPIERYELFNPNTESFNRFITAFDVYLRFRDA
VYKLFETTIASLNLLSERDMHVTRDIRSINDEYDKFKVHNETLDEHRALESNQKETLEKKLVEAKNLFNEYKH
KKNEIEGELEDVKHALSNLLSKTKERCLFDDLSDFIIDSAAVEKILGELDNEFQIKNGKIGLLKKNVTDLRLK
YDNIVRSCTFLEELGKNLQHHHNVTIPHVDSDLQELKKKAQCDSLGEKIKILTENRNFIVNKLEHLKKLQKQ
KLGREQEESQLRINLAKRFAEDTEKSSDLIREKISETNERIEREGVVLDTLYRQNTEELHNLNRKIKHVATATK
CYNERIMYINSKLQE

>NUF2_Theileria_annulata [459]

MSDSSHVLVIHDLFKSVRPDEILEDLQKFGVSVSASAFRNPKPEESLGLYSLAIQVVFGKTIHDIKPEELSGKL
YETKTEVQGVDFSSENVDFLKNAIGNLRLWRYSQLRHVKVGLGEIERYELFNPNTAKSFNELISAFVVYLRF
QALYSLYESSIQKLDNLAEIDLRLDENLKNLVNENDKFKNSSLKEVVEQESLLKLKNELENRMIVANEEFNN
ARETKKKLEIEKEKINSVNDTMLSKSKARFTFDDLSQHFSTNIETANRLYSELNAEYISQSQRLEALVSDVE
RLNTLHGSLVETLETFELLKSNEEHYNEVIIPHKSLELSKRLSSQTESLKLKINQLTRTRDSISKNIELQKTLQ
SEKLKKLSDLQAKVDEAHRFAQESSNSESGLKVKIYDMRNLASQLNTKAQDVRKNSQELFSNLSQRIQTL
YDSIDSYNETIKLLNKSLTNS

>NUF2_Cytauxzoon_felis [401]

MKNVSIIQELFKRVRIDEILDDLKELGVHVQYDTFKQPKCEDSLGLYSLAIQIIYGKTIIDIRPEDLYNKLKVLPID
INGLNYNENDIEFLKCGIGNLRLWRYCQRLHESGLNMIRQFELFNPTSESFIKFITSFIVYLRFREALYNSFEI
KISSLDLLAERNYKLTHDVKLMNDEYNKLKSYKNNLNQNNNLIQRKKLLEIKLKEVKNKRDQALKKSIEISKR
RESINTINDLKFKIDSINKLKDNLKTKLKILKDNNSMTIKINTLKDGNKYLNIIKKLSNHYDITLSPNIKKNTNY
NHNYKRNKLILMDELNKLIDKRRLLIKDKEELQKNTIEEIDKTRDEITKINSNLKESKKNAILIFNKLNEKFTPVI
NASKEYRKYVNDITNQLNNLVI

>NUF2_Nephromyces [334]

MHRQMSRNSNFVNLYPFEGTDTRGLSTVTPQHTERFSYRHSASHYATSTPQLKNSHRLSIEEIRHELEK
LSIDSKPLLSANPPKCEEFHCLLTRIVEDVGLDKAMVRIEEKEGCSIPGFSQPQLFGGNTNRNVEDESNLY
VSAIGNLRFRLCEYVCRDIFSPTIKIDHFFNPNRRAIIQILQAYIHFLRFHDEIHTEFKEGLDSMQQLAEQIKVV
DTHVQNIQIDYEKLVNRHSQLESETKETSKEIENLTNECKLLKGQLQRYCDTNQNLQLQIKQEKEITDLEY
QISKTSNEIKTLQSQLTDSPEQLEKEQIAKEEHSKMKRKYE

>NUF2_Toxoplasma_gondii [608]

MAAPGLPQPGSLSTLAGYPSSTYSSQNSPFFPSADRGGRMTLEEIKRDFTKHKLECPDTMWREPQPEQV
QGLYSFAIESIFGLTVNDVRIEEVTGDVRSCLPSIDSLQFLSQDGRLHAKAIGNLRFIRLCQRLNRLVGLPES
RETFASPTASGVQRFASAVCMLLRLRESLCRQFESQIQQRASLQQSLQKLSQNAQLVEQELLRFRAERQT
QQPLAQRQKQQSELEELRQRHSELGALMEEFKERQAVHGRLELELGDLVLELMNLKQEREELHDQVV
HSPEKLMERRDELRVQQKHLDQALQLELENLAASQQKLLAFAKAVKKAKKAMEILSAHRDQVLAPHLGFRS
DMRTREKLFRELGEQKEQLSKAVQDLQAEREELARQLEDQERKKDEEETELRGHLARAKREVEERKKALA
DQQDMTAFLREAELKLEEKLAEQKQRHKLLVDAIEEEIQKVYAAFLTYVSQMWFIRSQMPLSLDLSQSASFL
AEHSEKTHRGSPACEREENDDFLASLLAESSPSSLGEEKENLPQSRPTAAADAERRLADFSPEKGRKRL
SCLDEETEEEARMHERERESDRLFPAAREPREDGDFPMYGHAE

>NUF2_Besnoitia_besnoiti [616]

MAAPGLASQAFYSSAAFSSQGSPLYPSDRGGRTMLEEIKREFAKYKLECPDTMWREPQTEQVQGLYSFAI
ECIFGLTVNEVRIEEVTGDVRSCLPSIDSLQFLSQDGRLHAKAIGNLRFIRLCQRLNRLVGLPDFREMFAA
PSAAAVLRFASSVCSFLRLRENLLKQFEVQLQQREALQQNLEKISEDQQRVEQELLRFRAERQTQLPVSQQ
QKQRREELEAEELRKRHGELGALMEEFAERQAVRGRLELELGDLVLETNLKQEREELHDQVVHAPEKLM
RRDELRMQQKLLDQQQLQQLERAAASQQKLLAFGKALKKAKKALEILSEHRDKVLAPHLAFRSDMRTREKF
FREISEQKKKLTQTVHELQTERARLNRELETLEKKTEEEEAEELRQRLRTKQETEERKKMMINQQEMTAALL
REAEEDLEDRLAQKQAGHRALLHAIIEEEIQRVHAAFLTYVSQQLHFLRNQMPLSLDLSQSASFLADHHAKRFS
GFSPSVDDGGREDLCARDADYLSASLLESSAPFADDASEAGDGKENRPLGGTHADAHTRGVGGEAEGADS
FAEAKELRGPGGAHSAASPAKAKRISSLDELEEAELRMEIDLREGGGEDEG

>NUF2_Neospora_caninum [609]

MASPGMSQSGGLSAHSIYPSTSFSQGSPFFPSTDGGRTMLEEIKREFAKHKLCPDTMWREPQPEQV
QGLYSFAIESIFSLTVNDVRIEEVTGDVRSCLPSIDSLQFLSQDGRLHAKAIGNLRFIRLCQRLNRLVGLPDF
TRETFAVPSAAVLRFASSVCTFLRLKETLWKQFEGQIQQRAALQQSLQKISQDAQLVDQELLRFRAERQT
QHPVAQRQKQRRDELESELRQRHSDLGALMEEQFAERQAIHGRLELELGDLVLELMNLKQEREELHDQVV
HSPEKLMERRGELRLQQKHLDQQQLQQLESLATSSQQKLLAFAKALKKAKKAMEILNSHRDKILAPHLGFRS
DMRMREKLFREIGEQKEQLSRAVQELQVEREKLSQLDEQERRKEEEEMELRGNLSRRKQETEDRKKAL
VNQQEMTTAFLREAERLEERLAEQKERHKLLVEAVEEEEIQKVYSAFLTYVSQMWFIRSQMPLSLDLSQSAS
FLVDHHAEKLQTFSPAASRDREENPDDFLASLLSNSSDDSLAPSALREEEKENQAQPRQPEGDRGRRDF
RTLDSPEKTKRLSFLEETETDVRMEDLEPEHEGRLRTSPRPAESR

>NUF2_Hammondia_hammondi [608]

MAAPGLPQSGSLSTLAGYPSSTYSSQNSPFFPSADRGGRMTLEEIKRDFTHKLECPDTMWREPQPEQV
QGLYSFAIESIFGLTVNDVRIEEVTGDVRSCLPSIDSLQFLSQDGRlhAKAIGNLRFIRLCQRLNRVLGLPEFS
RETFASPTASAVERFASAVCMLLRLRESLCRQFESQIQQRASLQQSLQKLSQNTQVVEQELLRFRAERQT
QQPLAQRQKQQNELEELRQRHSELGALMEEFKERQAVHGRLELELGLVLELMNLQEREELHDQVV
HSPEKLMMERRDELRVQQKHLDAQLQDLENLAASQQKLLLAFAKALKKAKKAMEILSTHRDQVLAPHLGFRS
DMRTREKLFRELGEQKEKLSKAVQDLQAEREELARQLEDQERRKYEEETELRSHLARAkreveerKKALA
DQQDMTAFLREAEKVEEKLAEQKQRHLLVDAIEEEIQKVYAAFLTYVSQMQRNQMPQLSDLSQSVSF
LAEHADKMQPGSPLACEREESDDFLASLLAESSPSSLGEEKENLPQPRPTAAADAERRLADFSPEKGRKR
LSCVDEEETEDARMHERERESDRFPAGRELREDGDFLMYGHAE

>NUF2_Cystoisospora_suis [690]

MSSSLYGGPRSSSASPVAVGDTVHTQMTSNSLN SHATSTSPPSFSSFASLQFPAQKPSPVPPGGTGLPPS
SSSPFSLPASHTSSGFFSNAQAASGAGAGTNASFFLPSTSQGGGGRLSLDEIKREFAKYKLECPDSMWKD
PQTEQVQGLYSFAIEMIFGLTVNDVRIEEVTGDVRSCLPSIDSLQFLSQDGKLHAKAIGNLRFRLCQRLNR
VLGLPEFSRETFASTPGVLRMASAFCSFLRVREGLMKSFESQLQQREVLQQNLQQLSGNLQMVEQEL
VRFRAERQSQEATVKQQKEQREELEGEIRDRHTELGRVDEFKEKKTLYGRLELEVEDLRLLEMNLNRQEK
SDLHDQIVHSPEKLMMERRDELRQQQKRLEEQLQNLERVAGRHQQLQSLCKALKKAKKALNLVNEHRNQI
LGPHAAFRSEMRTREKHLKEVSENQRQLNQRVQQQLQNQRRELAKQLEELQDRCDAEEREMRRQLTETRK
EAEELKLALAAQQEDTTSSLSQAKDIEKLLHEQRERQDALVRAIDEQTEKLYAGFQTYISQVEFLRGQMPLS
LDLSQSASFLVDRHPEKSHGLGSQDT SPEKSLERHSQTRQSSFLDENESLNSNRP AHTLGQASGSPCSPR
REDVSGVSFSPA KDRAKNEGGGLSIVDLQEEE EGSAEPVEGETQVTETDGERGSSGGG

>NUF2_Sarcocystis_neurona [993]

MLQQPLGGAAGSSSSAGQGGSLLTSFSSIASCSSPLRELQLSARPLGGAAAATQQQQQLPSFCGAA
VAGVAAPSGFSRMSLASSPLAAAHRMHSPQLL RASGVFPAAAASPPPSCPSRSFLLQGTDPQAAAAAAA
TPGGGGAAAATAAPGGGLSFFSSI AAAA VEAQARA AAAAAGTGAHSAATTNAAAASPATAALGAAGLL
LPPGRLSLEEIKRELAKHKECPESMWKEPETEQVQGLYSFAIEAIFRTLSDVRLEELAGGDVRSFLPSLES
LHFLAQDGRLHAKAIGNLRFYRLCQKLN RVGLPEFTRETFASTPVVLRFASACCSFLRLRDGLLRQLEP
LLQQHSHLQQQLYKVKSDLRALQHELHKL RGEQQMQQPIAQQQKQQEAELHVEIDRLKKTLEPGLVEEHK
RKLQQDSRQAASLSQQQNAQGGVPSCCCCWCCTARLVSLTGQKQKIRTFCVIGVCLAAALAAPLSSWG
RSLAVLLSGLCCSCPRVLTALFRLQQEVETLVLDLSTLKQERDDLLSQVIHSPKKLTARREELRQQQRQ
QDKQLLQLESACAQQQQIQLMERGTKKVKKALHFLSTHLEAVLKPVQVTTRSESRLREKHYKELKEQRRA
LQQTVQQLQQQQREGLQQQLQELQEAVDSEAQQQQQLAQAAQQHVQQQTDRLLKQQEETVLLRQGTEQ
QAALLQEQQEVRRQQQLQAVEAETQKLLAAVQTYFAQVDFLRSQPLDLLVSQPPSSAARPASTDSLSTRD
LDEEEEQQTAAGEAAAAAAFVTRAGEHDCSAETAARSSAAGISTKDTGEDLPLCSSNAITSVVKRKTGE
MESSDN SCTFAISGCSTVGAQPEKATTGAAAYGQTVSVGTLQPRQAAAAAQP LEAAATGRPAGRDGAE
L VVRTESLGNEYAQDEASGERSPSPRTSVDCEQPAAPALHASNEDHAAFCDAATAGEHTAAAAGR TA
ATFDSPADAT

>NUF2_Eimeria_tenella [514]

MPSLAAAGGPEVKRELLRRGLECPESLWKPQAE EVQGLLSAAIKFVFGLSMTQIRVEEAAGEAKAAAPDS
RPFLTLEGRTHLKGIGNL RWFRYCQKLA KVGLADFSRAFVFSPTPQTA RL SAALCCFLAARDAVA AAAA
LPLAEALQQHQ LAAARA EDEL RQTEQQLQQLKADR DAQQQHA ALLH KELQ ALRNTVSSLEVELRSTEDKR
SLQQKQQQRLRQELQEAQELLRLQQQREDYMDQIVHSPNKL RQKRLELQEEQRRQDTRLSSLSKQRQQ
QQLLQQQVARALKKALKKAERILTEHQHNHLEPCVAARQDLRACEKTQKELQQRRQQQLQQQQLQQEYQ
QQQQLLREEEQCSILSREL RERVQQAKEEAE SFWSSLPAQ QEFIRLLQQQQQLQLSIHRKRERHA ALL
AVEEAQQQIEASFCSYLSQMEK CRRALPPPATLSESLANIENE PPTLLLQLQQQQQQEQQQKQQQEEAEEE
FLSAHETESNFPPHKPWLVGAPN

>NUF2_Eimeria_falciformis [705]

