Characterization of the Role of *aeneas* in Primordial Germ Cell Migration and Blastoderm Cellularization

Dissertation

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by

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2007
meiner Familie
Affidavit

Here I declare that my thesis entitled "Characterization of the Role of aeneas in Primordial Germ Cell Migration and Blastoderm Cellularization" has been done independently and with no other sources and aids than quoted.

Roland Graf
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### Abbreviations

<table>
<thead>
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<th>Description</th>
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<tbody>
<tr>
<td>aa</td>
<td>Amino acids</td>
</tr>
<tr>
<td>AAJ</td>
<td>Apical adhesive junction</td>
</tr>
<tr>
<td>Arp</td>
<td>actin related protein</td>
</tr>
<tr>
<td>BAJ</td>
<td>Basal adhesive junction</td>
</tr>
<tr>
<td>bp</td>
<td>Basepairs</td>
</tr>
<tr>
<td>BSA</td>
<td>Bovine Serum Albumine</td>
</tr>
<tr>
<td>c-DNA</td>
<td>copy-DNA</td>
</tr>
<tr>
<td>DAB</td>
<td>Diaminobenzidine</td>
</tr>
<tr>
<td>Da</td>
<td>Dalton</td>
</tr>
<tr>
<td>DNA</td>
<td>Desoxyribonucleic acid</td>
</tr>
<tr>
<td>Drosophila</td>
<td><em>Drosophila melanogaster</em></td>
</tr>
<tr>
<td>DTT</td>
<td>Dithiotreitol</td>
</tr>
<tr>
<td>EDTA</td>
<td>Ethylene diamine tetra-acetate</td>
</tr>
<tr>
<td>F-actin</td>
<td>Filamentous actin</td>
</tr>
<tr>
<td>FC</td>
<td>Furrow canal</td>
</tr>
<tr>
<td>FCS</td>
<td>Fetal calf serum</td>
</tr>
<tr>
<td>Gal4</td>
<td>Gal4 Transcription factor</td>
</tr>
<tr>
<td>GFP</td>
<td>Green fluorescent protein</td>
</tr>
<tr>
<td>g</td>
<td>Gram</td>
</tr>
<tr>
<td>HRP</td>
<td>Horse radish peroxidase</td>
</tr>
<tr>
<td>JAK</td>
<td>Janus Kinase</td>
</tr>
<tr>
<td>k</td>
<td>Kilo</td>
</tr>
<tr>
<td>LB</td>
<td>Luria Bertani</td>
</tr>
<tr>
<td>Abbreviation</td>
<td>Description</td>
</tr>
<tr>
<td>--------------</td>
<td>-------------</td>
</tr>
<tr>
<td>l</td>
<td>Liter</td>
</tr>
<tr>
<td>LOF</td>
<td>Loss of function</td>
</tr>
<tr>
<td>µ</td>
<td>Micro</td>
</tr>
<tr>
<td>m</td>
<td>Milli</td>
</tr>
<tr>
<td>min or '</td>
<td>Minutes</td>
</tr>
<tr>
<td>mRNA</td>
<td>Messenger RNA</td>
</tr>
<tr>
<td>OD</td>
<td>Optical density</td>
</tr>
<tr>
<td>ORF</td>
<td>Open reading frame</td>
</tr>
<tr>
<td>PBS</td>
<td>Phosphate buffered saline</td>
</tr>
<tr>
<td>PCR</td>
<td>Polymerase chain reaction</td>
</tr>
<tr>
<td>PFA</td>
<td>Paraformaldehyde</td>
</tr>
<tr>
<td>PGC</td>
<td>Primordial germ cell</td>
</tr>
<tr>
<td>PMG</td>
<td>Primordial midgut</td>
</tr>
<tr>
<td>RE</td>
<td>Recycling endosome</td>
</tr>
<tr>
<td>RNA</td>
<td>Ribonucleic acid</td>
</tr>
<tr>
<td>rpm</td>
<td>Rounds per minute</td>
</tr>
<tr>
<td>RT</td>
<td>Room temperature</td>
</tr>
<tr>
<td>RTK</td>
<td>Receptor Tyrosine Kinase</td>
</tr>
<tr>
<td>SAJ</td>
<td>Spot adhesive junction</td>
</tr>
<tr>
<td>SDS</td>
<td>Sodium dodecyl sulfate</td>
</tr>
<tr>
<td>sec</td>
<td>Seconds</td>
</tr>
<tr>
<td>SGP</td>
<td>Somatic gonadal precursor cell</td>
</tr>
<tr>
<td>STAT</td>
<td>Signal transducer and activator</td>
</tr>
<tr>
<td>TGF</td>
<td>Transforming growth factor</td>
</tr>
<tr>
<td>UAS</td>
<td>Upstream activating signal</td>
</tr>
<tr>
<td>ZA</td>
<td>Zonula adherens</td>
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Abstract

The communication and mutual influence of epithelia and migratory cells is a field of high scientific and medical relevance. Many cellular processes underlying the establishment of cell polarity and adhesion are implicated in migratory cell movement including metastasis in cancer.

