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## Secretory organelles of pathogenic protozoa

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### ABSTRACT

Secretory processes play an important role on the biology and life cycles of parasitic protozoa. This review focus on basic aspects, from a cell biology perspective, of the secretion of (a) micronemes, rhoptries and dense granules in members of the Apicomplexa group, where these organelles are involved in the process of protozoan penetration into the host cell, survival within the parasitophorous vacuole and subsequent egress from the host cell, (b) the Maurer's cleft in *Plasmodium*, a structure involved in the secretion of proteins synthesized by the intravacuolar parasite and transported through vesicles to the erythrocyte surface, (c) the secretion of macromolecules into the flagellar pocket of trypanosomatids, and (d) the secretion of proteins which make the cyst wall of *Giardia* and *Entamoeba*, with the formation of encystation vesicles.

**Key words:** parasitic protozoa, cell secretion, Apicomplexa, trypanosomatids, encystation vesicles.

### INTRODUCTION

The protozoa kingdom comprises a large number of species, including some which are agents of human and veterinary diseases such as malaria, leishmaniasis, Chagas' disease, African trypanosomiasis, amebiasis, trichomoniasis, giardiasis, toxoplasmosis, coccidiosis, theileriosis, and babesiosis, to mention only those more important. Some of these protozoa, as is the case of *Trichomonas*, present a simple life cycle. For others, however, as occurs with Apicomplexa (which includes *Plasmodium*, *Toxoplasma*, *Eimeria*, etc), and some trypanosomatids, the life cycle is relatively complex, displaying several developmental stages in the vertebrate host and, in some cases, in invertebrate hosts. These protozoa are also of interest from the cell biology point of view since they present spe-

cial cytoplasmic structures and organelles, which have been studied in some detail in the last years providing new information of general biological interest. In a previous review we analyzed organelles involved in the metabolic pathways (De Souza 2002). Here, we intend to review, from a cell biology perspective, organelles involved in secretory processes. We will not emphasize aspects well covered in a previous review (Becker and Melkonian 1996).

### CELL SECRETION IN APICOMPLEXA

Since the first studies on the fine structure of protozoa belonging to the Apicomplexa group, especially studies carried out with *T. gondii*, *Eimeria* and *Plasmodium*, it became clear that the anterior region of the infective forms (trophozoites, merozoites, sporozoites) was highly specialized, forming what is generally known as the apical complex

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(Fig. 1). In addition to cytoskeleton components, such as the conoid-sub-pellicular microtubules complex, two organelles were initially recognized and designated as micronemes and rhoptries. Later on another structure, which can be found in other regions of the protozoan body, was recognized and designated as dense granules.

Apicomplexan protozoa present a typical endoplasmic reticulum with associated ribosomes. In *T. gondii* the ER is distributed throughout the cell and the nuclear envelope itself provides a significant fraction of the ER. Vesicles with a fuzzy coat bud off from the nuclear envelope and the ER. In *Plasmodium* the ER is less well developed. Several cisternae of the Golgi complex are observed in the anterior region, just above the nucleus of *T. gondii*. Vesicles with a clathrin-like coat bud off from the trans portion of the Golgi complex. Proteins such as COP I and II, Arf 1 and Sar 1, involved in vesicle formation at the ER-Golgi complex system, have been found in *T. gondii* (Review in Joiner and Roos 2002, Ngô et al. 2000).

#### THE MICRONEMES

Micronemes are small, cigar-shaped organelles that are restricted to the apical third of the protozoan body. Their number varies according to the species and the developmental stages. In some species are hardly seen while in others are so numerous that correspond to the most abundant organelle found in the cell (Fig. 2). The organelle is surrounded by a typical unit membrane and presents an electron dense matrix due to its large protein content. Indeed the organelle is intensely stained when the protozoa are submitted to the ethanolic phosphotungstic acid technique, which reveals basic proteins (De Souza and Souto-Padrón 1978). At present, we still do not have a clear explanation for this labeling pattern since the known micronemal proteins have isoelectric points lower than 7.0. All proteins found in the micronemes are synthesized with an N-terminal signal sequence that mediates their entrance into the secretory pathway by translocation across the endoplasmic reticulum membrane. A mutagenesis anal-

ysis of the C-terminal portion of MIC2 showed the presence of two conserved amino acid motifs mediating the target of this protein to the micronemes (Di Cristina et al. 2000). One motif is a tyrosine-based signal and the other one consists of a stretch of acidic residues.

Many of the micronemal proteins are glycosylated as seen by labeling of the micronemes when thin sections of *T. gondii* are incubated in the presence of gold-labeled lectins (Carvalho et al. 1991).

It has been shown that when the infective forms of Apicomplexan parasites touch the host cell surface they trigger a process of  $Ca^{2+}$  release and the discharge of the content of the micronemes at the junction between the parasite and the host cell (Carruthers et al. 1999a, Bouchot et al. 1999, Vieira and Moreno 2000) which then mediates parasite attachment (Carruthers et al. 1999a, b, Carruthers and Sibley 1999). This process takes place in a few seconds and the released proteins are not incorporated together with the parasites but instead are capped and released from the posterior end of the protozoan (Carruthers et al. 1999a). During redistribution on the parasite surface, transmembrane MICs are thought to connect external receptors to the submembranous acto-myosin motor that provides the power for parasite motility. Chelating of parasite intracellular  $Ca^{2+}$  inhibited both microneme release and invasion of host cells. What is the origin of the  $Ca^{2+}$  used by the protozoan? There is enough data indicating that the  $Ca^{2+}$  used for protozoan motility, micronemal secretion and cell invasion comes from organelles found in the protozoan. Fluorescence microscopy of protozoa labeled with the calcium indicator fluo-4 showed that cytosolic calcium levels underwent dramatic and rapid fluxes (Lovett and Sibley 2003). Several protozoan organelles, such as the endoplasmic reticulum, the Golgi complex, the mitochondrion and the acidocalcisomes may store calcium (Moreno and Zhong 1996, Review in Arrizabalaga and Boothroyd 2004). It has been shown that there is an intracellular calcium release channel with properties of the inositol 1,4,5-triphosphate/ryanodine receptor su-