MAAPTQQQQQLLQQTVQQQMEQQPQQQQSLYPLPAALQPPPPCMNVESSTAPAISQKVQLLSSPAAA
AAAGAATAATTATTLIPPLPASTLSPLQQQQQQHQRHVSPPY PGFSAGGALHPFYAGGIQQPLQQQLQQQ
QPQQQHVSVYGGVSSPLRQRAT SFFASDLGPLPTL PSSSSSSSSSSSSSSGRLS LDEVKRELQRRGIECP
DSIWRDPQPEEMQGLVSVALELVFGATLQQLRVEEITGEV KVSQPDTRPFLTLEGRT HLRIGNL RWFRQC
QRLAKGVGLSDSRAHVF SPTPQTVGRFATALCSFLRVKD AILAEAGPLAEALQQTQIATANAEEDLRQTE
EQLVRLTADRDAQQQHAMQLRKQQQSLRSTVSSLEME LKSAQERRALQEKQQQRLNQELREAMEELAGL
QQQRKD L TDQIVHSPRKLKQKRKELQDEQREQDARLAELGRQRRQQQLQQQVSRAKKASKAERILTT
KTSLLEPCVAARQELRSNEKTQKELLQRRQQQLQQQLQQEYQQQQQQQLLREEQQCSDISRELRQRVQ
LAKEEADAFHKALPAQQAIRSLQQQLQHLEVS VQRKRKRHAELLQALRAAE LDVVA AFQAYVSQLHVYR
RSLPLPPTLGQSLALS DTEENEPPLHMQQQQPQQQLHASDAEQQEMQSLKPREPLQWPQHSTGTVG
GPSSGPL

>NUF2_Eimeria_praecox [310]

MARPSTQQQHTDYLQPHL TEHSLSEVAPLQQQQQQQQQQQQQQQQQQQQQQQQPSL YP VAA APPSNN
IFSSLTSAAPS FALGP AVTPSAAAAAAAATPAATPA AIPSPSPMANL GSLQQQQQLQQQQ

TYGDVRQNLKIAERRAASEIKGKEELTKEVQLRTCHENLKFDDLQNQHEKEIIRLETEQEQQSQQKLEG
RVELSSFRQQIEAVQENHIKAATALQADAEAQNRLLDELEESHAILVRGFEQYNRIVDEMRSKLPLPLSLSM
SVTCKSTDTEVDNENSQAEPAMRNTMDVPCDDYRRTLLSHLGTPAAVAQPVFVSAAAACTPKHPQTQ
PTSRVMENSPSLKRQSCEGLDDDTIFSPLYPSRRIGR

>NUF2_Pterospora_schizosoma [558]

MYSTPERRNPVSNDRNQIKRELQNLHISLPSSFLWEPKPSDAQEFYRMVIDYVYEGIQLPLNNCSSALPR
DDFKLYSEMKNWHVVLPKLCMFRCQCQTIATKILGCEDFTLHDITRPDPIRLLSFCQAFTNFVRFKEDAKMHS
SARRQNIDELNNNIEKEKEKKANEELYSALRCKLTEREGKICWENQRKREIQSRILATHEKYLIKEKLMEIL
AESERMNKKLTTEHNFLEAKKKNEELLQRAAKSPRRLKKALEDLEISINIKFEQTEGAENKLKYLKLRSHSY
KVGEEQVDSAYQLLRKLKLHENITNKLQEQQVECEVQELKEKIQKQEEGLSSKKHELNIVKKKVEDTKAL
WRKQVETAKAKYSEAESLFNKNKILTEQKQKLHQLVANKDAMMNELSNISNIFEKEAEKCNQLQRSLLQS
VRKGQKIIAAGQKCSSLRICNRREIICRWCTDSAAAPTEYWMTV GALVN VGRWNPMSTNDKLMILNNNNNN
NNNKTSNFIVAANKNVSSLCDNNNNNNKTSNFIVAANKNVSSLCDNNNNNN

>NUF2_Monocystis_agilis [570]

MEKKTLQKLLFRQGVETSPDFFDAPKREEIRIFWDGIISLVFGPHVLLCEVDSVQNALYASYIKETRSRALQL
QSSDHDFVDSSAHSWSIEELLPDRGSRQLQALAIARIIFQCLRHFGLFGIRNISVQLVARPPTHALIMFGNAI
VNWMRFQEQTNEEMAEWNNVNKAEMVKAKRTERDEFRSAVRNIKEQVEQQLPQKQQLEWAVEEMKS
EVCALHNDFNQVKAQEVKVTRYLEQKGEQEKKRAAKNHRLRLEIASMKERIVESPEQIVEALKSLTAKVEFE
QTAAKHESRMLPINQQHRSRVAEDLGKRYELLWSFVEERARIVRENDKLKQQIDVQRKRVSSKGTMNEYM
TQERQRADLVSTQSKLEKARAQKIEYEKMMQMKEEEAVKSTTERKHTEEANERIRREGEENLETTRSIAE
ETRKSEAAAAALIQEYLEASREEDAALSRRVEGIVERYSGENLQSNDVMQQILNDDVAKNFVVKASMRMY
TPATQLGVTRNNTEAKPNASTSAQFINRAGEGLRGRENDESESETCRALGRQFKQMTLKDKVNENRKSE
GK

>NUF2_Selenidium_pygospionis [473]

MASRGRIRTPRGQTGAFVLPVLDFAGVRHDLEQFGIDAALLENWHNPTSQFVEELYSAAVKHLLDGPEELR
CEEMSLDAGPMHIQGVGNLRLRLCSKVCDLIGVDDFTITDLTRLDPERTYRFACAFVNFRVYLEKRLQRH
HPGLRESESVLA LASTAATSQEIGDSAALKALEQTPNIETEIQEQQVSVTGLRTQVVAKLAQLRSLEEELAQ
RVHKEGQLEERYKEQQEELATNQALARDLEDQVVQSPNRLKQIAEISEKVEHEGQKKEAICYKKSLLKND
QLLIEKVEKRLNKKLRLVGLVKEKIQGANTVWKSLSATITTEERRQIAERLKQDEAELRRMLEAIAEKKEES
RARLSALRVT AETDLANRKNHSQTEKERMEKLTAQHIAEEEVETLEETSAGLDELIVAVGEQLEMLAQTL
GKTFTQYRSSLK VEMGKVDLHQNYRRGVLPKLNQSLTQSADDE

>NUF2_Lecudina_tuzetae [305]

VSTQSTWGPTDIPLKIQLLWRCRRHTLLAMYSTPKRRLQVSDFDRNALKRDLANLDFIVPHSLLWEPTAAD
AMNFYQLVCSYVFEDDDAGGGDENELEVRLGLDSLSELGNCHRFSRMRLFRQCQRFYRSIGDEDFLGDL
LHPDPHRINHFCNSFVQFCFKEDNIERLRQKKEEIDMKKNITALDAELATLRDRETELELVKRDTQAEDD
ENERQHAMVKEQLLALHSNFSSCETLSNANASAGKLASELKHI EGLVNAAKSRQDELECEVVRSPGRLKK
SYEQLVQATETEEQNLRQTNR

>NUF2_Cryptosporidium_parvum [481]

MRLQSKQKYQFPDLEMGELMNELDMLGFEGVGSNFWESINHEIAVELYMNCLSI ALEIDTEDIRPEELIGQLP
SSAAGIISENGKSQIKPIGNLRLRYCKILWVMIGIDDSMNDIYRPTPDRIYSFLCGFVNLMR FKEDRWMTY
KNEFYIEEEILDSDVKSNEQIKQKKEDLNNIRVRYNEQSGEIANRRRDNQEYQEKMRS LHGEFLQNQQELK
RLTQSEHDLKEQLKDVEFRITTGNQDIQDLKDQVVQSPERLNTLEELNKSLENERKQIDQISIKNNELKER
QNLLQKTEKRLGAKTFLEQTISGIKDANNIKQSIKEIEHHIEKDWTIEQTTEERLLLQTVEQISLRIQNNQQ
HYE SLIEEAQTLNQEKQFEGQQEFLDVQSSEAFSFERQAELEMEKEIQLKVSHEKAINALNIQHSQNN
LTNYKKHLISKLDINSKYNLKLDET SKENSTINSIENTIVMQHN

>NUF2_Cryptosporidium_muris [496]

MGPKFRYQFP ELDLNELMHE DMLGFEGVGSNFWDNITSEIASDLYMNCLSI ALEIDMEDIRPEELSGQVLLT
AIDLSDQGKHHIKSIGNLRLRYCKILWNMIGIDDSMNDIYRPTPDRIYSFLCGFVNLMR FKEDRWMTY
EFYQIEEEILDVKDKANEQIKQRKHD LANRLKF NELSDEIQSRR IENQQLQEKM RNLHGEFLQNQQELK
QLDNDTQEALKDIEFRITTGNQDIQDLKDQVVQSPERLNTMEELNRSLESDRRLIEQTSKRHN ELQEKL
LRKVEKRLDKAKSFIEQIFQSIKEANNIKQNIK DIEHHIEKDWTIEQSTEEERLLLQTVEQISLRIQNNQQ
SLIEEAQTLNQEQKQFEGQQEFLDVQSSEAFSFERQAELEMEKEIQLKVSHEKAINALNIQHSQNN
YRNELIQRLQLFCNYKNYSKIKTSESIFYNKEGRDNII MGRFTDENDENN SHLN RGECD

>NUF2_Cryptosporidium_meleagridis [481]

MRLQSKQKYQFPDLEMGELMNELDMLGFEGVGSNFWESINHEIAVELYMNCLSI ALEIDTEDIRPEELMGQL
PSSAAGIISENGKSQIKPIGNLRLRYCKILWVMIGIDDSMNDIYRPTPDRIYSFLCGFVNLMR FKEDRWMT
YKNEFYIEEEILDSDVKSNEQIKQKKEDLNNIRVRYNEQSGEIANRRRDNQEYQEKMRS LHGEFLQNQQELK
RLTQSEHDLKEQLKDVEFRITTGNQDIQDLKDQVVQSPERLNTLEELNKSLENERKQIDQISIKNNELKER
QNLMQKTEKRLGAKTFLEQIISGIKDANNIKQSIKEIEHHIEKDWTIEQTTEERLLLQTVEQISLRIQNNQQ

HYESLIEAQTLNNQEKKQFEQGQFELVQSSEAFSLERQAELMEKEIQNSKVSHEKAINALNTQHSQLLNITNYKKCLIGKLNDINSRYNLKLLDETSKENSTINSIENTIVMQHN

>NUF2_Cryptosporidium_hominis [481]

MRLQSKQKYQFPDLEMGELMNELDMLGFEVGSNFWESINHEIAVELYMNCLSLAILEDIDTEDIRPEELIGQLPSSAAGIISENGKSQIKPIGNLRFLRYCKILWMIGIDDFSMNDIYRPTPDRISFLCFGVNLMRFKEDRWMTYKNEFYIEEEILDSDVKSNEQIKQKKEDLNNIRVRYNEQSGEIANRRRDNQEYQEKMRSLHGEFLQNQQELKRLTQSEHDLKEQLKDVEFRITTGNQDIQDLKDQVVQSPERLRTLEELNKSLENERKQIDQISIKNNELKERQNLQKTEKRLGAKTFLEQTISGIKDANNIKQSIKEIEHHIEKDWTIEQTTEERLLLQTVEQISLRIQNNQQHYESLIEAQTLNNQEKKQFEQGQFELDIQSSSEAFSFERQAELMEKEIQNSKVSHEKVINALNIQHSQLNVLTNYKKHLISKLNDINSRYNLKLLDETSKENSTINSIENTIVMQHN

>NUF2_Cryptosporidium_andersoni [455]

MRPKFRYQFPELDLNELMHELDMLGFEVGHNFWDNITSEIASDLYMNCLSLAILEDIDMEDIRPEELSGQVLLTAIDILSDQGKHHIKSIGNLRLRYCKILWNMIGIDDFTMNDLYRPTADRIYIFLCGFVNLMRFKEDRWMTYKDEFYQIEEILDKKMRNLHGEFLQNQQELKRLTQLDNDTQEALKDIEFRIATGNQDIQDLKDQVVQSPERLKNTMEELNRSLESDRRLIEQTSKRHNELQEKLNILQKVEKRLDKAKSFIEQIFQSIKEANNIKQNIK DIEHHIEKDWTIEQSTEEERLLLQTVEQISLRIQNNQQHYESLIEAQTLNNQEQQKYVQCQVLLERQGSEAFTLERQASLERDISNLSRNHEKTINMLTKQHSYLVEIFHKYRNELIQRQLFCNYKNSSKIKTSESIFYNKEESKDDIIMERFTDENDENNNSHLNRGECD

>NUF2_Piridium_sociabile [263]

SAFPFSILSPEDILGFLRDIQFDSTLENLIKPDQHVQDILFTLLCVASKTAESPLALNGNWSQSFRFPDLHESAPLSAMLVKLRFLASVCKIDDFSLRDLYPSSTLSSGNDHHHNNSSRLFHLLSGFINYIRFQQMHENNYLDTFQALEKMQKSSLQEQQEWAKKSEGEVVRRLRCKLAQKLDLVEELRKKKYETDEILDACQSEAASLQSEWKA RKATGDKCSEAFHEAEVVTKEKKALLVELRSQVSSPEQLQKTLDDA

>NUF2_Vitrella_brassicaformis [628]

MRSSASLSLRPASTFNTFADFSNDALLALLHENGVDDADMDKLHKPDEKFIVRALAALLCAATDTSEEILAIPAVCGSDAFKWPPPLHDESVVTNLVSLRKLMAVVLVNDFLSLVDLYKCDSQLCYLLSGIGNFLRFRAFRDDFFDPIQQRIDGLQAEMDSQEAANRQQAGELERLKGQVSAKEMAVRLLEQREADIEGTLSPRLHQAHELQM EWKDMRTKAELKTDEKTNLDFTTRMTDKNVQHLKQQIVQSPERLKESVEESRVQCEEMERQLNEQHRHEQHQRRKDTHLQRAEQQQQNKHQLKKYQENLSATSALSALSAQLSALRTRLDETAQREQEEQEQAWSLKCD EAAGNIAHEKEAHGRVADATERLERAMEVAGRKRGMVETTALKTAGRTELEMQLQEIQKQEKEDILSKES EFLAAHSSLTSIVHRTHTSTLMPLLSALSSLTTLTHHTHTDTETTSTQPQPQQEQQQEAQPQATLTPSL SGSTEAHPHSITPTQHNDEASAPHTPRHTEPSTGKAVAHHETPEFGGGGQQGGNQRDVAGGGQELEG ERAGVEGGSRGGDGIALSVGCPSVAPSVGISPGBTSELQASSDFLQATPGGQHWQKEDGSRFDF

>NUF2_Colpodella_sp [488]

IFCVKSIHEMMKSQMPFLPKAQDICSSFKEFDIEISMTDITNPTVSGVLNLYTRVVAHALGYNMNGLVQLQFN EENSLQEPELYDSFAQRSKVFLYLKILSDTVGFDFCITDFMRPESDRLIRFLSLLQFMHKHQHKLNFREIS SNLLSCSQAFEVAQRERQNAEVRRQEGEVKAQAVRQRISELRLQYEKTAALRALKEHQRSSLKEDAG TTLIEVKKRTEQLYRDAEILKQRAENTKSKIAPNPSALVKERDRLLIEKESANAAMRQAEEIAVTISRREILDK SLAKLNKRALAIKECEELVQMIVKQSVDQKDLLTNAELAAQKLQKLTEEVEAYSRTIESSKRNLISVDTIYADR IANLNAQLAEELETAAAEREPEVEREIVKKIQIFESQKFENEKMAHVEDEAKRIAEEFEDKQSKLYDLVETITK FINQIQGQNLVNASQLNPSNSMQPNFMADSGKNFQIADSSQHSIPVAHN