Primordial germ cells (PGCs) prematurely cross the epithelial barrier of the somatic blastoderm during its cellularization in Drosophila mutants of the aeneas (aen) gene. The objective of this work was the characterization of the biological and molecular function of the Aen protein.

In this thesis I show that Aen is required in the somatic cells for the integrity of the blastoderm epithelium. Aen co-localizes with secretory vesicles and with proteins of the dynactin complex required for vesicular transport. Biochemical interaction experiments as well as co-immunoprecipitation experiments reveal that Aen interacts with Arp1, an important component of the dynactin complex, suggesting a function of Aen during secretory vesicle transport. Furthermore Aen interacts biochemically and functionally with Slow as molasses, which is required for blastoderm cellularization and guided PGC migration.

In summary, the results suggest that Aen encodes a novel conserved adaptor of the Dynactin complex, required for Slow as molasses (Slam) transport or function in a process essential for the epithelial integrity that consequentially inhibits PGC mismigration. Therefore the aen mutation opens new ways to address important questions on tissue invasion by migratory cells.
1 Introduction

Cell migration and the generation of structural and osmotic barriers are important aspects of life. In the eukaryotic domain the means of locomotion of unicellular organisms have laid the foundation for the mode of cell migration found in multicellular organisms (reviewed in Schneider and Haugh, 2006). Even though the molecular mechanisms, that allow the movement of a cell are conserved to a high degree, the implications of migration and the molecular mechanisms underlying the guidance of migrating cells have changed considerably with the transition from protozoan to metazoan life. In multicellular organisms, cell migration no longer serves the purpose of locomotion. Nevertheless, cell migration is a prerequisite of metazoan life, being required for the development and maintenance of a structured and functional organism. During embryogenesis, the formation of the mesoderm (e.g. in Sea urchin, Drosophila, chicken) and the closure of the neural tube (chicken) are exemplary for the involvement of cell migration in morphogenetic movements (reviewed in Gilbert, 2006). In adult organisms, cell migration is involved in wound healing (fibroblasts; reviewed in Li et al., 2004) and immune response (leucocytes; reviewed in Zen and Parkos, 2003).

Accordant to its importance for the functioning of a multicellular organism, aberrant cell migration is implicated in pathological processes, the most prominent example being the spreading of metastases from malignant tumors (reviewed in Sahai, 2005).

The formation of epithelia acting as physical barriers is a second important characteristic of multicellular organisms. Apart from protecting the organism from physical stress, pathogens, desiccation
or osmotic pressure (epidermis, gut epithelia, lung/trachea), other important functions like gas exchange (lung, trachea), the resorption of nutrients (gut epithelium), the transport of metabolites (blood vessel system) and finally the excretion of waste products (kidney, malpighian tubules in Drosophila) are mediated by epithelia (reviewed in Williams et al., 1995; Nation, 2002). To exert these functions, epithelial cells share two main characteristics, apico-basal polarity and tight adhesion. These attributes are essential for epithelial integrity and function (reviewed in Tepass et al., 2001).

In the concert of morphological movements during the development of a higher metazoan organism, the vital processes of cell migration and the maintenance of epithelia must be reconciled. Mechanisms have evolved that regulate the passage of cells through tissues and that allow for rearrangement of adhesive structures in between epithelial cells. However many of the processes and factors involved are not fully understood.