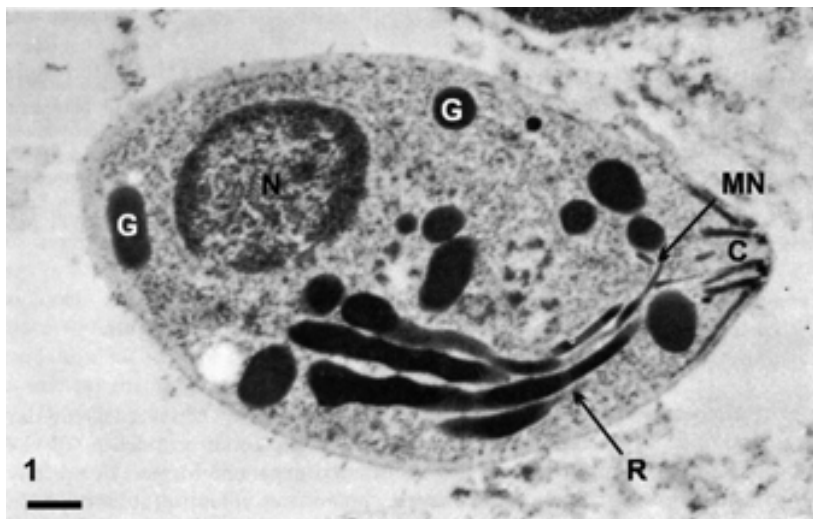


Fig. 1 – Transmission electron microscopy of a tachyzoites of *Toxoplasma gondii* submitted to the ethanolic phosphotungstic acid technique, which labels structures containing basic proteins. In addition to the nucleus (N), staining of the dense granules (G), Rhoptries (R), Micronemes (M) and the Conoid (C) is observed. Bar, 0.3 $\mu$ m. After De Souza and Souto-Pradr3n 1978.

perfamily (Lovett et al. 2002).

It has been shown that isoforms of phosphoglucomutase, a cytosolic enzyme, are implicated in Ca<sup>2+</sup>-mediated signaling events. One isoform, known as parafusin, plays an important role during exocytic activity in ciliated protozoa (Zhao and Satir 1998). A protein called parafusin-related protein has been identified in *T. gondii* and showed to be localized to an apical subpopulation of micronemes and to be redistributed during invasion of the host cell by the protozoan (Matthiesen et al. 2001a, b).

Several micronemal proteins contain one or more adhesive motifs found in mammalian proteins such as Epidermal Growth Factor, integrins, thrombospondin and kallikrein. Four of them, all designated as MIC (MIC1-4, from micronemal proteins), have been studied in some detail.

MIC1 has a size of 60 kDa and contains two degenerate repeats similar to an adhesive sequence found in thrombospondin and known as type I repeats (TSP-I), and is able to bind to the

host cell surface (Fourmaux et al. 1996). It has been shown that it is a lactose-binding lectin (Lourenço et al. 2001). It was also shown that its N-terminal portion functions as an independent adhesin and promotes association with TgMIC4. A galectin-like domain interacts and stabilizes TgMIC6 (Saouros et al. 2005). Single deletion of MIC1 gene decreased invasion of fibroblasts and slightly reduced virulence of the parasite to mouse (C3r3ede et al. 2005).

MIC2 has a size of 115 kDa and comprises an adhesive N-terminal integrin-like A domain that has been implicated in binding ICAM 1 (Barragan et al. 2005) and glycosaminoglycans (Harper et al. 2004), which are ubiquitous sulfated proteoglycans found in the extracellular matrix. MIC2 also displays six thrombospondin type-1 repeats that also have the potential to bind glycosaminoglycans (Wan et al. 1997). It has been shown that a second micronemal protein, known as M2AP, facilitates the transport of MIC2 through the secretory pathway (Huynh et al. 2003, Rabenau et al. 2001). These two proteins seem to form stable hexamers consist-

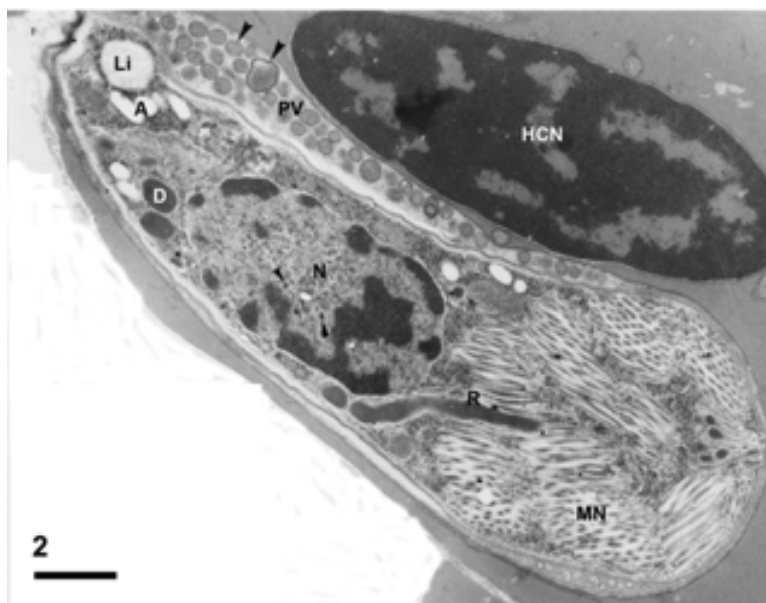


Fig. 2 – Routine transmission electron microscopy showing a trophozoite of *Cyrilia lignieresi*, a haemogregarine found in erythrocytes of a fresh-water fish. Structures such as lipidic inclusions (Li), amilopectin granules (A), dense granules (D), rhoptries (R) and a large number of micronemes (M) are observed. Spherical bodies are seen within the flagellar pocket (small arrowheads). HCN, host cell nucleus. Bar, 1 $\mu$ m. After Diniz et al. 2002.

ing of three  $\alpha\beta$  dimmers (Jewett and Sibley 2004).