>NUF2_Colpodella_angusta [493]

AEYMGIFCVKSIHEMMKSQMPFLPKAQDICSSFKEFDIEISMTDITNPTVSGVLNLYTRVVAHALGYNMGL VQLQFNEENSLQEPELYDSFAQRSKVFLYLKILSDTVGFDFCITDFMRPESDRLIRFLSLLQFMHKHQHKLNFREISSNLLSCSQAFEVAQRERQNAEVRRQEGEVKAQAVRQRISELRLQYEKTAALRALKEHQRSSLK EKDAGTTLIEVKKRTEQLYRDAEILKQRAENTKSKIAPNPSALVKERDRLLIEKESANAAMRQAEEIAVTISR REILDKSLAKLNKRALAIKECEELVQMIVKQSVDQKDLLTNAELAAQKLQKLTEEVEAYSRTIESSKRNLISVD TIYADRIANLNAQLAEELETAAAEREPEVEREIVKKIQIFESQKFENEKMAHVEDEAKRIAEEFEDKQSKLYDLV ETITKNFINQIQGQNLVNASQLNPSNSMQPNFMADSGKNFQIADSSQHSIPVAHN

>NUF2_Colpoda_aspera [466]

KNNINAGVLQSNFQMIHEVKILKEQEVIEWFLRELNFELEDNRIINRPDPEFVIDVLSACALTCLSKKKEDFSAIDFKGFNQIVNNEGQLESITLFNQFYAVEQMMEILNIQDFSLFDIIDPQPKLRYYISVIRSYLRHKEVEEDVQREHLIKNKELDEQVASLTQQLNKTAKENDALQDKTNLQMPIIHDKKREIDFIENESRLKREALDILKREKLDFEKEKEELENRVNVENVKIEEGRNEINQLKSQIVQSPEQIVHTLEEMNDEIKKEEVVSTLEGEIAKQNLVNEIFSL QQALDETFLKLMDDIKSSLNKEKNGKKNLKDINKEASTLRDQVEKNKNELQEVLSIKTTEERTENIEDQLQKRKEQVRTTIAQMMDGKAAVQRIYEEQTMNHRNLEKTNLEKELEKTSLEMDRDSMFEEGARLYDVLQEAVM GLHEELSYLFQEGSSKFVNPFETSAKP

>NUF2_Chromera.velia [222]

ADAGRKRDSTMPPRRPLSIEEICSVKRSTKMLAKEDHERLRGHLEKPKPEIVQELYRAFVCECLHLAPEDLKQMRPSAVRYPDLLERC GGVARLHLLIFAKILCQIVGIDDFVLEDMGKPDQRTTHRIFSEICRFIRHSEGRA

QLRVQLEEEQQRLVEERKERRLLEDAKREREELKMREKLEGAEERQREVLMRQELKRKAEREEGA
ARDIDLLK

>NUF2_Platypteron_vivax [590]

MENRRPFTIPMVPFEELVGVRDLLGIKELTSASIEKPTADIVRQLYSVSLEQFWDPREELGQMSFNSLRY
FKYPEIHSESIPVLIFLRHMHQHFLTACGVVDFSLQDLFKPEPKRLRKHLMAIVNFARFREERIQQWAAVQDE
RDELLLERAKLQERSEKFAEEVQIQRQKLEKDLPAIQNCTRKEASAQQKIEQLKRAQVLLQQDLEAKVTQFN
ELESRLKVEDDNLALKQEVQFLNTQVVHSPEKLREQVNAKKAKQYEMKCIADIQAHQRCVAERLRLFEQ
GKQRLSIIATQQQTAKTFEDIREAQQSCKCELEVQQTKLVEDVRRKIQLVDKRLALQQEQLAQQEEWF
EKQKLELGEELIAAAKALQEARESQSRCVIETRHLAEDNKLLRKQIEEVACKHRGEMETLVNACGTLLRAVS
TYGRQLEERLLCDGSNKPVTQLPNSANMISPWPTNKIQRAATTVARPKCKVDVEQPKTVAKPYHGLSTPD
PLVVDEQEDLETPPIRQISELNLDCKTESKTVTFVVTATEENKEHVEDSNEENVEKKPECINSPELKSSSEM
CEKTPGGTRWSVPLAMR

>NUF2_Digyalum_oweni [291]

MTSANDKRAPFSIPLVGNEDILNISRDLGVPLTAQMEKPTAELVKQIYVSSLEQFWDMPKEDLAQMSYN
SLRFFESPEIYSEAVPFLVMIRHLFVSLTMSKPEPKRLKRHISAMLNFARFRQERELAWAAVQDERDEV
LAHSRMTEKVEKSKEELQLSRSKLEVELQDVQECCRKKIADTEHKMEQVDRGTEAAKQQVGMKVENFNALD
SRVREDDAIRMLREEIKFLQTQIVHSPDQLRQHLHEIKACKLKENSEAQLANEHTAASDTVTLQNAEQV
LLRVN

>NUF2_Perkinsus_marinus [327]

MVRGSFPKLTAAQIVEVLKDDFAVIVPPKLKACILALQPEDLSACPAEILRITYDALLVGALDLTEDELNVPAE
FLGQRAAHESTTPIMLMSRQVKWTYTLLRASEHIRSVCPPVACSVVSPYEDTKRTRMLVSALINYIRYKYI
VLAQPKISSTEQMRADEEDALVEVQEKFVEHSKRALQDVKATRERLKPSLTAKHDRHVLDLCLGKAKE
ALVTSIEEHGRVKEELTVGSSERRQIEEKKEELQHVADLVENPERVVEKHKKSEERAALRRTLDEKQAK
LSMAREKTSDWRETNRFEGLMKKARGIKADKEKMEK

>NUF2_Symbiodinium_microadriaticum [694]

MESMKRSPTFLVGTVQQKDILGEFQKREQHYTEEYSKNPLCFSARAFQQPHPAKVRMGYKDADATLTSP
MLLGVCDCGSQLEEFQMDPSLLPNELLRTCEELAMLQLMPDTNIAPQDQYRGPISSLKEAYQETTSYGSTT
VLLAALDNSTRIGHKLHPMIAVLSIGDCELLMLRRTNGRQSELEAVFHTEMQRIDYNVQTPLQLARVDERID
EEFDESIALELVGSSCPGQPPPATARAVSTEGRTYNMLKGFAFKILAPKDTGVLNMIGVHPAVAADNIDRPS
ADNAMAFFNALAEFAYDMDSQQVKAQMPAVTPHPEIYDEAMDFLTIFKLSRQLAMINLVDDFNFKDFWDPV
PKRFRALLSGMINFCRYKEAKVVITGMKEDVQALDSTRLEMVDKLNQVDAELSAAQERHNAELQDMWTA
ENEAAEAKAINDKLTRQRNSADRVEDAERKRTSVKERVRQGEQRIEQLREQVTALQSQIAESPEGLEKEI
EELKSGVCQLKAVLEEKANQRRAHSQRDQVVCRLRHLESYKDELTRVGSVAANAENAKQRAAAAREDLA
NLRQSLEATKQDGTELEQLSKSVVADNERAKQVHAERMEQLEVRRQAAQQHQUELQAKRSEEQRQLHQL
QSQRLELEAEVAIRRAHAAEMGELRVMWKAVIDKAESYNLSLDALFHEHCHEDGTVL

>NUF2_Tetrahymena_thermophila [455]

MSQQTPQSQYQQHSNCQQDITLSNQEIFTIFKNKGFNIDSEKAISEPTQEFFFNNILVYSAFLILRVPLEEINQS
EMCNQLDLEMENGVKAFQIYQKMKYIKQMLDLVGIEDFCSYDIIKPDKKVRRRLVAITQYILYQETIISHTQI
LDKNRADEQKIKRYADELEVLFHSDQLKDRKEAEKNSFQEKMQFYEKLTAEYQDLLKQKQILIKQKSEYQ
NSKLQMEKSIHQAAQEDIQKLIENNQKEKMKIVKSPFKLKDQCLSLQNEIEQQMKELDLIKKDYLFCIMKNNYN
DKLLQKIRKHLTLPSLTLEEMIKKESEANKILKNAKNEIKEQLKEEVSHQSYQKISDYHMDLIAQLDQKNKLA
EEKKKTYEDEINNMLKEQQKLKEELSEVQKTIYEIQQMQANLEQKISEEESSHQMYMNEKKSQMEIALKFK
EYKNDVFLLLNESEQID

>NUF2_Aurantiochytrium_limacinum [447]

MAAFRQHEQQSFSPLLVDDIELMCMEELQVLSAKDLEKPSAPTVRAYESLCETCIGVTRDELNPLED
AMSRIEFPELHEESIPQLTYLRYVQKLLGAAGVHDVLLDLTKPDPKRFRRNLSAVINFAKFREERLMECEEL
TNNTDKLLERKDEIGKGCTDIQEKENRVRREKREAEQPRVEQLKHDCRDLEAEKLQSHKQHASLQHETKQLK
AKSNELKDLVSTLRFQKLNVVQETERMEGNIVRSRPERLKSELADLRERLDGERDENSRLSENLEELRAKDK
ALARVEEEMERLHSVLMELVGMKGCKQALKETKNMNSSIRDHDQVLRQLRSQRDETEKQIDITKDRVTRL
ANTKSLKVAASHALSESKKEREQLQKSNKSLRAKLAERKDEIENVENELAEQNTKHEKEVEMHRTAFKHL
ENVIEETWEPLNKAMAEAS

>NUF2_Bigelowiella_natans [431]

MKTTFPILKIREIVKCLRELHISLSTEEIKKPTRKKMMEVYEKFLHHMMGVSKEELRQPAFDSDMDCFEYPELH
EDSVGEFSTSFSYFLLRLFHAVGLTRASMRDFMFPKSKRTIKTLSALINFAKFREEKLPLYTKFCTDTEELQNL
EKVAIENDSFKANLAQLKEQREQDAPQIEKLSQETEELAGKIEKLNELHASLRTDIRSRKTALEKTDERISTS
KFEILNVKEENNDLKEQLVQSPERVLKDLTKTINVEKQELQESAECARILQNKVALLQKKLAKRIATLKEYGKI
MKTICKLQRDIKDRKIVGENKEHVKSTNITVDHMKRQISQYQTKLFELQEQQFDEKRQAAKLLNEAMKKK
MESMHIGAQEKAKIEHNTMVYKTRKHELELAKKTHSQMQHWKDNYADLLKTVRRYHDHLRNKITAN

>NUF2_Arabidopsis_thaliana [440]

MSAYEYPRLSRSIDIITALKDAQIASVTEDLKTPTSDFVSELYTRILYLDALDEEKGQVDFEALEQLENPDH
HATSMQAMKLYCKVKDMLEMLDCPLPISFKDLLRPESSRTEFFISALLNYGLYKDSKMDLIRPKAEELGLLD
EQRKQCEAKVAQLNAEIGEDEAVERDLPVQELEANIEQLNKKILELNQQMSLRATFQKMREKSTQMDN
EISKAEDLVETVQENANLRSQIVQSPDKLQGAEEKLVLGETKKAQSAMVTFQEKAAILEVFEKALKKIL
KSSSQLQLINEQVTNAKTVKEFKALKDKLSEDGVAYKSLEAKVVERERIVEQLNESLKQLEKEKAVMFDD
WTKQLNELKVEVESRRRELETRQTNVESVAMVDDNTAKTNQRQSGEAKVKKLAKYEEIVKQFHEYTV
SFDAFLPSL

>NUF2_Trichomonas_vaginalis [407]

MSTTLNLSKQKIIDISKGLGIANADIIFNQGQTQFTQMLDLVFVQMLVEYYPEHAYKLPKFALRPEPIPDTFAQ
GIDFLNKSLLLKHLTNELNLSFGIKNYHDEKFLTKFFSNIINLGRFNESYIKAEQNAARERMAIVTDELNQIEI
KWQQTLEEEQRKQQLLTEMADAEDIDNKYESLITAIDKQQKKNEALQQKIDRQNIELQSKKNQIQENNDEIER
LTNAIKSLDDILNVKPDEERQRLEKEQSNLEALRTSKSLEDEKAKVQQELDSFNEFYNLLFTNMPDFQKYG
ENASKKKQVDMQLQDQIARRREDMNNLQQAIQEMQKKKDAIYEIDNKAAEAKNISEQIQNAKKLRLMAKEK
HKKELANYRYQIDDVRSQLDAFNRELWKMMNEIDQQDKESEV

>NUF2_Giardia_intestinalis [484]

MTSHLKTKKSLSSRMDGLTQPFTDQPMEVMTDEQLKEIHEKKIIDAINGSIFGLAKVVKETQHIEPVATLDQ
GPLKPSSVLATLMMFYNTAYLLPLSKVIDDIPPQFKALFSEHYAEKYQYILNIYAQYGILKEFLADCKYPGFS
FTDIFMPAFKRYKAQLHNISEFLWFREAVIEILHSALTEYKTAYTQNKPVLESASALKGISRTDETVKLRETA
SQLPSRIDPLIQRYNRALESQQQLDKENSEFTNRILVLTNELELEGQVKKKTEEYESEKSTLITKKQNDAVL
QDIQAAEALGNDLRTQCDALLSSINIFEQHNNNDLLGITDKITEILDADQYSSSSDNLQAQLESLQVMKDTVN
TLDAQINASKQEIENMNNAMQYGSAVEQEQDNPSLRADEEEELRRLTEDCEGLEATYAKLSQLVRALQQEIN
DENAKITRVRVDTAEKIDQLSDKANEMMSMFVAQLTGMISKLSSM

>NUF2_Naegleria_gruberi [428]

MGVVPTLTQENFTKPNPDLALKVFGSCMEKFLDETIEDIKQRNREYISQAPTQGESLDVYTNILEFIEVFKKI
QKVLVIGKLSDFSVDIIPKEPKRFKLALY AFLNLRFREQIIEVYNDIISEFDNLQNEKQELLIALERDNNALNS
YKEQEKVNEGKLKILGDKVREMTQAIQEMNLKDMELKVTLQNVQAENEHLDKEIDISHHHLEVVTKENQSL
KSQIVSNPDDLKKKISLLRTCIETSHRELKTLEEEIQQLTLSDMKNAEQFTKILSKSIVLAEQLEELRNKKKEIT
KSNKTKQESIEKLQSKLMEFDNLKKQEQKQNQSLSEKIERIEGQFYEKLERAMSMENDALKEKEAQEAHCR
EIKEKNEAAKQRILELERYRSQMAQRYKSEIENIVNGCRSVEVGTDNFHNNVISAVSQKNKH