Following discharge MIC2 is proteolytically cleaved by proteases with release of its ectodomain from the parasite surface, a process that seems to be involved on parasite invasion (Carruthers et al. 2000, Brossier et al. 2003). Recently, proteins belonging to the rhomboid family of intramembrane-cleaving serine proteases, designated as TgROMs, were detected (Brossier et al. 2005, Dowse et al. 2005).

The binding of MIC2 to the host cell surface may establish connection between the host cell surface receptor and the cytoskeleton machinery of the parasites activating the gliding process necessary for the penetration of the parasite into the host cell. It has been shown that *Plasmodium* sporozoites, which do not express a homologue of MIC2, known as TRAP, fail to glide or invade host cells (Sultan et al. 1997).

MIC3 has a size of 90 kDa and possesses five partially overlapping epidermal growth factor (EGF) domains and an NH<sub>2</sub> terminal chitin binding-like domain, which probably are also involved in the process of parasite association to the host cell surface (Fourmaux et al. 1996, Soldati et al. 2001). It is a disulfide-linked heterodimer comprised of two 38 kDa isoforms (Achbarou et al. 1991). Single substitution of two critical amino acids in the chitin binding-like domains of this protein abolished its binding to cells and decrease parasite virulence (C  r  de et al. 2005).

MIC 4 contains six domains and it binds to host cells (Brecht et al. 2001). In the case of *Eimeria tenella* MIC4 is a transmembrane protein with a molecular weight of 240 kDa, containing 31 tandemly arranged EGF-like repeats in the extracellular domain. These repeats have calcium binding consensus that seem to be involved in make the

molecule to adopt a protease-resistant, rigid structure that could favor its interaction with host cell ligands (Periz et al. 2005).

MIC 6, which is a transmembrane protein, forms trimeric complexes with soluble micronemal proteins such as MIC1 and MIC 4, functioning as an escort protein. Deletion of MIC 6 prevents targeting of these two proteins to the microneme (Reiss et al. 2001). MIC 8 also seems to work as an escort protein to MIC 3 (Meissner et al. 2002).

In *Plasmodium falciparum* a micronemal protein, known as EBA 175, has been shown to bind to sialic acid (Sim 1995) a molecule, which plays a fundamental role on the process of parasite-erythrocyte interaction.

More recently several other proteins which do not present recognizable adhesive motifs have been identified in the micronemes of *T. gondii*. TgMIC 5 has homology to the parvulin family of peptidyl prolyl cis-trans isomerases and may assist in the folding of other micronemal proteins (Brydges et al. 2000). Other proteins, including Tg MIC10 entirely devoid of cysteines, were recently identified (Hoff et al. 2001). MIC 11, a 16 kDa protein, was recently identified in several coccidian parasites. During its traffic through the secretory pathway it is proteolytically cleaved with removal of an internal propeptide, resulting in a mature form containing a  $\alpha$ -chain and a  $\beta$ -chain tethered by a single disulfide bond (Harper et al. 2004).

Members of the genus *Plasmodium* present several developmental stages, which are able to invade different cells in both vertebrate and invertebrate cells. For instance, merozoites invade vertebrate red blood cells, the ookinete invades epithelial cells of the insect vector while sporozoites invade the salivary gland epithelial cells of the insect and when inoculated into the vertebrate host, traverse several cellular barriers until invade hepatic cells. Recent studies have shown that *Plasmodium* presents several genes coding for proteins, which present a membrane-attack complex/perforin-like domain. One of these proteins (PLP1/SPECT2) found in sporozoites was shown to be localized in

the micronemes (Kaiser et al. 2004) and is necessary for cell traversal (Ishino et al. 2005). Genes encoding proteins with similar properties were identified in the genome of *Eimeria* and *Toxoplasma*.

#### THE RHOPTRIES

Rhoptries are long, club-shaped organelles connected by thin necks to the extreme apical pole of the parasite (Figs. 1-4). At their basal portion the matrix of the organelle shows a spongy appearance while the neck region is uniformly electron dense making it difficult the distinction from the micronemes. Their number varies according to the species. Several can be seen in *T. gondii* whereas only two, often designated as paired organelle, are found in *Plasmodium*. The organelle is surrounded by a typical unit membrane and is heavily stained with ethanolic phosphotungstic acid (Fig. 1) (De Souza and Souto-Pradón 1978). Cytochemical studies have shown the presence of glycoconjugates and lectin-like molecules (Carvalho et al. 1991) and  $\text{Ca}^{2+}$  (Pezzela et al. 1997) in the rhoptries. All proteins found in the rhoptries were synthesized in the endoplasmic reticulum and passed through the Golgi complex. Members of the ROP2 family contain multiple independent targeting signals (Bradley and Boothroyd 2001, Striepen et al. 2001). ROP 2 displays both YXX $\Phi$  and LL motifs. Deletion or alteration of these motifs abolishes protein delivery to the rhoptries, with its accumulation in a post Golgi compartment (Review in Joiner and Roos 2002). There is some evidence that rhoptry proteins after leaving the Golgi complex are first accumulated in an intermediate compartment, a type of immature rhoptry which is an acidic organelle, with a pH of 3.5 to 5.5 whereas the mature organelle has a pH of 5.0 to 7.0 as determined using the DAMP technique (Metsis et al. 1995, Shaw et al. 1998). Based on this fact and the involvement of both secretory and endocytic pathways in the rhoptry formation it has been considered that this organelle is equivalent to a lysosome (Metsis et al. 1995, Ngô et al. 2004). More recently, a novel  $\text{Na}^+/\text{H}^+$  exchanger, designated as TgNEH2, was localized in

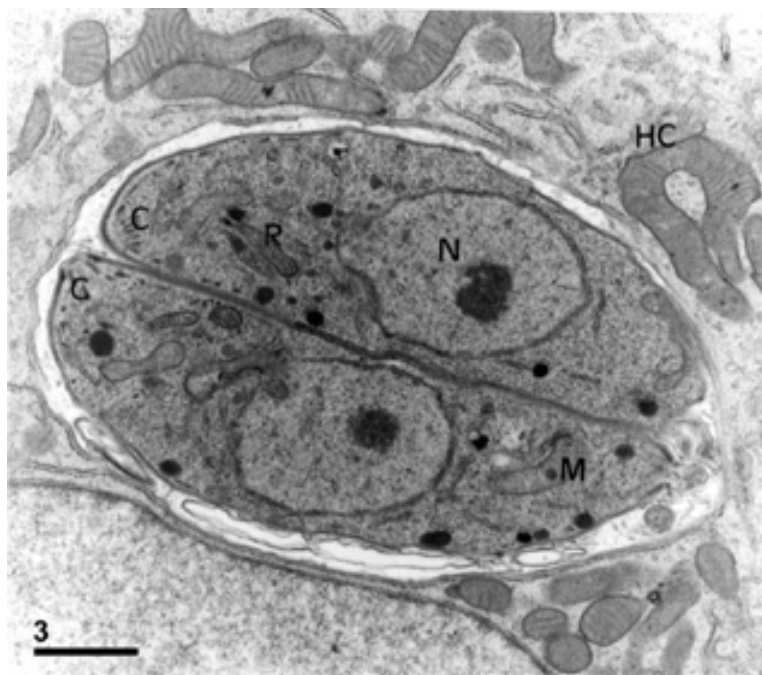


Fig. 3 – Two trophozoites of *Toxoplasma gondii* within the parasitophorous vacuole. C, conoid; HC, host cell; M, mitochondrion; N, nucleus; R, rhoptries. Bar, 0.4  $\mu$ m.

the rhoptry of *T. gondii* (Karasov et al. 2005). This protein may be involved in rhoptry pH regulation. This organelle presents as a characteristic feature the fact that when its lipids and proteins are exocytosed through the duct the organellar membrane is retained and an empty organelle, which can be easily identified by electron microscopy, remains.

Immunocytochemistry provided evidence that the rhoptry content is not homogeneous. For instance, some proteins are localized in the basal region of the organelle whereas others are located in its apical portion (Review in Blackman and Banister 2001).

Secretion of rhoptry proteins takes place immediately after adhesion of the parasites to the host cell surface. In the case of *T. gondii* kinetic studies showed that release of the proteins is completed in about 1 minute and that the proteins are internalized and will make part of the membrane lining the parasitophorous vacuole (Sam-Yellowe et al. 1988, Saf-

fer et al. 1992, Carruthers and Sibley 1997). Several rhoptry proteins have been identified and characterized. ROP1 has a size of 60 kDa and exhibits an extreme charge asymmetry with a highly acidic N-terminal domain and a basic C-terminal domain suggesting that it participates in protein-protein interactions (Ossorio et al. 1992). It has been suggested (Schwartzman 1986) that ROP1 corresponds to the penetration enhancing factor previously characterized in *T. gondii* (Lycke et al. 1968) and for some time it was considered as a key molecule for the penetration of host cells by this parasite. However, it was shown that ROP1 null mutants invade host cells normally (Kim and Boothroyd 1993). ROP2, which has a size of 54 kDa, ROP3, ROP4 and ROP8 are antigenically cross-reactive and may have overlapping functions. ROPs 2 and 8 have been characterized in some detail and cDNA sequences have shown that although they are highly homologous with one another they share no significant homol-

ogy with any other proteins in the databases (Beckers et al. 1994, 1997). Both possess single putative transmembrane segments 75 amino acids from their respective C-termini. OP2 is secreted during the process of penetration of *T. gondii* into the host cell and is found in association with the membrane lining the parasitophorous vacuole. Its N-terminal domain is exposed on the cytoplasmic face of the parasitophorous vacuole and may be involved in the association of the vacuoles with cytoplasmic organelles of the host cell such as the endoplasmic reticulum and mitochondria (Sinai et al. 1997, Sinai and Joiner 2001). Little is known about the other ROP proteins, which have been identified using monoclonal antibodies (Leriche and Dubremetz 1991). A novel rhoptry protein, designated as BRP1, was recently identified in nascent organelles found during the first division of bradizoites, but not in tachyzoites (Schwarz et al. 2005).

The rhoptries of *T. gondii* have been isolated by subcellular fractionation procedures and biochemical analysis showed a lipid to protein ratio of 0.26, thus indicating their richness in proteins. The cholesterol to phospholipid ratio was 1.48. Phosphatidylcholine was the major phospholipid (Fousard et al. 1991). Proteomic analysis using mass spectrometry of the fraction identified 38 novel proteins. At least 11 of them were localized in the rhoptries, as shown by immunofluorescence microscopy. Some are localized in the bulbous basal portion of the organelle while others are restricted to the neck portion, an observation which points to the existence of different domains in the rhoptries. In addition other proteins such as toxofilin, Rab 11, kinases and phosphatases were also found in the rhoptry (Bradley et al. 2006).

The rhoptries isolated from *P. falciparum* merozoites showed a large number of proteins (Etzion et al. 1991). A large number of rhoptry proteins have been identified in *Babesia*, *Plasmodium* and *Eimeria* (Review in Sam-Yellowe 1999). Many of them present as a special feature the ability to bind to erythrocytes.

#### DENSE GRANULES

The dense granules are spherical organelles distributed throughout the cell rather than localized at the apical complex, with a mean diameter of 0.2  $\mu\text{m}$  (Fig. 4). Its matrix is uniformly electron dense due to the high concentration of protein. Kinetic studies have shown that secretion of the dense granule content takes place after parasite invasion and localization within the parasitophorous vacuole persisting for several minutes (Carruthers and Sibley 1997). In contrast to secretion of micronemes and rhoptries, which takes place in the apical region, dense granule secretion occurs at the lateral regions of the protozoan. The secreted proteins associate with the membrane of the parasitophorous vacuole and with the parasite derived intravacuolar membranous network. Proteins are delivered to the dense granule by the bulk flow pathway (Coppens et al. 1999). Proteins from which specific targeting signals for other organelles have been deleted are localized in the dense granules (Striepen et al. 2001, Reiss et al. 2001).