>NUF2_Dictyostelium_discoideum [469]

MSKYSFPILEDDEITSDEMGHRTFMEELNRPTSAMIREFCENILVFTGYNKSTSHHPSQYGMKDQLTQGIE
LFTQDGAIIGEITFLRAMSNLMKVVGFLDFTYKDIYNPKAARVKKIFSGIINLGKFMELKISIYSDLKLNKNSCE
KRDITLGKEQKLLELENKRMKKHEKDLIVNGLIEQHQQQDQRLKQANHYQNEQCKIILNKTNKNESIKQEIL
NLISSIENTIEECARMQVLIVPSDKIKKVLYDMKQKLALKKEGLKEFEPSMSKLQNKSKQLEKINKNIRTLE
LVETYQKENKLYKSTKLSKEATKVNQDQRQINELDIRKFNDDLLSDESLKKNRDNYITKKSELINLQS
ELKNQKAIIEKERESTQIKIDHLIKE DIVRTSIAKLKSNEKEKREIFKSFSTLVDSIHTYHLQLFNLMENNLLID
YHQQQQQQQQQQQQQQQQQQDHN

>NUF2_Saccharomyces_cerevisiae [451]

MSRNQDVFPILDQELVICLQSCDFALATQENISRPTSDYMTLYKQIIENFMGISVESLLNSSNQETGDGHL
QEENENIYLDTLNVLVLNKICFKFFENIGVQDFNMTDLYKPEAQRTQRLLSAVNYARFREERMDCNSFIL
QMESLLGQLRSKFDDYNLIQQQLKQYEDVGDNIPDEQELQKLEEQNKELEIQLKKLTQKETLSIDYNDYKI
SKQSIFKDLLEALSFQIVELESNRDKLIKISNTDMEELSEGIKELNDLLIQRKKTLDDLTAQQKNLQDVTTFETII
SELYDVLRISSEVQESNRTETELVGLKQNLINNKLKLMNVLETGIMYKLEILQEQLDLQLKNLEKLSQDTKEE
SRLNDTKLMDLQIKYENEIKPKIDKTDIFIQEELISGKINKLNDEIKQLQKDFEVKEIEYEYSLSGHINKYMN
EMLEYMQ

>NUF2_Homo_sapiens [464]

METLSFPRYNVAEIVIHIRNKILTGADGKNLTKNLDPNPKPEVLHMIYMRALQIVY GIRLEHFYMMMPVNSEV
MYPHLMEGFLPFSNLVTHLDSFLPICRVNDFETADILCPKAKRTSRFLSGIINFOHREACRETYMEFLWQYK
SSADKMQQQLNAAHQEALMKLERLDSVPVEEQEEFKQLSDGIQELQQSLNQDFHQKTIVLQEGNSQKKSNI
SEKTKRNLNEKLSSVSLKEIQESLKTIVDSPEKLKNYKEKMKDVTQKLKNARQEVEKYEIYGSVDCLPS
CQLEVQLYQKKIQDLSDNREKLASILKESLNLEDQIESDESELKKLKTTEENSFKRLMIVKKEKLATAQFKINKK
HEDVKQYKRTVIEDCNKVQEKGAVYERVTINQEIQKIKLGQQLKDAEEREKLKSQEIFLNLKTALEYHD
GIEKAAEEDSYAKIDEKTAELKRKMFKMST

>NDC80_Plasmodium_berghei (PBANKA_1115700) [589]

MNNHSVYKFNPTMSVFNNNTLNNSIYISADKKSKSRSLNTLSFYKPNASVYVKKIDKTDKDNVKTLIRYLG
WKNYSGTISPNLFKNPSMTDLINIWNFIFKHDPLIEVNKDNYGEVVLFSFYRDIGYPYTISKSTLVAPTTGLQY
NTHLSALAWLCQLLIFEAECAFNDINEEKDVNFYDFNEDCEIKMEDFILHSYRSYVNKEEKNIKDMLSAKLEK
ELERLENDIEKHEDIQEKKKKIEEIKSNMKENEELIKRNVLREENIKIKSLHTNSLEETIKLEK DIEICKKANID
EKNKTEEILEEINKIKETLQKQTLNKAQFIQMENIDKNKEKIKENIKNEIKSLNNNDHPILSEKLNRRNNDLKKLA
KSVNDKLTEILQILNIYNNLSQSNWNNVSTISINIDGLTIDTMLNVKWKENKKNIKYIEKDKKELSNSINIEEN

ENQIKTIENDIEQIKDCIYDLEESIRKTNENISNFISDSNNRFNQIIRQNEETLEEARSKQNEANEMFKDVIASK DDKINTLNKIKMENEQLFKEKLSTLQKACSVLIELKSYSKSYISIVLDEKKKELELYKNLHQSVTE

>NDC80_Plasmodium_chabaudi_chabaudi [590]

MNNHSVYKFNPTLSVFNNNTNLNNSIYIPADKKSRSRSLNTLSFYKPNASVYVKKIDKTDKKDNVKTLIRYL GWKNYSGTISPNLKNPSMTDLINIWNFIFKHVDPLIEVNKDNYGEVVLSFYKDIGYPYTISKSTLVAPTTGLQ YNTHLSALAWLCQLLIFEAECAFNDINEEKDVNILYDFNEDCEIKMEDFILHSYRSYINKEEKNIKDMLNAKLEK ELERLENDIEKHEDIHEKRKKIEEIKSNMKENEELIKRNKVLREENIKIKSLHTNSLEETKLEK DIEICKKANQD EKNKTEEILNEINKIKETLQKQTLNKAQFIQMNMENIDKNKEKIEKNEIKSLNNEHPILSEKLNRRNNDLKLA KSVNDKLTETILEILNIYNNLSQSNSWHSINTISINIDALTIDMLNVKWKENKKNIKSYIEKDKKELANSINIIEDNE NQIKTIENDIEQLKDCIYHLEESIKKTNENINNFINDSNNQFNQIIRQNEETLEEAQNKQNEANEMFKDVIASK EDKINTLNKIKTENEELFKEKLSTLQKACSVLIELKSYSKSYISIVLDEKKKELELYKNLHQSVTE

>NDC80_Plasmodium_falciparum3D7 [591]

MNNHSVYKFNLTHSVFNNTNLNNSIYVSGDKKNSRTSLNNLSFYKPNASVYVKKVDKSDKKDNVKILIR YLGWKNYSGCITPNLFKNPSMTDLINIWNFIFKHVDPLIEVNKDNYGEVVLSFYKDIGYPYTISKSTLVAPTTG LQYNTNTHLSALAWLCQLLIFEVECFNDINEEKDSLSSYDFNELNEIKMDDFILHSYQSYINKEEKNLNDMLRTN LEKEIDRLDNDINNKIEDINEKKKKVEEIKNHMKENEELIKRNKVLCEENKKIKQLYTNSLDETEKLEK DIEICKI SNQDEKNKTCENILDEIKKVKDILQKQTLNKAQFVQMNMENIDKNKEKIEHIKNEIQALNNEYPNLSEKLNRRNS DLKKIARNINEKFIEIVENINLYKNISTSQWNEIHPININIDSTTVNDMLNVWDFKSNVKQFIDKDEQELKNS LNLIEENEKQIKVTEKDIQDLKQEIMDKENLVKQMNEDTNNFLNESTKTFNQIVAEENEKMLEEAEKLSETNE MFSEVRTSREQKENELEKIKEENEKKFKEKLSVLQKACAMLIELKRYSKSYISIVVEEKKKELDLYKDLQKSV VD

>NDC80_Plasmodium_gallinaceum [590]

MNNHSVYKFNPTLSVFNNNSSLNNSVYISTEKKSRSRSLNALSFYKSNASVYIKKADKSDKKENVKTLIRYL GWKNYSGSISPNLFKNPSMTDLINIWNFIFKHVDPLIEVNKDNYGEVVLSFYKDIGYPYTISKSTLVAPTTGLQ YNTHLSALAWLCQLLIFEAECAFNDINEEKEMFSDDLNDNEIKMDEFILHSYKSYINREERNLNDIMNAKLD KEINRLENNINKKEDISEKKKKIEEIKNHMKENEELIKRNKVLCEENKKIKLLYTNSLDETEKLEKIEICKKSY KEEKEKTEKIIIEEINKVKEILQKQTLNKAQFLQMNENIDKNKEKIEKIEKNEIKNLNNEYPNLSEKLNRRNNDLK KLAKNINEKLIEKENINLYKNISISEWNKINNININIDSFTVDNMLNMNWDFKSNVKQFIDKDEQELKNSLNLI EEHEKQTKIVEGEIKYLKECILEEENLVKKLNEDTNNFLNTSTNKYNQIVSQNQKMLEEAKEKNEANEMLK EVLASKEEKENELNKKMENEKIFKEKLSILQKECAILMELKNYSKTYITIVLDEKKKELELYKDLHKSIVE

>NDC80_Plasmodium_knowlesi [591]

MNNHSVYKFNSTLSVFNNNSNLNSSVYISGDKKSRSRSLNALSFYKSSASVYVKKVDKTDKKENVKTLIR YLGWKNYSGSISPNLFKNPSMTDLINIWNFIFKHVDPLIEVNKDNYGEVVLSFYKDIGYPYTISKSTLVAPTT GLQYNTNTHLSALAWLCQLLIFEAECAFNDINEEKDMNCSDYDFNEDGEIKMEDFILQSYKSYINKEEKNANDVLN AKLDKEIDRLQNDVSAKSEDIVEKKKKIEEIKNHMKENEELIKRNKVLCEENKKIKLLYTNSLDETEKLEKEIEM CKKANQEEKVKTEKIVEQINKVKEILQKQTLNKAQFLQMNEDIEKNKEKIEQIKNEIKSLNNEHPTLSEKLNRR NSDLKKLAKSVNEKLSDILQNLNLYRNISSEWNKINNVSINIDSFTVDNMLNLNWDFKSDVKSFIEADQEE MKNSLRILLEDYEKRIKLLEEEIDDLSKNIQDSEARVKKLNDTDFKVTASSDKFTRIVAQNEKMLNEAKEKSD KANEMLKIIAEKEEKENEFSKI TDNEKMFKEKLASLQQACAHLVELKMYSKNYASIVLDEKKRELELYKEL HKSVE

>NDC80_Plasmodium_malariae [591]

MNNQSVYKFSSTLSVFNNNANLNSSVYISGDKKSRSRSLNALSFYKSNASVYVKKVDKTDKKENVKTLIR YLGWKNYSGSISPNLFKNPSMTDLINIWNFIFKHVDPLIEVNKDNYGEVVLSFYKDIGYPYTISKSTLVAPTT GLQYNTNTHLTALAWLCQLLIFEAECAFNDINEEKDLNFSDDLSEDNDIKMDDFILSYKTHINKEERNANDMLNA KLDKEISRENDINTKNDDIIKKKIEEVKIHICENDELIKRNKVLCEENKKIKLLYTNSLDETEKLEKDIEMCK KNNEEEKNKTEKILDEINKVKEILQKQTLNKAQFLQMNENIDKNKEKIEHIKNEIKNLNNEHPNLSEKLNKNN DLKKLAKNVNEKLNEIMQNVNMYKSISISQWTKINNININIDAFTVDNMLNINWKDKKADIKTYIEQDEHELKN TLNLIDEYEKYITTMQDQINELKSNTQKEDAVKLEDEIAKFVIMSEKKLNQIVSQNEKMLGDAKEKNNKV QVLKEVLESKEEKENEFNQVKMENEKMFKEKLSALQKACALLIELKSYSKSYISIVLDEKKRELELYKDLHKI VTE

>NDC80_Plasmodium_ovale_curtisi [591]

MNNHSVYKFNSTLSVFNNNGNLNSSVYISGDKKSRSRSLNALSFYKSNASVYVKKVDKTDKKENVKTLIR YLGWKNYSGTISPNLFKNPSMTDLINIWNFIFKHVDPLIEVNKDNYGEVVLSFYKDIGYPYTISKSTLVAPTT GLQYNTNTHLSALAWLCQLLIFEAECAFNDINEEKDVNMSDDFNEICEIKMEDFILSYRSYISKEEKGNDMLNA KLDKEIEKLKYDINVKNENILEKKKKIEEIKNHICENDELIKRNKVLFDENKKIKSLYTNSLDETNQLEKEIEMCK KTNIEEKNKTEQIVEDINKVKQILKKQTLNKSQFQIQINENIDQNKEKIEKNEIKNLNNEHPNLSEKLNRRNNDLK KLSKSINEKLT DIVDKINFYKNISQVWKWNKILTININIDSVDNSMLNLNWDFKSDVKSFIEADQEE EDLENEKIIQDEINLLKEKILHKENEVQRINEDNTFVNSSRERYNKLVSENEKILEEAKRKNDEASDMLKEV QSSREEKEIELNKKMENEQLFKEKLSLQKACALLVELKSYSKAYISIVLDEKKRELEMYKYLHKSVSE

>NDC80_Plasmodium_vivax [591]

MNNHSVYVNFNSTLSVFNNSNLNSSIYISGDKKSRSRSLNALSFYKSSASVYVKKVDKTDKKENVKTLIR
YLGWKNYSGSISPVLFKNPMSMTDLINIWNFIFKHVDPQIEVNKDNYGEVVLFSFYKDGYPYTISKSTLVAPTT
GLQYNTHLSALAWLCQLLIFEAECAFNDINEEKDMNCSDNFNEDGEIKMEDFILQSYKSHINKEEKNANDVNL
AKLDKEIARLQNDVTSKSEEIVEKKKKIEEIKNHMKENEELIKRKNVLCEENKKLLYTNLEETKKLEKEIEM
CKKTNQEKEVKTEKIVEQINKVKEVLQKQTLNKAQFLQMNEDIEKNKEKIEQIKSEIKSLNNEHPTLSKLNN
RHSDLKKLAKSVDILSEILQNLNLYRNVPSEWSKINVSINIDSFTVDNMLNWKDTGKVSKFMEADQ
EELKNSLRLMEDYEKRIKLLEEEIDLKEDIQDSEARVKKLNDDTDQFVTASSDKFSRIVAQNEMKMLDEAKE
KSEKANEMLKEVIAEKEEKENFTKIKTDNEKFREKLATLQQACALLVELKIYSKNYASVVLDEKKREELY
KELHKSVVE

>NDC80_Plasmodium_yoelii [590]