Several proteins have been identified in the dense granules. GRA1 has a size of 22-27 kDa is relatively abundant and remains either soluble within the vacuole or loosely associated to the membranous network. It binds  $\text{Ca}^{2+}$  with two EF hand motifs and therefore has been considered to be involved in homeostasis of this ion within the vacuole (Cesbron-Delauw et al. 1989). GRA2 has a size of 28.5 kDa and after secretion is tightly associated with the membranous network (Mercier et al. 1993, 1998a) through two amphiphatic alpha-helical domains (Mercier et al. 1998a, Sibley et al. 1995). The expression of an HA9 epitope-tagged form of GRA2 by stable transformation of *T. gondii* showed that it is correctly packaged secreted and targeted. Expression of deletion mutants lacking either of two amphiphatic alpha helices resulted in the production and secretion of proteins, which did not associate in a stable way to the membranous network (Mercier et al. 1998b). Based on the observation that a GRA2 null mutant of *T. gondii* presents an



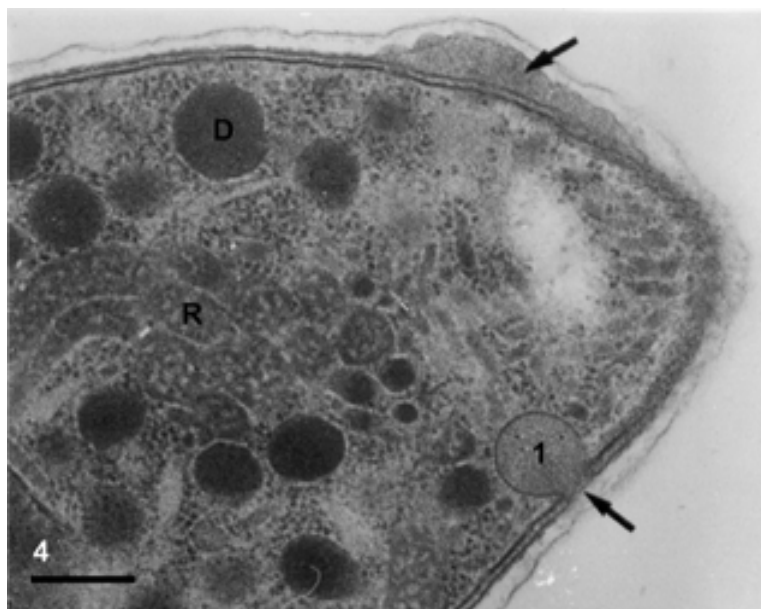


Fig. 4 – Secretion of the dense granule content (arrows) in the lateral side of the tachyzoite form of *Toxoplasma gondii*. D, dense granules; R, rhoptries. Bar, 0.35  $\mu$ m. Courtesy of JF Dubremetz.

attenuated virulence to mice it has been suggested that this protein plays some role in the virulence of the parasite (Mercier et al. 1998b). GRA3 has a size of 30 kDa and forms multimeric complexes that associate with the membrane lining the vacuole through hydrophobic interactions (Ossorio et al. 1994), although there is no predicted membrane-spanning domain for it. However, it was recently shown that GRA3 is actually an artificial chimera of 2 proteins. One, with a molecular weight of 65 kDa, shares the C-terminus of GRA3 and the other, with a predicted molecular weight of 24 kDa, shares the N-terminal region and is recognized by antibodies previously shown to label the dense granules. The corrected GRA3 has a N-terminal secretory signal sequence and a transmembrane domain, which explains its insertion into the membrane lining the parasitophorous vacuole (Henriquez et al. 2005).

GRA 4, 5, 6, 7 and 8 each have one putative transmembrane segment and it has been suggested that they may constitute the molecular sieve that allows the passage of molecules smaller than

1900 Da across the vacuolar membrane (Schwab et al. 1994). These proteins have a size in the range of 21 to 32 kDa. The level of GRA 7 expression is lower in a less virulent strain of *T. gondii*. An additional form of GRA 7 with reduced mobility, probably due to some modification of the protein after exocytosis, was detected on the surface of intact host cells (Neudeck et al. 2002). How this protein reaches the host cell surface is not yet clear. More recently a new protein, designated as GRA 9, was identified. It is a 41 kDa protein, which associates with the tubular network found within the PV (Adjogble et al. 2004). In addition to the GRA series the dense granules also contain two closely related isoenzymes of the nucleotide triphosphatase NTPase which are involved in breaking down host-supplied di- and triphosphate nucleotides participating in the purine salvage pathway, for which *T. gondii* is auxotrophic (Asai et al. 1983, 1995, Sibley et al. 1994) and in the initiation of the release of the parasites from the vacuole (Stommel et al. 1997, Silverman et al. 1998) in a process which in-



Fig. 5 – Stack of flattened lamellae (arrows), which form the Maurer's cleft found in erythrocytes, infected with *Plasmodium falciparum*. P, parasite. Bar, 0.25  $\mu\text{m}$ . After Przyborski et al. 2003.

volves depletion of ATP and increase in  $\text{Ca}^{2+}$  concentration (Stommel et al. 1997, Silverman et al. 1998). Two protease inhibitors have also been identified in the dense granules (Morris et al. 2002, Pszeny et al. 2002).

#### THE MAURER'S CLEFT IN *PLASMODIUM*

Erythrocytes infected by *P. falciparum* show the presence of a cytoplasmic structure, which is labeled when the cells are incubated in the presence of anti-malaria antibodies, and observed by fluorescence microscopy (Tobie and Coatney 1961). Transmission electron microscopy of thin sections of infected erythrocytes revealed the presence of stacks of flattened lamellae of long slender membranes with a translucent lumen, usually located below the erythrocyte plasma membrane, and designated as Maurer's cleft (Fig. 5) (Trager et al. 1966, Langreth et al. 1978). Variable aspects of this structure have been extensively described in several *Plasmodium* species (Review in Przyborski et al. 2003, Lanzer et al. 2006). This structure has not been observed in other members of the Apicomplexa group.