MNNHSVYKFNPTMSVFNNNTNLNSSIYISADKKKSRSRSLNTLSFYKPNASVYVKKIDKTDKKDNVKTLLRYL
GWKNYSGTISPVLFKNPMSMTDLINIWNFIFKHVDPLIEVNKDNYGEVVLFSFYRDGYPYTISKSTLVAPTTGLQ
YNTHLSALAWLCQLLIFEAECAFNDINEEKDVNFLYDFNEDCEIKMEDFILQSYRSYINKEEKNIKDMLSAKLEK
ELERLENDIEIKHEDIQEKKKKIEEIKSNMKENEELIKRKNVLREENIKIKSLHTNSLEETIKLEK DIEICKKANLD
EKNKTEEILEEINKIKETLQKQTLNKAQFIQMNNENIDKNEKEKIEKNEIKSLNNEHPILSEKLNRRNNDLKKLA
KSVNDKLTEILEILNIYNNLSQSNSWNSINTISINIDALTIDTMLNVKWKENKKNIKSYIEKDKKELSNSVNIIEENE
NQIKTIENDIEQIKDCIYDLEESIRKTNEINNFSNDSNNKFNQIIRQNEETLEARSKQNEANEMFKDVIASKD
DKINTLNKIKMENEQLFKEKLSTLQKACSVLIELKSYSKSYISIVLDEKKKELELYKNLHQSVTE

>NDC80_Babesia_bigemina [600]

MSSLRNFRSALDSQTTSSTARHQAAQPAAKSPLTNKRECVQCILAFLAARDFHPCATAKELLRSPPPLQTLIDI
WNFLFSYIDPNAHTKENMAVEVPKFFKEFGYPHIMKTSHLRPTADHQWEANLVALSWLCKLLLQECECF
DKPFEVTKKRHGSPYMGDFGAHGKATISNTMVAQFVTEQATKHFQLYLKGEENSQQLVEQFHARDVTEL
AGVQDSVDGKIRELEELRGQTLQEELESFARTKEWIEKANVELQRIEELTASVRGTCVQAAETLKRHKQ
TLRDEEAALAAQEEQNADIEKLIQQDINKNVQELNNTIRALKVRIADGNRKIKELEHDIASSKTTAQTMES
QLVRLQKALASTHESIRNFLVNNGRHAESWRSLGQLRINPQGINSEILGVEPATYNGVLQDVINKDREATH
ESSEARTELEQANQHLEALGNEIARETSEILRETSSLIKNQHVEEKDARVQLEASELAAVIRAAANDELVSRL
DNRDLRGLTGAQUESTKRELEAVEAEVAVAHAQLQQDAAAQEAWQVFVNRFKEMFQIARSSKESNCHAL
RGDRRLIGLEHRLFSGGDDTQHEASETTC

>NDC80_Babesia_bovis [598]

MDSILSNPRNVKAATGEPASSVTRTMQQSVMKSSIANKKECVQCILAFLGKGFQPCSAKDLLRSPPPLQTL
LLDIWNFLFRLLDPSANVTKDNMVEVPKFFKEFAYPHIMKTSNLRTPTADHQWESNLIALSWLCKLLLQE
ECFGKSFNDNAVKRQHLPMSGINAQATISTSADVHFVTEQARKHFNLYVRGEENSVELMADFQSAISDLILST
QDSVDKMMIELEVVRDKALSLQDELENYSRTKEWMKTAKELRQRIEELTASIRGSCLA
HVSQDLGVYHCKPQV
EALKSHNHTLREEEAALSLQEEQNRELERIIIEKQDINKNAVIELNQAIRKLKAHISDSNRRIKELQHEIVIGGT
TQLAEDQVMRAGSALESLHESMKNILANKGQYAEAWHSLEPLKINCKGHTSDILGVPPSAYSNNVDEMIN
RDRDMLRMSRDTRTALELSNQELESLYNSIAKEISDLRKVSSLFQTQLREEKDSALQAETNVANSIRATK
MEELVGTCVVYNILKERAKSELAGIQEAVAQAQNRLREQESQAQTKWQETVNLKEAYTTARNTKQENRE
MLQDLIDTENRILQENKSRQQNTA

>NDC80_Babesia_microti [551]

MELFQGLKKKVEGEIKSTSKE TENDIKLILKFLEWRKYPTPKSLENPTVNDIIDIWNCVIQTDPMIIVTRNNM
AMIVPSFLHIFSCPFTVNKASMAAPHLGAGLSANLHFIGWICKLIIYEDYCFCFQVPNQKFYNDTDLIATNITA
EEAAISYVKRLLIGDNYHIAMRSGNDNTATDSMVKNIIHTIRDYIDEINTVIQSGVQLEKIQHELTQLKNIVGLR
DELKARINTLNNNDIENIDGAVVKGKRDLSKKEQEIEALTLKLEQETKLNKLNEEAELELQKIIITSQKLSKNEIEK
LKTELNAKDTLASIDFKKKVVQSIPTLENEIENITKELYLKQCEIMTTQHKVSETIAVHKSDYAIPWIKLDAP
RINISAETINSIIGDELKRYESHLEPLMKRNLQYHDAKQTLININEYVKANKRNLDAQNELESLQNSIISANK
LLDLRSRSLDEHDSRFENLLKESKDRIKANWNKAKQSLAKVATKHAELSKILTEKKLYLSSFWNEQVKQIELI
VQLIRTAKMTNYNKLKELNKYKENILANLK

>NDC80_Theileria_annulata [504]

MIQQFLPHKSQNTKKLNTKKECVNFILHFLAWKGYNACSVSDLRSPPPLKVLLDIWNFLFRLVDKNVTITKEN
MAVEVPKFYNDFGYPHIMTTSHLKPTAERQWESNLVALSWLCKLLLQEHENFKDFEKSTMTQTDSYM
FDAAFKMNIKNPSTYKTVTEWATEHLQLYINVVEKRKSIPNDFEELNKVLSRLEKSVNSKTQLNHLRTVS
EELQKEIQEFDDLKERVNTKIEIKTIEDLTNKVKDYCNRMKNELELRESILTQETNSLIQMENSKEFNDKMN
LKIVSSKSYKIIQDGKNIKEINQDINQLKLNLCNSKKMRELENDRLSLSSEVSKKNLYVKSLSASGIDTEPW
KEFEVLLKKIDPHSPDHSKSLSDLMNLANDIDSESLQGGSVSDLQKTLKSLEQEIEDLRVTESDYSNSTTF
NKEKSHLISENEYLVSELTKELENTRNDLANKRLLESRSQANYDLQKYIKGLREVSYCLKSIYII

>NDC80_Theileria_equi [555]

MTYSHGLKATSTPLPLGTFKRNVVKKECVNTILQFLAWKGYQLCSERELLRAPSLGVLLQIWNILFKFVDA
NVEITRENMTVEVPRFFKDFGYPLIMETSHLRTPTADHQWERNLLALSWLCKLLLQEHNFEISAPRSNVNI
PSTIPKANISNKLLENSITELTTKHYQRFINGCETSKELEAFSAELGSIRADLQFDIDKKMELFEQLRNEVSQ
LQDYICNFGNVKHIRIKVLSSDLEQIERATRNLKDSFVISEHELESSKHLRKESSHLLRLQEKNKEINDMIYNN

INGVNLKDLNDRIYTLRAEISTLVERIKEYESEIPLLSSNIISMSNTLLKLHASSSSLEGIKSALLANGIEASSWVNLEKITLNTEANAADEILGISIDSFEAKIEEIFSSDSQKFQETTWAIKQTNDTIAELEIAISLENEINITNKKLEN SQDSANVSRLQAETEDIVTMIKSCAQENLQTCKDKLRSVLEELQGANDQLELEKANAKKKLESYLEAIRTIV QVCHEQKLKNCHAIKGLVSHKVATIEGLSPLLKDNG

>NDC80_Theileria_orientalis [549]

MSTKKVISRKECVTILNFLAWKDYHPCSEKDLLKSPPNVLLNIWNFLFRLIDSNVNITKENMAVEVPKFYN EFGYPHTMKTSHLRPTADHQWESNLVALWLCKLLLYEHQNFDKEFENKTMLRQTDSYMNFNVASKMNIK NTLTSMTITELTNKHFQFYIKSEETGKSIANDVDAKLNMIAKLQSSVDSKKSELETRSSSVKMQNDIEFD KLKERVSTLSLELSQIETLTENIKDYCTRLKRELDSSKHILNEETNYLLELESKNKEINDMMKTRIDGNDIRGL NIAINELKISLSNSVSKNIKMLEMEIPTLSSNTKHNLAWSLSSVLFNLIPSQVSKTYQLIMNSLSTHGIDVSY WKKLDRLLKSVDHSVDKLLSITDNCLKIVDEALVSETLTFKNESEVKIQHNETLSLEKSEKLLKEEIAELQA KESEYFKSSLFNKEKSQISETDYIVSIIKSTAKDEMVFKLIISLLFQENSKRELENATNDLQQAHKRLSEEKFQ AKSNLNRYLNNIKELVEKVEDYKSRNCNMKELV

>NDC80_Theileria_parva [140]

MMQQFLSQKSQNTKKLNTKKECVNFILHFLAWKGYNACSVNDLRSPPVKVLLDIWNFLFRLVDSNRITKE NMAVEVPKFYNDFGYPHIMTTSHLKPTAERQWESNLMAWSLCKLLLYEHEVCKLLGKNIEFSQGF

>NDC80_Cytauxzoon_felis [523]

MNSIFGKGIGGLSRGRDGRSIDDKNECVDVIIKFLISKGHNICTFRDLLQPLPVELLINIWNLFRIDPNSTII TNENMSVEVIRLCKEFGYPLSINSNQFVIPIDESEWTKTLKLVWLCKLLYYEEHCFNKEDEDDVKDEPVLSS FIPNKGSKINISNVMIAKCLTDVTTKHYLYVNGRKDGQMLMNDVNDELNRISINLQGSIDDKMKLLELLRGN ILDIQDDKESFDDVNSLEAEIEKLQLEVNDLNDVSYLEEYRDLLNEEEVKEKYENLKDELKKLEVSSISDNE NELIKLKDELDECRCNVKESYEKMNSTYEKVINLLLNDIDASSWLNDKIELNFNGNTVDKTLGINCDTFMKI IDDMIATEALTYSSIHKAKAQEDDEIELKKRIDTVNNEIEILQQQKSSSFNNGNIGNIGHSAADAPEGGTPGD RGGSIHLINRMDFMKSEYNLLLDYKNLNNEVKRIFNEYIESLSSLLTIEDDKRRTENYIKCIIEDDKRILENIE NVN

>NDC80_Nephromyces [609]

MANLGDSYTYGGPTGYTEDPLYKERQRVSLSEASFLKNPLTSGPRNTINRRNSDQRNVPPDLIKRERMKT LMRFLSWQEYPSEIFNDLQAPHRSLLISLWNFILKRVDESIEINTK NANEEVPRIFKELGYHSAISKSSMVAP GTGHQWFHHLTALSWLCELLIYENAIFPSISQGSGVKISLDNSFGTSLINSKHADPTTKEI LISTVTLNKLVS KSFIPCMTGNENKDAIKNAVLSQLLEVVDILRASHQKKANKLKELDLEHKRLQSIQDHIQKSSEINEMQKKY NHFEDRVEEIRKQIPMADKRIKMNEKSISSKNIELEREISTVTQLRDQLQKQLSHNEVTLKNTIQDHDAIR KYNETLETLDENRLLKEQITNYCEELEYCNSINEKLLISRRNVKKLKSPSIKLWDQLLAFSFTFDSPTVDK MLNVEWKQKELLENIKEEETYLQKIKSELSAKSVPNQQKQLEILIEKTKILESDENNKKALEDLEMEMNTK LQEINKIKEEALNQLEEEAELTSKLSRSQAHSDGLQEAQKQTQKFQECIEKWENIEDMCKDLSHAKINSI NGFQRLVDIKIEKLHKLNTDEVK

>NDC80_Toxoplasma_gondii_ME49 [728]

MHDPSDLRGLASGASALGG SAYPLSSPLRSTSPYRADRLAAPSSTLPLFASPMRSAAGDSRLRAFGQENE STHSFLDSHGGFSTASC SAAGSAARETFRREVGQFSFYKSSLPPGSAGTVGRGVGKGPGGVQQLLGNE EAKRESVKTII RFLCYSGFPQLSPKIFVAPPNLLVEIWNHLLRACDDSVQVTNENANEEVPRLFKELGY PLTIAKSSMQAPNSAHQWPLHLHALSWLCELLIYESEEVFSRDPPLLNP AIEKKDAL SAGAVVGEEMISRM LH YYPQSGNGRDLTHLQQSLHSRLQRDVEQLESSIASRERQLARAQQELRRISAELEAN AALPPEIDRLCGDL EKLKGDIKQQRSA SEQQEKEIDAKQSRRNALQSEIRQVAETQQLDAQVKEQGLSKAEVEKIRSEIHLRE RVALRSKDVEGQQQLLAGVEDEINRRGEELRVTT C NSLLQRCM QDTETAKFASDTAWASLSRFATSAT LSGDSSSSFASSRGSSESRRSSSRPWAPQSHSEPEAKGNADEHLATHEQLLGPWKA WKAELLRLVQSD KQVDAQSRRTDEKTIGEIEKLKKARQQLDSTRRTEERKISVLHEDLKLEEQQKQVLSSEIETLQRGV S EHRLAVEAELLKAERQANDLRLLYEKTEKENDAEVAARIDNLKKIQQDGLAAKREVLHALQQLVQNKEHAC VGYEKDLLAVVSRSEKYTLGC

>NDC80_Hammondia_hammondi [728]

MHDPSDLRGLASGASALGG SAYPLSSPLRSTSPYRADRLALPSSALPSFASPMRSAAGDSRLRGLGQENE STHSFLDSHGGFSTASC SAAGAAGRETFRREVGQFSFYKSSLPPGSAA TVGRGVGKGPGGVQQLLGNE EAKRESVKTII RFLCYSGFPQLSPKVFVAPPNLLVEIWNHLLRACDDSVQVTNENANEEVPRLFKELGY PLTIAKSSMQAPNSAHQWPLHLHALSWLCELLIYESEEVFSRDPPLLNP AVEKKDAL SAGAVVGEEMISRM LH YYPQSGNGRDLTHLQQSLYSRLQRDVEQLESSIASRERQLTRAQQELRRISAELEAN AALPPEIDRLCGDL EKLKGDIKQQRSA SEQQEKEIDAKQSRRNALQSEIRQVAETQQLDAQVKEQGLSKAEVEKIRSEIHLREK VALRSKDVEGQQQLLAGVEDEINRRGEELRVTT C NSLLQRCM QDTETAKFASDTAWASLSRFATSATL SGDPSSSFSSRGSSASRRSSSRGWAPQSHSETEAKGNADEQLATHEQLLGPWKA WKAELLRLVQSD KQVDAQSRRTDEKTIGEIEKLKKARQQLDSTRRTEERKISVLHEDLLKLEEQQKQVLSSEIETLQRGV S EHRLAVEAELLKAERQANDLRLLYEKTEKENDAEVAAKIDS LKKIQQDGLAGKREVLHALQQLVQNKEHAC VEYEKDLLAVVARSEKYTLGC

>NDC80_Neospora_caninum [721]

MHDPSNLRGLTSGASSLGPSAFLPLSSPLRSTSPYRADRLAPAPSASSSFPGFSSPMRSAGGGDGRSRG
FPGQAGSAATVGRGLGSKKGPQGVQQLLGNEEAKRESVKTIIIRFLCYSGFPQQQLSPKVFVAPPRNLLVEIW
NHLLRRACDDSVQVTNENANEEVPRLFKEGLSWLLFLCFLRGPSRSRLLGLSFDCCPSSYPLTIKSSMQ
APNSAHQWPLHLHALSWLCELLIYESEEVFSRDPLLNPAVEKKDAMAGAVMISRMLLNHYPQSGNGRDLTQ
LQQVLYSQMEREVEQLESIASRERHNLRAQQELRSISAELEANAALPPEIDRLCADLEKLNDGIKQQKTAS
DQQDKEIDAKQKRRSALQTEIRQVAEETEHLDQVKEQGISKAEVKIRSEIHALREKVALRSKDVEAQKLQ
LTSVEDEINRRGEELRITTRTGNTLLQRCMQDTESAKFPSDAAWASLDRFATTASLPADSDSPSSSLGASA
RSRSDGRGRSASQLPLADGRERGGRISVTSATDSPLATHEQLLGVWRWKAWKAELLRLVDSDRHFDAQSR
RTDEKTAEEIEKLKKTRQQLESARRTEERKISVLREDLLKVEAQRTQQHQVLSAEIETLERGVSEHRQAVEA
ALLKAERHANDLRLLFETREKENAEIAEKDDLRIQEDGLSEKRRILHALQDLVQKKEQVCVEYERDLVA
AVARSEKYTQSC

>NDC80_Sarcocystis_neurona [689]

MEKGKRDLCRECSFYKSLLSSDPKSNYYHHGSVGGGHGSSSVGRSSKKGSAAAGGSAGVSSSS
ASSSSNVGVQALSNEELRRACSRТИAQFLAYSGFPQAQLTPRQLFAAPPRNVLLDIWNHLLRAYDDSVIT
PENANEEVPRLFKEGYPLTIKSSMQAPNSGHQWPLHVHAELWLCELLIYESEEVFSRTPALAPPGVHTPP
VAAAFEDLLPGTSADDVVCQIAVEQYAQLQTRDVAALQENAEIRLRKDVQIMEAAAATRAEQLAVATQET
QDVTAELEANAALLPEINHLTSECQKLQEAVQQQRRIAIQVEKAIAAKEEEKKNLQTELQAVAAEAQLEQRV
RAQGMSRAEMEKITETACLREGVSLRSKEIDTQQQLLARLEEQKNQHAEELRVTTSCNALLQHCTQHA
ENVKFDSQPGWASLARFSTVSSLISL SAGRSPRTSRTSSTAAGAGAPPTAAASEEEGEGAAAATAATL
EKLLGFDWKPWAKSLQKLMQADHQSAAQCLKDDERVRAETEKVKKNYQQLVSMRKTEERKLLVLREDLD
AAAAAQREQQQQRSQAQEHQGLEQQQLDELRCNVEGAHAAQQQLQQAQQLAARQHDAQQLREGLTTL
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>NDC80_Besnoitia_besnoiti [729]

MHDSLHLSAGGPSSALPPSSPLRATSPYRIDRSAPFPSSGAAFVPPFSFSSVASPVRFAAGAEGAERGF
GQTVRGGYAGNPDVGDARNSFFDATGGTSMIAAGGSAREKVRREVQCSFYKSLLPAGAAAVGRGGK
KGPGGVQQLLGNEEAKRESVKTIIIRFLCYSGFPQQQLSPKVFAPPRLSLLVEIWSHLLRKACDDSVQVTNEN
ANEEVPRLFKEGYPLTIKSSMQAPNSAHQWPLHLHALSWLCELLIYESEEVFSRDSILNPSLDRHEDRMR
SGILVGEEMISRMLLHHYAHSGGGRDLAPLQQTYAKIEGEVEHLEKSISARERQLTRAQQELRSISAELEA
NAALPPEDRLRADLEKLNEGLKQQTTAFAEQEKEVESQQKRRSAMQSEIQVAAETKRLEAQVKEQGIS
KAEVEKIRGDIHALRERVALRAKDVAQQQLQTGLEDEINQHGEELRITTRTCNSLLQRCAQDAETAKFASQ
AAWLSLTKFSTSASLSLDSPSPSRGAGSASLDGGGSDAAGLAHERLLGVWRWKAWKADLQLRLVES
DSSSDAQSRREDERVAAEIEKLKKTRQQLESSRTEERKISVLHEDLQKVEEQGAQQRQVLSEEIEALQRG
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RACAELYERELLNVVARSEKYTQKI

>NDC80_Cystoisospora_suis [134]

MQAPNSGHQWPLHLHALCWLCCELLIYESEEVFNRDPLNPLSQLQSSHTLSSPLSSSSSSAAAANGRADSLA
GGVSGEEKISRLLNYYAQTGHGRDTNQIQTICKSNLFFFSSFSSSSSSLLLLFFFSSSS

>NDC80_Eimeria_acervulina [592]

MQEEKKVMKLVGGTVVGLLVPSKGTYKLLASAASAHNLESQQPREMSDFSYKTVNTTDKSLHSTYRRGK
RSAEATKLKSLQFLSFEPGPLSAKALSSPQRSLLIDIWGHIFRRCFDDSVKITAENVNDEVPKLFKEIGYP
FPIKSAMLAPGTGHQWPHHLAALSWLCELLIYEENEFFFQEEEADMQAARGDVVGRMFAAAYPQLGKDP
QALRAAVLGQLHGMNREAETQQQLQQQLQQRLAAVEAEIRDNASLMEAAEERDSLLLQQK
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NECLSSMSQLAVSAKLPTAEDWSALPKMETAVAAAPAAAACDIKRATNEILGLDWQELKKGMQKIMQQE
VTAKEETHQQQRQRGIREIEKTKKAIKDLLANRKTLEKRMQLVQDDLQQQLQQQAEQQQLVEAERQHLEG
VRETKQQYQQQQQLQRLVQQQQQLLEQEERQQQQAAAAMEALQQQQQALQQKELLLLQLRSLTAT
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>NDC80_Eimeria_brunetti [564]

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FYQEEEAAEMQAARGDVVGRMVAAYPQLGKDPQALKAAVLGQLHAMNREAETQQQLQQQLQQRQQ
QLAAVEAELENASLMESLATATEAERDSLLLQQKAVLEEVQKLGDRINKAAGQGIDKAEVERIRHDVSRLREG
SHRTKEIESLSEGLQTLIDKVADDSEKLRIAGQSVNECLSSMCQLAVSAKLPTAEEWSALPKMETASAAAAA
APAAAAAAAAACDMKRATNEILGLDWQQLKKGILQIMQLEGAKEETQQQRQRAAREIEKLKKVIKDLLAS
RKTLEKRMQLVQEDLQQQLQQQQQLLETERRHLEGMRRETQQYQQQQQLQHLVQQQQQLDQ
EERQHQQQAAAAIEALQQQQQEAQQKQMLQQQLRFIADEKVALEQQQLAELAEALGPPPPEEEGSS
S

>NDC80_Eimeria_falciformis [627]

MEPLIGGPTSPRLNTPPLHPDSQQQRELSEFSFYKAMNTGDCKALQSTYRRGKRSTEATKLKSLQNQDS
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ISKSAMLAPGTAHQWPHHLAALSWLCELLIYENEFFYQEQQPRELSEFSFYKAVNPADKALHSTYRRRKRSAEATKLKSLQTQD
RNLVLGQLHAMNAEAAQQVQQQLQQRQQQLAVVQAELRGNASLVESIEQLTRQEDATRASIAETKEE
ILHQESRAAAAAGERDLLQQQQKGLLDEVQQLEERVKAQGIDRAEVERIRHDVSRLREILAHRTKEINGLSE
ALQTLVDQVADESEKLRMASDSVNECVGISSQLALAVKLPTAEAWASLPKMETFAAETADSPAAAAAAAND
VKSATNKILGVDWPALKEMRLIQLGEVQAETHQQQRQRAEREIEKLKRAIKDLVASRKTLEKRMQLVRED
LQQQQQQHQAEQQQLEETERQQLEGIREVKQRYQQEQQQQLKVKQKQELLQQERQQQEAASAME
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>NDC80_Eimeria_mitis [529]

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WPHHLAALSWLCELLIYENEFFYQEEEAAEMQCLGSRRELLIYENEFFYQEEEAAEMQAARGDIVGRMFAA
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KDISSLCRCIADTRAEISQESLAAATEAERDSLQQKAVAEEVQKLGERIKAQGIDKAEVERIRHDVSRLR
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>NDC80_Eimeria_maxima [575]

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QQPSTQEQQS

>NDC80_Eimeria_necatrix [646]

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PFPIKSAMLAPGTGHQWPHHLAALSWLCELLIYENEFFFQEEETSECSGPADVVGKLVAAAYPQLGKDPQ
ALRTAVLGQLQAMKRDAETAQQQMQQQLQQQQQLAGIETELRDNASLMDTIEQLRREADSLRRQIAETR
TEISNEESVAAATEAERDSLQQQKAALEEVKRLEERIKAQGIDKAEVERIRHDVGRREGVSHRAKEIEVLS
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AGSGGSAAAERAAAAVAAAACDVKRATKEILGVWDTELKEMLRVMRLEEEAKEETQQQRQRQLGKEIEK
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>NDC80_Eimeria_praecox [469]

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FFYQEEEAAEMQAARGDLVGRMFAAYPQLGKDPQALRAAVLGQLHAMNREAATQQQLQQQLQQRQQ
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RVKAQGIDKAEVERIRHDVSRLREGLSRSKVADDSEKLRAGQSNRKTLEKRMQLVQEDLQQQLQQQQD
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>NDC80_Eimeria_tenella [645]

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PFPIKSAMLAPGTGHQWPHHLAALSWLCELLIYENEFFFQEEETSECSGPADVVGKLVAAAYPQLGKDPQ
ALRTAVLGQLQAMKRDAETAQQQMQQQLQQQQQLAGIETELRDNASLMDTIEQLRREADSLRRQIAETR
SEISNEESVAAATEAERDSLQQQKAALKEVKRLEERVKAQGIDKAEVERIRHDVRLRDGVSHRAKEIEV
SEGLQDLVDKIADDSEKLRAGQSVNDALSNMAQLAVSGKLQTAEAWSALPKMETVPAAAAAAAAAGGAAAA
AGSGGSAAAERAVAAVAAAACDVKRATKEILGVWDTELKEMLRVMRLEEEAKEETQQQRQRQLGKEIEK
LKKAIKDLLASRKTLERKRMQLVQEDLQQQLQQQQNEQQQLQEAEREHLEGVREAKQQHQQQQQLQRLV
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PSPNQEEPPQ

>NDC80_Cyclospora_cayetanensis [481]

MEPLIGGPTSPFRANAISAASSILPESQQPRELSEFSFYKAVNPADKALHSTYRRRKRSAEATKLKSLQTQD
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DIVIVRRFLSFSEFPGPLSAKALSSPQRSLLIDIWGHLFRRCFDDSVKITAENVNDEVPRLFKEIGYPFPISKS

AM LAPGTGHQWP HHLA AL SWLCELLI YEN EFFF QEEE IAD FYEGH AAAA AER DLLLQQ QKG VLA EVR QLE
ERI KA QG QID KAE VE QIR HDIS RL REG LA HRT KEIE SLNEA VQ ALD QV NEC ISSM SQLA VSK LPT AE AWN AL
PR MTP AAGG AAAA ADRIA AAAA ACDV KRAANG IPLL GREV QGE AL VR LL VPQL GLD WAS MKKE MLR
LMQ LEEAAKEETHQQRQRAEREIDKLKAMKDLLAVSAVGTRDFTVYGVS

>NDC80_Eleutheroschizon_duboscqi [633]

MRAYGDSRSPYPSQGGPQRGGEGFSLYERGRARPKVSVQSMSFWKPANSQDTAGTAGGGGTRKLLDA
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LFRDIGCPMQLSKSSMPAPGAGHQWAHHLYALSWLCELLIFETEAFFPDLSLDAEIVSQRWKAGLQHQHF
SDAELNRGGAIRRPPTSGPDRGLASIIANAFTQLQEGKDAQATRGNILKQIETIDRSRQTLKQRRAFREA
TDALETANSECQANAKLAPDLAAMEADRQKFELQHHHEHKIASLTDDIQEIEANKKELEEQLRTVTAECQ
LQHVIQNQGICKDEMVKITHDLNIANRQLQTIAELEKQNNDNALAHDKINNLDNQIITSRAVNCLISGCRTA
ATRLPTSKVWAELKTFISRDSAADLNEMAGLDWKTWTAVKKNVQDDAAELNKAEVRLAQENKAHVDNVK
DIQKLKRNKSLEQTKLDTLQMDVDAENTQHEDRNNNLSQRQIEIRTNIAGLQDDLEWQLGKEKSALQECSA
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>NDC80_Cryptosporidium_andersoni [686]

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SLSSPTRTLYLEVLFQIINQYDPKIKL SRPDEDIPRFFKDIGYPFTINKTTIAPGAPNTWPQHIAAMSWLCELL
DYEQATFPYLNFSKEGELNFV PDLTGTSTLANAFTGTAGSMNADHMSIDS AKLLSQTLNKRLSESYKLF LG
GQSDGGLLAETLEAYCRERKGQAQQAFDRKKQDLEQM QMEMQKIQHELQEGD LFLQKNHTLSQDILKME
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RSDREKAENWALELQIEELSTKLYQLSRLWNNKYQEIIITSNEIRNHEIGSFNSLQNLSMSILNSDELNN
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VCDKREMRQLEQI QE KVKKYKTEFEQKKDEALKRLETARSRAVAKLEQQLVLAENHAKNELQQCKQVINSD
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>NDC80_Cryptosporidium_muris [686]

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SDREKAENWALELQIEELSTKLYQLSRLWNNKYQEIIITSNEIRNHEIGSFNSLQNLSMSILSSDELNKET
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>NDC80_Cryptosporidium_hominis [696]

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QDLLKMEHAVGQCQNSIRQMENSCNDTENILNIKKKELRDIEDEIEKLQKRIANQGIQRDDVSRVYQDMKVK
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SGSNDKSQGV DGIIGQNWNSNMKH SCKRCIVEVEMWKRKLKTQEVFSEIMKSTKDTI IQKKQACDALEKKI
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>NDC80_Cryptosporidium_meleagridis [695]