Several parasite proteins have been shown to be localized in the Maurer's cleft. They are synthesized in the parasite ER and then transferred to the cleft (Blisnick et al. 2005, Marti et al. 2005).

Subsequently they are transported to the erythrocyte surface. For some authors the Maurer's cleft is a well defined structure continuously supplied with vesicles budding of from the membrane lining the parasitophorous vacuole. Subsequently, vesicles bud of from the cleft and migrate towards the erythrocyte cell surface, with secretion of their contents into the medium and incorporation of some proteins in the erythrocyte plasma membrane. Vesicles with a diameter of 100 nm have been identified in infected erythrocytes. Homologues of COPI and COPII proteins, which have been shown to play an important role in transport of vesicles in mammalian cells, were identified in *P. falciparum* (Adisa et al. 2002, Wickert et al. 2003). Other groups consider that the Maurer's cleft is part of a continuous network that connects the PV to the erythrocyte cell surface. This view, based on images obtained by confocal laser scanning microscopy, has been confirmed by three-dimensional reconstruction of thin sections (Wickert et al. 2003). According to this view the secretory route would include insertion of proteins in the PV membrane, which then would move along the membrane network by lateral diffusion. Only at the end of these tubular network vesicles would they bud and fuse with the erythrocyte plasma membrane. A recent review covers

well the process of protein transport and trafficking in *Plasmodium falciparum*-infected erythrocytes (Przyborski and Lanzer 2005). Another recent review (Marti et al. 2005) analyses the functional role played by several parasite proteins which concentrates in the Maurer's cleft and are involved in processes such as cytoadherence and import of serum proteins.

#### CELL SECRETION IN TRYPANOSOMATIDS

Members of the Trypanosomatidae family present a well developed endoplasmic reticulum-Golgi complex system with the formation of coated and uncoated vesicles which subsequently migrate towards a specialized and polarized region of the cell surface, known as the flagellar pocket, where they fuse with the membrane (Figs. 6-7) (Reviews in De Souza 1984, Landfear and Ignatushchenko 2001). The flagellar pocket corresponds to a specialized region where most of the endocytic and exocytic activities take place in the trypanosomatids. It is important to point out that the secretory vesicles do not present a dense content and for this reason they are not easily distinguished from the endocytic vesicles which form at the flagellar pocket region and are involved in the uptake of important macromolecules or macromolecular complexes such as transferring and LDL. At least three distinct groups of secretory products have been identified in trypanosomatids based on their fate: (a) one group contains integral or peripheral proteins which are inserted into the flagellar pocket and subsequently migrates to other regions of the plasma membrane. The most evident examples include the synthesis and secretion of the variant surface proteins (VSGs) found in bloodstream forms of *Trypanosoma brucei* and which is involved in the process of antigenic variation (Review in Borst et al. 1998) and cysteine proteinase (cruzin or cruzipain) *T. cruzi* (Fig. 8) (Souto-Padrón et al. 1990); (b) The second group are accumulated in special organelles, as the megasomes of *Leishmania* (Figs. 9-10); (c) the third group includes proteins that are released into the flagellar pocket where they remain as sol-

uble proteins, as is the case of cysteine proteinase (Fig. 11) and proteophosphoglycans in *Leishmania* (Duboise et al. 1994, Foth et al. 2002), while others polymerize within the pocket, as is the case of acid phosphatase found in *Leishmania* (Stierhof et al. 1994). How these different proteins are sorted is not yet clarified. Studies carried with *T. brucei* have shown that VSG lacking its GPI anchor is not efficiently secreted. The proteins may be then mistargeted to the lysosome and is subsequently degraded (Triggs and Bangs 2003).

#### CELL SECRETION IN GIARDIA

As in all eukaryotic cells, trophozoites of *Giardia lamblia* synthesize proteins, which are incorporated into its plasma membrane as well as are secreted into the medium. These proteins are synthesized in the endoplasmic reticulum. Ultrastructural and cytochemical studies have shown that this protozoan does not present a typical Golgi complex system, although structures resembling this organelle can be occasionally seen (Lujan et al. 1995a, Lanfredi-Rangel et al. 1999, Marti et al. 2003, McCaffery et al. 1994). Certainly the more elaborated secretory system in *Giardia* occurs during the process of transformation of the trophozoites into cystic forms, when there is formation of a cyst wall (Review in Marti and Hehl 2003). This process plays a fundamental role in the life cycle of the parasite allowing the development of forms which resist to drastic environmental conditions. Several studies have shown that cyst wall proteins are synthesized in an area of the endoplasmic reticulum where the cisternae is modified, forming a dilated region known as the cleft (Lujan et al. 1995b, Lanfredi-Rangel et al. 2003, Gillin et al. 1996) (Figs. 12-13). Apparently there is no participation of the Golgi complex in this process. The cleft is continuous with the endoplasmic reticulum and lacks an electron dense content. Gradually the cleft widens and becomes filled with a homogeneously dense material formed by the concentration of the cyst wall proteins (CWP) (Figs. 13-14), as can be shown by immunocytochemical



Fig. 6 – Anterior region of an amastigote form of *Trypanosoma cruzi* showing the flagellar pocket (FP), two short flagella (F), the basal body (BB) and the kinetoplast (K). Vesicles are seen close to the flagellar pocket (arrow). Bar, 0.25  $\mu$ m.

localization of these proteins. Although continuity of this structure with the ER is evident, glucose-6-phosphatase, a classical enzyme marker of the ER, is no more found in this structure (Lanfredi-Rangel et al. 2003). Subsequently the dense vesicle, which is now designated as an encystation vesicle (ESV), increases in density (Figs. 15-17) and migrates towards the periphery of the cell (Figs. 18-19). Another view of the process suggests that ER vesicles containing CWP use to each other to form the ESV (Marti and Hehl 2003, Marti et al. 2003). Analysis of the genome data on *G. lamblia* led to the identification of orthologs to factors involved in vesicle tethering and fusion, as soluble N-ethyl-maleimide-

sensitive fusion proteins, Sec-1 adapters, Rabs and the COPI complex. Two syntaxin homologs and two Rab GTPases, designed as Rab 1 and Rab 2, were identified (Marti et al. 2003).