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TLNNPTRALYLEVLFQIISQYDPRIKISKPDDEIPRFFKDIGYPITINKTTIAPGAPNTWPQHIAAMTWLCELLD
YEQAVFPHLNYSKDGEFGGYMADTSLTSTLSNIFQHNTSMVNGFPNFE GNNNLDSSKLLSQTLNRRLS
SYKLFLNGQNDGGLLNETLEAYCRERKNQAQQALEKKKHELEQM NMEI QKINHELQEGE LLLQRNHTLSQ
DLLKMEHAVGQCQNSIRQMENSCNDTENILNIKKKELRDIEDEIEKLQKRIANQGIQRDDVSRVYQDMKVK
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GSNDKSQGV DGIIGQNWNSNMKH SCKRCIVEVEMWKRKLKTQEVFSEIMKSTKDTI IQKKQACDALEKKI
NLDGDL SVC D KREMRQLEQI QE KVKS YKAE FERN KDEALKRLEKA KSRII QLEQ QSNLQESHSRNEIIQCKQ
VIQSDAEAI FK LKEYI VQNL SNSL NSIANLKNQV KLRNN SKLIEVEK WKK

>NDC80_Cryptosporidium_parvum [696]

MSSNAFRPFGQNSMRRTGFRDNESEGDLSQDVMTSFSIGQNKTQSPTHFATSSLLKGGSNNINNTSNAWAYSSSRGGNSQRGANAVGTSSMTGNMSSAAASSSKGRSISDSRREGLKILTFLSKRGYPNQINPKTLNSPTRALYLEVLQFIISQYDPRIKISKPDDEIPRFFKDGYPITINKTIIAPGAPNTWPQHIAAMTWCELLDYEQAVFPHLNYSKDGEFGGYMADTSLTSTLSNIFQHANSNVANGFPNFEGBNNNLDSKLLSQTLNRRLSESYKLFNGQNDGGLNETLEAYCRERKNQAQQALEKKKHELEQMNMIEQKINHELQEGERLLQRNHTLSQDLLKMEHAVGQCQNSIRQMENSCNDTENILNIKKELRDIEDEIEKLQKRIANQGIQRDDVSRYQDMKVKRQTIRNLRTERENTEKENWEIELSIEEHSTNLYNLSRAWNSIYQEITNITSNDIRTNEVCSFNLLQPLSLNISYSGSNDKSQGVTDIIGQNWSNMKHSCRCIVEVEMWKRKLKTQEVQFSEIMKSTKDTIQQKQACDALEKKIKNLDGDLSCDKREMRQLEQIQEKFVKSYKAEFERNKDEALKRLEKAKSRIIQLEQQSNLQESHSRNEIIQCQVIQSDAEAIFKLKEYIIQNLNSLNSISNLKNQVKLRNNSKLIEVEKWKK

>NDC80_Lecudina_tuzetae [261]

VSTQSTWGEMNTPSRSQGGRLHGGCSHRSQDGGRSAPRLTASGASPVRSGYEARWTASRLHGHQDQRTPLVVSGPMSTVAMVSKNSLSRKHMHDPSRNEAIRTVLRYLTTRGFPRQVSAQELQSPAKALYVSMEF LIQVDNQSRIKDPEEDICRFFKDVAACPLTITKSAIKAPGAPTTWPTLLMAMSWLCELLIWDEMVCQPMMDGAHSALDIPEEQRTFDQQIQVCYRRMLEENLDEAGAREVLERQREKDHAAR

>NDC80_Pterospora_schizosoma [645]

YQRRVHGTSQSSGQVPHTFDMNIIDGVRPATAPSRRHSRYQNESGIVTSYIGVPSRAVAPLIDCLSYKAENIQIVMSFISKFPDFMGRHGLNEQSIHSPTKTVVAIMDFLMHQFDASIIMQIEEDVPKFYRDMGCPVLITKSYVTAPGTPSTWPTLLMAMAWLCELLRYEQSISNHSCPIGAVSEQNTSFNHVILKAYQLKTRCSTQKELNAQQILEQYCESSESMTSTLNEIKSKLADVRQKTRDCRDQLENADTTALRNEDLEEDEKKLKIAYAKCCSNYNECGANKMSTDLYEKKKMLDNIEKSNAELTDMIKAQNICVKDIVDMETEVSTQQQRLEDVTSKYDNVDNKILQ LLEKNKKQCQGITVKLNSDLIPCTTNQKIIESTEVGDTCMHVIMEVERALHKWRSYYTSLEDKNVYESCHNRLRSSAADIPEDLLKPVAANIENQIVHFEQEQQRIDEQVMKYKKNIDDSLRRLKEKQREIKNIEASINSKKHEIESAVAFHERECSEIKNDTQEKKQKMHSCLGDDIANKVKHATTCKDLKIRLEVAQKEHYKNLKNLNHVVVKMTSTAMDQCQRECESKLESFAVSSSKTCKSYYQVCSEAYRLLENVSSPPAINKEQCCKHITHATDACIK

>NDC80_Selenidium_pygospionis [742]

MHSFQPQARTPVRQSNSTQRSAGASSLLGFNTPGTGRRGLLRQSHRSASRERPATVDRGGGGDSAYTS PRLHGGPCTFGSTDGAMSAPRKPTNAMVAQIPVGAMSVGRHFVGSHTGQSGLGGFTKSSEARREGAE PSKKESLKNVVRYLTNHNYPSPVTAKSLSSPTRAAYLDILMFLIARVDPLVPSTFTKVEEDVPRFFKDMGYP ERINKTSMIAPGAPNTWPQHVAAMSWLCDILLYEEEVFPSLCECWGTGPGTPSGTPNGRTGSEIVEGTDFS NGQGGHWEGVTRAIADAYRVYMLTPADEATSVKEEAANDELIEGLIEAKKAQFERAQRKKQSELDNIKNQIIDLERQTKTYDDLSESIKDARVQLTELETMSRVQHELKQKISLVEQKTAQVEALNDINVSTSRCDELENTIS RQGINRVEVRQIDSQDLKVKKRLLSLRSERTSKEKEVYELQVQIREKIDALGGHTKAAVEHSSLATVKHY PDHPSTSHSSTSNAASQRSSVASSSPELRGIRPLEMDLSMITAHLTSIEDTKEAREERRFTVDDVFKVKWK QHKEQLTRCINKEQSAQFQLEREVASVSESCAVVEKDISKRANNVELQRQVESNEEARLFESQEEESIK RVEEQVKAVSLDIASYKSNVQDKLANCQVSKGSIESEIQEKEALLERILLEARTQFFRCAKSLVELKEFTESGFAGAVKQCEAISKSLPEPPRSDQGLSDSVIE

>NDC80_Colpodella_sp [85]

SGAGVRKDTRPLTEKKWQETNLKNLIQFLISRGPYHSLTSNSLKKPQMVEFASLMKFLVTEFDPDFDFGTRNFNEDVEKFIKAIGYP RNFNEDVEKFIKA

>NDC80_Colpodella_angusta [66]

KNLIQFLISRGPYHSLTSNSLKKPQMVEFASLMKFLVTEFDPDFDFGTRNFNEDVEKFIKAIGYP

>NDC80_Colpoda_aspera [616]

MSTRYRASIGGMSFGGKLPTDMNHSRSTSSNTRQTIIGYNPKHPQRSASTGRESLGAGGLRADLKPVSPFSKPKRKLESKIHKDKLAGLINNIFEFLSSHNYEAAISKKGLGSGNRNIFLSMLGFIISFLDRKFAADIMVWN EDEVMKLLKFIGYPFSFPKNFFAAIGAPSWNQCLHVLWSWIVELIKYFEFVKTVKKKEPAEKTDSDLNDYVQKKFVKGDLMENEGQNELTEKLIKRNQHLQQLCNDYKLKIEIENIEMMQNTLHDYPDPSALQNKISHLKDEAFNIEQQIARQKNENSPLYESVNRRRHLENLNTDLDRLKAEKQHLSAMIQNQPLTIEESERLNKEIQCENNRSKLRQTQKEKMKQRDVLMQSKEELVNELESRAKEFNNTIQSEREIFTKCSGFQIKLNENALMDPRADDQI DSDEQVFIKHLGCGISPKEEVRPVLREILDATNERVRNMEEEVTNQKFLCAQLGDQKQELAAKIDEQRFRKLKVTEIHDRVEALSREHVNFTEKREKIQKEIDLKREKVEVESENGVLTRYNTLTSELEQLQETEANRSGITNMLANDVERALVHKKEIIESLRDIDLKFGKIKESFEDFLNQ

>NDC80_Chromera_veila [733]

MESPLMLHRNVHTANLRFVTMGVGNRDKDPRPLANAGWQSRAITSLVDFLVNHDFPHQINQNSLKRPT RTVFMVVAKHLISLYDPAWHWSDKPDEDVAQFFKMAGYPYEIHKSIAAVGAPNTWPPLLGALSWLVDLL QYGESVGITAQLAEKSGLSLISSVPEYDEKAILARTLREDAIETWDLLERGISSPEHDRKLTYLKSHPVRSQQLNIEYVRQERECCENDIAEFERKCRQASEEQREHSKVRKEYEEMKRREEERLRDARGPLAERSALQERV VRLSKERDAKRRECESLRRERVNSQPMREDFESTRRGIEKTRVLAQREAELAEVDRDAAAEREVIRDIGFELDAIAREVRAADDDELVQTAQSSKGFFIDKPEAHPRHLLDCPARICRDLEGGAEDAETEGLGEGDVNSVLGRWKDQVKPLLKDADLLVFTKHRENLKIHEELQSDRLQLEREEQALTYEKDRLARRVAETQAEREREETAA

QEIEKNFAEAEAREAALERDIAEEGKVQDLKEQEKMERTLRKAQEDGEAARATAQSRVNAMSDGLAQL
RQEERQRFAATEAARAVREKLRSSFEENLVPLLENLSARVADLAERAGVNLDSDSSMQEADEGAAAERE
GHDAGRDSAQVSSGPSGDSPDCDMGCEQSASASASSSARPENSGETEGDGANGCCFFFEPVDDDV
GVKGTTTEEVTEGQVCAEEGEEEQTGDCLV

>NDC80 Piridum sociabile [386]

MADENTGTTTPRRRPTAVGPSRVISARQPGMATTCLRPINQTHCPMVTPQPGRGAANNVPVSSTQLRSSA
MPSRATNNRPVVGKMSNAFCHQNGSKKASDPRDVSSRVFQLECFKDVT AFLQTYDYPHTISIKQFSSPTR
NDVLNITSFIFSIDPNWTMQKGLEVDPQFLKFICKYPTPIRPA LLNTGAPSTS IIIAAIQWLSDLAI FVTQHH
GTICGTNGGATADCNSTDQSEA QVSPLMLAQQALCNFTREAYSLW LQGEDEFQEQT SRLSSDLARANDILI
SRTNEYRQESKNLEASKAGLLEDIDQEGILQDSIRRAQTDVEKLQRELRLRDTKKHKGWPGYADKNQR
DLLECQNHVQSLLDEERRLRDKIANQDVT

>NDC80_Vitrella_Brassicaformis [673]

MDSHTPRRGLSIPPNSRSAASMRAPSRTPVPSAYRPLVGKPSLTGKASLGGKAGALVKGKRGAGLKV
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YRYDKSIEVDISSLFFRYVRYPNPISKTALLNVGAPNTWPSLLAALQWLIELCQYAEQLDDREDPFGVSDVEE
EGAHSSEASPLSPSSSLTSSLREQWVCLKETYALYMAGEDHYPEQDRLKAHYEQMESALQTRLQNA
RDRSAEQAASKLGYQEDLQKEDTLKASIARKQTDMEKLSREIRRLQENRQKKEGPWGEVARAQREVEAV
RQIVQESEAEEKRLKETVAAQEISASDVEHLLSEQRRYETKYKADRDDKAKLDHSVMQMEDIKLHQRQSSL
SSNSRKLQSLMSAIRHDGQLPSTDPTIGSFLALPTPCLMLQCDALQPAALEQHEKDKEWTDVVDSVLGV
DSQRFRDQLEGLIEQEAQHHSKLSAEADELQRAIDAMTKTIKQNQQAKDRSHCRYERHVDELQRLQKQHE
EEVNNINKETQATIKAKEQADEERSLRQIEEKELLRDQITAQREEWQQRAKTACDIKESVRMVDAAQP
PVVKALNDGFQALVDVKKHLSQEAAMPLRMPTFPTDEEPQ

>NDC80_Platyproteinum_vivax [363]

MSNPSNKEYMSIIHFLTLLDKHFSPVKKLEDEIPPFFKQIGYPFMITKTNLIAPGAPNTWPNLLGAASFVVEL
LQYDLERENLNEDEEHKDNIIINNQIADGYMAFMTGSDDIEKHDAISIADHFVSRNQEIQKECAMLQETIQRY
EEQIEVFKSDQYSLPALQQSLQVREEELLHAQHEAKKMQEAVCQKEDVIKDKEKVIEDQKTERSEVEQEIE
RLKIQVEKQEISREELQKMLSDRARLSDELDVLRNEKNGAQKELLDTESSLHHKGEDLQRMGRQINQEASR
ASATVLSQPSWDFEHLPTIDLNGRNIEALLGGNVSKLELLNDIRDAKKRLNEAHDQEGWSLQRQLQLLT
DEE

>NDC80_Digyalum_oweni [609]

LTMPQSIQGGSSRRSLGPPQRPTAGPRTDSMANRPSLNSSSLGRTLGVGGLNPGRGKMSDSRPLKDR
EYMQAQTRKLLDFLVSSNYPHPIAPKAMVNPSKFEFLSILTFLIMLLDPHYLPLKKIEDEIPSTFKQFGYPWMI
TKTNLIAPGAPNTWPNLLGAASFVVDLLIYDQERQIEGSSAYEGDNERKDTMFVNEKLASGYITWMTGNDDI
TCHDTSVVTYFHQRNEEIKNSIGQLQETLNQQEQQLLTFGGNEYSLPSLEVKLHERNETLKAKQHENESQA
SVLKHWEEKIRDKNNIIQEQLNEKLPLQNELEKLKHQLDSQEISREELQRMLTDRQLQNELERLRDEKDNL
NSRLWDLKDKSGSYEEILHKLARQISQDAAKLGKIPKSRTDTESGSIDLPKIDLTARKNIKQLLGGNMEVTID
ALEAQLKTGEALQQAKDEEWKGSEISMLETQQKQLIESIEWKTKRESAMKNEDALTQERFNKQLSEM
RNHLEEAEKGTEKNLAFFEAWLKKENQSQUELVLEEEESRLLIESERQQADKIVARVSDIVLKCQRKNNERI
HHSEVAMQSYVDQLQSNPLRLPAAPLTVLQLSHTIN

>NDC80_Perkinsus_marinus [634]