At the cell periphery, the ESV establishes contact both with the inner portion of the plasma membrane of the trophozoites as well as with the peripheral vesicles, acidic organelles that correspond to an endosome-lysosome system (Lanfredi-Rangel et al. 1998). The observation that cyst wall proteins are processed by a cysteine proteinase localized in the peripheral vesicles (Touz et al. 2002), suggests that fusion of the ESV with the peripheral vesicles takes place immediately before or simultaneously



Fig. 7 – Freeze-fracture view of the anterior region of a promastigote forms of *Herpetomonas*. A large number of vesicles (arrows) are seen within the flagellar pocket (FP). F, flagellum. Small arrows point to an aggregation of intramembranous particles, which form the flagellar-cell body adhesion structure. Bar, 0.12  $\mu\text{m}$ . After De Souza et al. 1979.

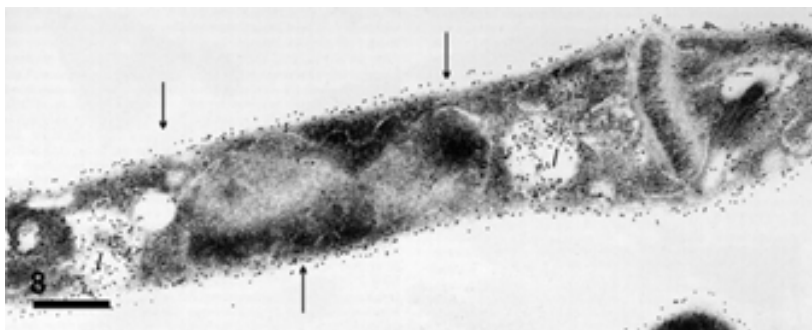
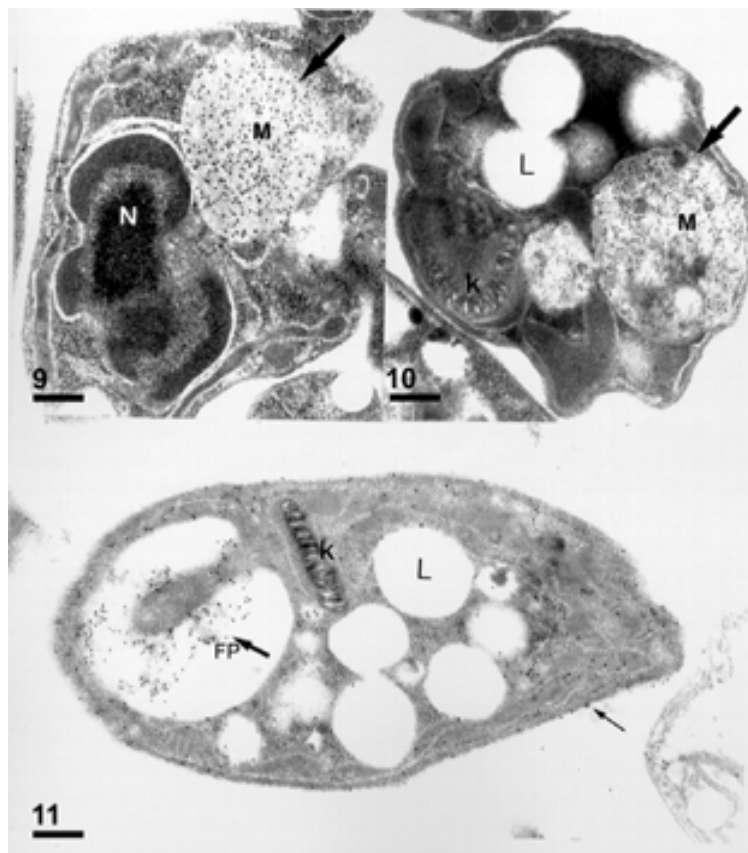


Fig. 8 – Immunocytochemical localization of cysteine proteinase in epimastigotes of *Trypanosoma cruzi*. This protein is synthesized in the ER and concentrated in structures which are part of the endocytic pathway of this protozoan (asterisks) and in the plasma membrane (arrows). Bar, 0.5  $\mu\text{m}$ . After Souto-Pradrón et al. 1990.

with the fusion of the ESV with the cell surface. It is important to point out that the cyst wall proteins are not glycosylated but they have potential N- and O-glycosylation sites (Lujan et al. 1995b, Mowatt et al. 1995). It has been considered that the cyst wall, which is formed by interconnected fil-

aments containing peptides and carbohydrate moieties (Manning et al. 1992), is assembled as consequence of exocytosis of the encystation vesicles (Erlandsen et al. 1996, Gillin et al. 1991, 1996). For some authors there is a dispersal of ESV into small secretory vesicles before secretion (Marti and Hehl



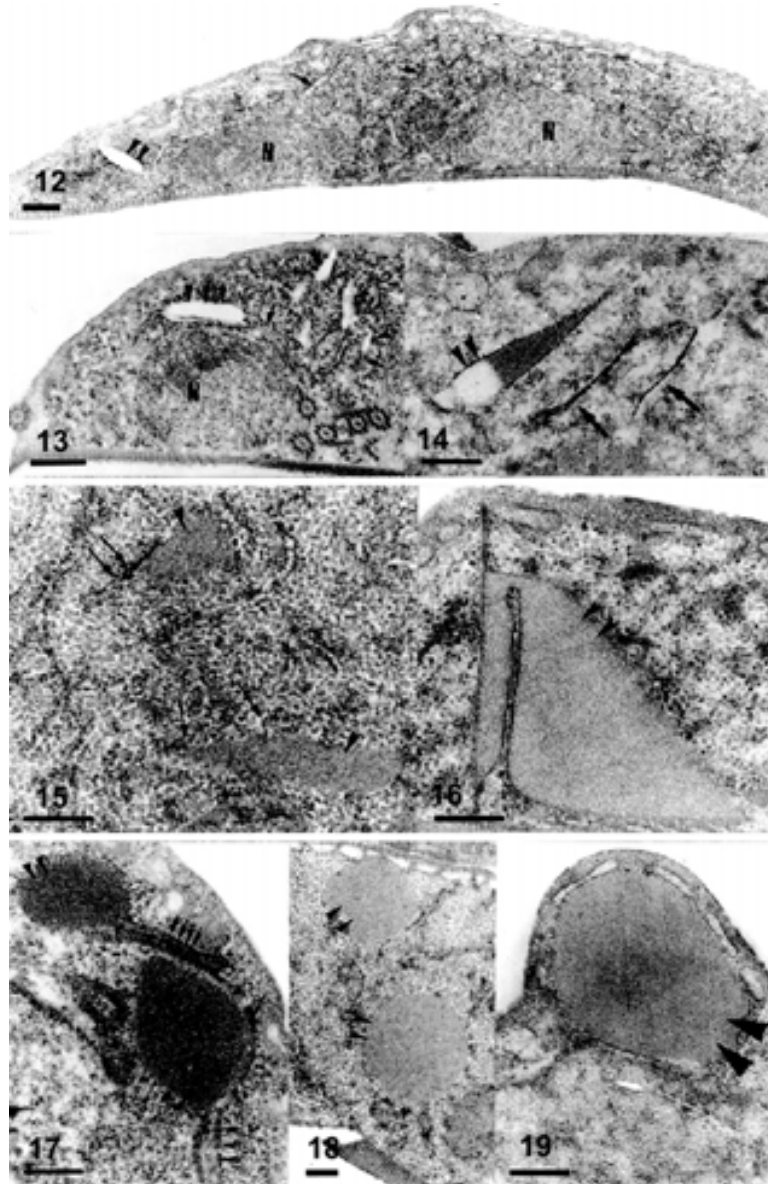
Figs. 9-11 – Immunocytochemical localization of cysteine proteinase in amastigotes of *Leishmania amazonensis*. This protein is stored in the megasomes (M) and secreted into the flagellar pocket. Labeling of the cell surface (arrow in figure 11) and of the cisternae of the endoplasmic reticulum is observed. FP, flagellar pocket; K, kinetoplast; L, lipidic inclusion; M, megasome. Bar, 0.25 $\mu$ m. Micrographs from T Ueda-Nakamura and W de Souza.

2003). More recently it was suggested that a typical exocytosis does not occur. During membrane fusion some membrane segments appeared to be disrupted and released into the extracellular medium where could be resealed forming empty vesicles (Benchimol 2004).

During the period of encystation *G. lamblia* maintains a constitutive pathway for the synthesis of the variant surface proteins. These proteins are not mixed with the ESV proteins. Therefore, the protozoan may have sorting mechanisms to distinguish these two export pathways.

#### CELL SECRETION IN ENTAMOEBA

Members of the *Entamoeba* genus synthesize several proteins, which are secreted. Cisternae of the endoplasmic reticulum and a putative Golgi have been identified by confocal microscopy of cells labeled with NBD-ceramide and by transmission electron microscopy (Mazzuco et al. 1997). Biochemical and molecular studies have shown the presence of an endoplasmic reticulum retention receptor ERD2, a cis-Golgi-associated transmembrane protein (Sanchez-Lopez et al. 1998). BiP,



Figs. 12-19 – Different stages of the process of biosynthesis of cyst wall proteins during encystation of *Giardia lamblia*. The process starts with the formation of clefts in the endoplasmic reticulum (arrows in figure 12), accumulation of dense material within the clefts (arrows in Figs. 13-14), with the formation of encystation vesicles (Figs. 14-17) which then migrates towards the cell periphery (Figs. 17-19). N, nucleus. Bars, 0.4 $\mu$ m. After Lanfredi-Rangel et al. 2003.

a known marker of the ER, has also been identified in *E. histolytica* (Ghosh et al. 1999). Proteins present on the surface of *E. histolytica*, as the Ser-rich protein and the Gal or GalNAc lectin are inserted into the plasma membrane via fusion of secretory vesicles to the membrane. During the process of encystation, which has been studied mainly in *E. invadens*, proteins such as chitinase, localized using an immunocytochemical approach, were seen in many secretory vesicles (Ghosh et al. 1999). During encystation of *E. histolytica* trophozoites large vacuoles with a densely packed filamentous content were observed. They contained chitin since were labeled when cells were incubated in the presence of calcofluor (Chávez-Munguía et al. 2004). It was suggested that these vacuoles are equivalent to the encystation vesicles described during encystation of *G. lamblia*.

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#### RESUMO

Processos de secreção celular desempenham papel relevante na biologia e no ciclo de vida de protozoários patogênicos. A presente revisão analisa, sob uma perspectiva de biologia celular, o processo de secreção em (a) micronemas, roptrias e grânulos densos encontrados em membros do grupo Apicomplexa, onde essas estruturas participam da penetração do protozoário no interior da célula hospedeira, na sua sobrevivência intravacuolar e no posterior egresso da célula hospedeira, (b) a fenda de Maurer, encontrada em *Plasmodium*, uma estrutura envolvida na secreção de proteínas sintetizadas pelo protozoário intravacuolar e transportada, através de vesículas, para a superfície do eritrócito, (c) a secreção de macromoléculas na bolsa flagelar de tripanosomatídeos, e (d) a secreção de proteínas que fazem parte da parede cística de

*Giardia e Entamoeba* e que se concentram nas vesículas de encistamento.

**Palavras-chave:** protozoários parasitas, secreção celular, apicomplexa, tripanosomatídeos, vesículas de encistamento.

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