MEPTTPARSEKAPRRVTINPMSAPKTTRKKSVGQRQQQQPGSGAKSGAVRDLYQTALPTSHTGLQA
SLSPKPVQATLLDPPRSHDGTVRLKDERPLQSKGYRKACAEYVAYLSKHGYRSDRMKPGWFDSPSTRE
FFSLCEFLLERIDPNLIVSAPQKKSTPEDLISIVRTKLLYPDVPKSALASVGAIITWPALLGLLSWLVEEADE
ADMRERINSVSDTCRAREKALGTAYTSVQAGQAETLEETAMLDKIEGLTRAQYEELRSENAKCLKIJD
NEAAIEKIKESGGDDPEQYERKNEEIRGCVAEKEREVGEIIRRREKEAAERQKWEERTAREKEKEKKLR
DRIDELSQRVDAQTYSHEDLAQLLARSKRLDEEARHLQEIQIARCDSDTNAEDAQLTEQDTLQRLCHQFN
ECIGAGGHPAIPRANFNPATEFLKRGFGSLAAPGLNIESLVDARVEDTRKQLNKAEDALREYVESREQAD
KQLRKAKKEIKTVAMKEAEWKETREALLKERADSEKKAVADAEARLVELDKQSCHKLHESSKAQKVVKIEDT
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>NDC80_Symbiodinium_microadriaticum [597]

MASTPKQKFRPSLGEPSRVSLRAQSPGASSVGSGEDIGTSPRRRALFPSGTGRTRALVSSRHSRDPRPV
GDRAFTAQCAKNVVELLSARGYSKTFSHDKLLKDPSTKEFYDVFRFLIAQLDPQLEVEGKMEDEVPSIMRR
LKYPVEVNRSKLQAISGPNTWPQLLAVLDWLTVLVRINDELIEPLAACQLGLSDVGDPDRDAGDHHLRLSL
HENYLNFLSGKDDSGDEERLRMIMQRLAEALQSEIQRLENQQVEMQQQLQDFQGEHDRLL ELQKMPAQL
EVEADRLRSIIQAQEHKVQTIEDEMLCKEAEEEREQQKEIETLQAKRAYLAEQVESQAYSRRDIERLHAERSH
LRQLFKDLKAEGERADEEVWELGMQEKSRAEEIGRLVRQVNDTVECLTHALTEEEADIFDFRVRVDLSEPS
DALASLAFDDQKTWAQAASSSHGERLQQAELALHELLEEQRGLQTSLAKEGEARHMRARHEQLTRIYQE
YREWSASQIDDAQKTTEATEDAVHEISIGSAAPSLRDAAEVDKLKLTASVQTQGAHERAQLEEQIRRDEE
RLEEHHRQLIMRELDSYAKASTALCEDVEAALL

>NDC80 *Tetrahymena thermophila* [671]

MRHTEFFRGPNDTSDGSSKASQSQKNSSNNNTHLFFKEGNYNANYNNSDKLNQSMNFSSIRMTNLNQPI
RNTVIQNTYKKDGNYKNNLNEKNIFILQDLLSFINLNIYQDSPISEQQLKKNPTQKIFFEIFWGFIIDPSINSKVI
SVDEIPQLMKFWNCPLQNKINTTIGAPHTLKQTQNLFWLYQLSKEVKMSSESLLDNQKSILQNSQSQSQ
RNDMEILNQYIAQSFRRKNDLLKGANFSKHNSSKNPQQNQSMLISQIGMSEDEKEAKDIADLLKQKNIYL
QQVIKELELKIELTQLSNKLNCLKQPDLSEVGKIEQQQDVINMEQELIYQLSSEIESNQKDIDNYHNIFNEKM
IAIHELEQEKSNIQKQITDQEYSIVEAEKKLKEIERLKKQKAQIDLQISQTSQRTVSYTQNEGQINEIHQLIND
YNRILNTSNEFLIQCQGIEAFVNEEKIRADNLDILTKGQLQIKSEDLKQGLDSFTNSKNNSKIIKSMINIDLKKVS
EQLAKLQDYQVSVKQEEISKQINRQIEEKTNINDLDIKINQLNIKSQQKQDELQVNFHKSEEQLQANS
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KKSFLINKS

>NDC80_Aurantiochytrium_limacinum [678]

MQSQPRRPPPLGRVSNSRLNAARGSLGGGMGSGLSGVGVAKKTRMSLGPGTLPRHSLGGGMGTAKKAK
SHKRMSMMPSMGSSLMSMQSNAQSQTSSNSRRQSSVSASRRKSIAVMGGGGSSALKSDPRPLTDKAF
RAQCETVIRFLSEHNYDQAISPKILHRPTGKDFEHIVLFLFRQVDSNRKFAGKFADEVTTMMKGLRYPFNIS
KTSЛАAVGSPHTWPSLLASLLWLVELVEYDELAGLQDDEDGDDEGEIDRIFTYIRKAYWCFLNGDDNRYE
ELDTELGNKFRERDVAIEEEIEDLERKQAEKDELETETQKQSLIPELEQRKRDFTSDKLKWDKLIRQFLENK
EIMVAKVREREAEVRAKEAELAQAEQAKAKVQIILDEQELNPADAERMTQEKRSLASCNTVLEQRDEIQK
ETWETELSITKRTKHIESLLEENYNAIAKSLKTQTVEAGQQRQGANIQLRLDTRRNVLMSLTENGDQGDRV
VDEMITSALQNLREDSIKRTHQIRTSNMELQAKVEENDEAQEAIRSEIREFEHHLRQAEANLQQDKASMEEE
LNQRASETEAVEEDIHLRQLEAEELPLRDAEATVEELRARVKSEADRHSKDIALGDDLNSAIDAFLSYKE
NIAAIADLNSSLQNQLNDANAFCESYKASIKPLCNSA

>NDC80_Bigelowiella_natans [616]

MSLRRRTLGVINHNERVSEDGNTKRRFSMAPNRRLDERKVGRGARFSVGRPSIGFMKGRKSFGRLSIAP
NGRPSIGRRSSMYNRSGKITDPRPLSDRKYINKNTRILVEYLTSHGYDKQITPKQVTAPSLKDFKNIVGFLFK
QMDPEFFFSQNLDDIRDILKIIGYPFNISKSSLSAVGSPHTWPPILAMLVWVIELLSYLEKLDAAEENQVAND
DGKIFFEFLASAYRNFLDEKDHSaedLEEGFREANEQRCAVVKDEIKRFEDENDELSKELQELSSTNTVEH
LQKKLSDHVSDLAKFEKLNQKLEKHLEVLMQKEEAAKSTLKELEGEKEQYVENQKNLKVAVEEQPLSRKEV
QMITQETKMKKDSIESLLQQKEELKRQVWDSEMLQCKEIGNVEEFVNEYNNLVYELGLKNASQATKNIDYQ
IKFDSAGNDIKEMVSKDLTHVIKKSLSHLETRAKSELRGLESELRDVKQETIRTTDLRSEAEASLNQHEQRLE
KIHEAYRTERKDMNKQCSESISAEETELHIHTRTRARATELRVSEQKIQEIEQLKVVLSEQEDIEQKTKDSI
LSSVEGLVRHKDWIVQTLTLDKATQGREGIESMCSGK

>NDC80_Arabidopsis_thaliana [568]

MRGGAAGKRRRTVGF GGAPP PPPSIEQQRHLFNSR DAS FASSRPSSIGL GGRGASDDRSSMIRFINA
FLSTHNPISIRGNPVP SVKD ISETLKFL LSALD YPCDSIKW DED LVFFL KSQ KCP FKITKSSLK APNTPH NWP
TVLAVVHWLAELARFH QHL VSN STS VP ED NS MN FF AIQS FG H FIR GE DD KV ND L DS Q FL G K LE A E K TS V AE
TIS GCE KIS GE LE AK L E SL R KG PS K KES LE KV KAD L E ND VN K F RT IV VE Y TDR N P AM E KV V E E K A K EL K A KE
EERER IS VEN KEL KKS VEL QNF SA AD VNR M R REL Q AVER DV ADA E VARD G WD Q K AWE L NS Q IRN Q FH Q I
QTLAIDCNQALRRLKLDI QFAVNERGETPAAVMGVDYKSVVKPALCSLCDGIKGSSAEKVEELVTLQHHKS
EMASKIESKRSLLGSIQLQINDLEEKMKLVKKETQELSTKCDLEAKTLVESVKA EALNLEVKEAAEFVKAS
ELRLQEAVKESEEEVQACAAQLFALIDSISKQKEYMDSKISEIKTG VADTASAVSEIYKANFKKNLGI

>NDC80_Trichomonas_vaginalis [507]

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LKLYQTGDEF P QDQ ET NYAETNFNNLLAFYHE FEEL GR DDD DG PI KERIA QT FEE QY DAAIKSCDK IED YQ
NQISFSEQKMEQM NQKAKIIHEKKS NIDT LQKEL TKL ISETEVLESQSSAM DLEM TKL QM E LRSQ NKY SEIV
KQRNSAETKVQELNFDANEVEKIKFKKEAI IKQI QT E LEK QSAT KKD IMN LRK DLE GAHE AYDRFAP YF VEN
MPPNHTLPDSNDKENRLSQAKALINALKAELS QEDPHEIEKQ LL DEQ SDIESQIAD LEKL QRS LEKE FSV S KP
KKKDESTDVLQEKLNVLNNQKKFEVERQKNIADAKAKLDEAVDALNSYKADAETAFNL LREVEMTGT I

>NDC80_Naegleria_gruberi [700]

MSFSKRSALRDITNTSGTSIHTGMHHHGGSAPLKS NLSIMNNNTTMNN SHSILN SHSLGNHLS TTSTA
TSTTTNNNNLLSTS KPPRESMIMKQTQ TLKNSK TSSSLNTS STTPNSNNI QYMLN AYEFFSQENISI EM
TPKTFREGDSSKLRYFLNYLFTHLDRNICVIVPTPSEVREMEKKNSSGNTMDARRTMNHLNSLSMN AS
MMNGMNHPSVQSSS TITTMTTINNKPPSISGLDEDL IELVDFKLPISLNKRHFTITPQSSRNWQTM IQVLY
YLTGLVKYSRERDFIERDLELQEVSIFFSFLSDLYAPFLENNQREMDRLVNDWRNKF NM DCTRSRENLTQ
KNLHESIKNRLNYF NTRQDPFHELEEKRIVLESQFLNLNESSRSNEEKLNSLLHFREITKAKIVSTNQQIDSKK
RDNEVCARQLEHQEITPLEAQALYREKSLYEEEERILKRTDYTRNQVENAQRHLSQCNEQITLTSQYNSI
VLKIDPNLRLSSNNAKWRIDDSKQKKTILQQFTTQLSQIQQLNETLAQKDEMIQVLNEHLQDVKESLTQ
ETNSFNNLESLLVQIKQQHQEEELFEYDRIKTIEDQSVQIHKSEKRNWKQEFENTELLIAQLEEMQREEQEQ
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>NDC80_Dictyostelium_discoideum [855]

MHNNNLTYQQYQQQQQLIQQQQQQQLLYQQQLLQQQQQQQQQQKQYSHGRPSTLGLPSRKPPPMQ
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PSKSNTISSNTTTAASSSSSNPSITLRQETRPVNDSKFQSQCRLRLQQFLSQSEKGYPRAITKDNLMS
EINGLMKFCLCHIDPHFKLPTEKPEEEIISFFKMMGYRSAGFTKRVFSQVGNAHTRGHLLGAIWMVELIEYD
QIIRQMDSHIEREILNGNTIDENGNLTTTSATNTNEDGNFNLTQIFSKHVFKSYEAFLKGDDGNHVDFQFLE
EKFNEVNELIVKECEESMERQKKIKQEIVNLTQYITNDDIDQLNFKLEQTNQEIIATHDLINDNELEDQKLDQEIDQNQTHFQELEIQQKEHQKKQEFESRIQIQQSNSVDAKRINQETFELKEELDKVQREQSRHDQLKQEKSKEIKQSINEIEKLIQSYKEICCKASLVTNDNLIIEKYRIIFTHTLNNNNNNNTNNSILNISNNIINNNNNNNNNNN
NNYEGIISISDLKLVIKNLQELNLEISQNIKIKQDLLKTSNLNDIQKESISDKEELARIKQKHDQKKNILKQEKDQDTFDITQREENIEQLKSSNKKLDDGVRELEKTIQLLESQREEHSRLKIDLKQSSEQMKLQYLKRIERFISHRQLIGSQIFDTLQMEELSLTQKILFPPPPPPPQLSQHQQLMQQQHQNNNTIDIDE

>NDC80_Saccharomyces_cerevisiae [691]

MQSSTSTDQHVLHHMDPHRFTSQIPTATSSQLRRRNSTNQGLTDMINKSIARNTISGTGIPGGINKNKRT
STVAGGTNGTALALNDKSNSRNSVSRLSINQLGSLQQHLSNRDPRPLRDKNFQSAIQEEIYDYLKKNKFDIE
TNHPISIKFLKQPTQKGFIIFWKLYLRLDPGYGFTKSIENEIYQILKNLRYPFLESINKSQISAVGGSNWHKFL
GMLHWVMVRTNIKLDMCLNKVDRSLINQNTQEITILSQPLKTLDEQDQRQERYELMVEKLLIDYFTESYKSFL
KLEDNYEPMSMQELKLGFEKFVHIINTDIANLQTQNDNLYEKYQEVMKISQKIKTTREKWKALKSDSNKYENY
VNAMKQKSQEWPGBKLEKMKSECELKEEEEIKALQSNISELHKILRKKGISTEQFELQNQEREKL TRELDKINIQ
SDKLTSSIKSRKLEAEGIFKSLLDTLRQYDSSIQLNRLRSRSQQLGHNVNDSSLKINISENLLDRDFHEGISYEQL
FPKGSGINESIKKSILKLNEIQRERIKTIEKDNTITLEKDIKNLKH DINEKTQINEKLEL ELSEANSKFELSKQENE
RLLVAQRRIEIKMEKKINDSNLLMKTKISDAEELVTSTELKLEELKVDLNRKRYKLHQQVIHVIDITSKFKINIQS
SLENSENELGNVIEELRNLEFETEHNVTN

>NDC80_Homo_sapiens [642]

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PDTKFEEEVPRIFKD LGYPFALS KSSM YTVGAPHTWPHIVAALVWLIDCIKIHTAMKESSPLFDDGQPWGEE
TEDGIMHNKLFLDYTIKC YESF MSGAD SFDEMNA ELQSKL KDLF NVDA FKLES LEAKN RALNE QIAR LEQER
EKEPNRLES LRKL KASL QGDV QKYQ AYMS NLE SHSAIL DQKL NGLN EEIAR VELE C ETIK QEN TRLQ NII DN
QKYSVADIERINHERNELQQQTINKLT KDLAE QQKL WNEEL KYARG KEAI ETQLA EYH KLARK LK LIPKG AEN
SKGYDFEIKFNPEAGANCLVKYRAQVYVPLKELLNETEEEINKALNKKMGLEDTLEQLNAMITESKRSVRTL
KEEVQKLDLYQQKIKEAEEDEKCASELESLEKHKHLLESTVNQGLSEAMNELDAVQREYQLVQTTTEE
RRKVGNNLQRLEMVATHVGSVEKHLEEQIAKV DREYE ECMSEDLSEN IKEIRD KYEKKA TLIK SSEE