In Drosophila, the formation primordial germ cells (PGCs) requires the positioning of distinct maternally synthesized factors in the oocyte. PGCs, formed by budding out form the posterior embryonic pole, are the first individual cells of the embryo (reviewed in Williamson and Lehmann, 1996). After the somatic lineages have cellularized, the PGCs move from their point of origin, towards the gonadal primordia. Thereby they actively cross epithelial barriers and other tissues. Several mutations, which affect these processes on various levels, have been characterized. Therefore, early embryogenesis of Drosophila melanogaster provides highly established and well accessible experimental system to analyze cell migration, the formation and maintenance of epithelial tissues as well as the interplay between the two cell types.
1.1 *Drosophila* development

1.1.1 Oogenesis in *Drosophila melanogaster*

*Drosophila* oogenesis shows the characteristics of polytrophic meroists (reviewed in Snodgrass, 1993). Nutrients and essential contents of the future embryo are synthesized and imported into the oocyte from germline derived nurse cells and somatic cells (reviewed in Snodgrass, 1993). During this process also the axes of the future embryo are defined by positioning of distinct maternally synthesized factors in the oocyte. This process is a prerequisite for PGC formation (reviewed in Jin and Xie, 2006).

*Drosophila* ovaries consist of about 16 ovarioles, which are surrounded by connective- and muscle tissue. Two to three germline stem cells (GSC) are located in the germarium of each ovariole at its most anterior tip (figure 1a) (reviewed in Huynh and St Johnston, 2004). The stem cell division gives rise to a GSC and a cystoblast, which then undergoes 4 incomplete divisions resulting in an egg follicle of 16 connected cells, the cystocytes (figure 1b). Egg follicles leave the germarium in posterior direction, forming individual egg chambers, which are ensheathed by somatic follicle cells (figure 1a).

During cystocyte divisions a continuous network of microtubules (MT) is established, that extends into all cystocytes through cytoplasmic bridges, the ring canals (Storto and King, 1989; Navarro et al., 2001; Lin et al., 1994; de Cuevas and Spradling, 1998). The pattern of divisions results in two cystocytes with four ring channels, the pro-oocytes (figure 1b; Lin et al., 1994; Lin and Spradling, 1995). One of the pro-oocytes adopts oocyte fate whereas the other pro-oocyte and the remaining cystocytes develop into nurse cells. The differentiation of oocyte fate is dependent on the on MT based transport mediated by dynein and the dynactin complex (Theurkauf et al., 1993; Grieder et
al., 2000; Huynh et al., 2001; Riechmann and Ephrussi, 2001). The dynactin complex which mainly acts as a cofactor of the motor protein complex dynein comprises three subcomplexes, a rod-shaped scaffold consisting mainly of Arp1 (Schafer et al., 1994; Bingham and Schroer, 1999; Schroer and Sheetz, 1991; Gill et al., 1991; Schafer et al., 1994; Schroer et al., 1994; Schoer et al., 1996; Imai et al., 2006), a shoulder and a sidearm complex, containing Dynamitin and a dimer of the protein Glued (reviewed in Schroer, 2004; Imai et al., 2006). BicaudalD (BicD) and Egalitarian (Egl) (Mohler and Wieschaus, 1985; Mohler and Wieschaus, 1986; Navarro et al., 2004), are both associated to the dynein motor complex and act as cargo receptors, but are also required for the polarization of the MT cytoskeleton (reviewed in Pearson and Gonzalez-Reyes, 2004; Riechmann and Ephrussi, 2001). When the functions of the dynactin complex and associated factors are affected, defects in oocyte differentiation occur. The treatment of ovaries with MT depolymerizing drugs leads to the failure of oocyte differentiation, resulting in the formation of egg follicles containing 16 nurse cells and no oocyte (Koch and Spitzer, 1983; Theurkauf et al., 1993). Similar phenotypes are observed in mutants of arp1, bicD and egl (Haghnia et al., 2007; Mohler and Wieschaus, 1985; Mohler and Wieschaus, 1986; McGrail et al., 1997; Mach and Lehmann, 1997; Riechmann and Ephrussi, 2001; Navarro et al., 2004).

After oocyte specification, the nurse cells produce large quantities of organelles, metabolites, mRNAs and proteins, which are transported into the oocyte (reviewed in Spradling, 1993). Also the somatic follicle cells surrounding the oocyte are involved in the synthesis and import of nutrients. The lumen of a mature egg is densely populated by yolk
Introduction

partic and lipid droplets (LD) (Spradling, 1993). The sum of compounds transported into the oocyte is termed maternal contribution (Spradling, 1993).

The majority of factors of the maternal contribution, for example hunchback (hb) mRNA (Lehmann and Nüsslein-Volhard 1987), are ubiquitously distributed in the oocyte. In contrast, factors required for the determination of the anteroposterior and dorsoventral axes of oocyte exhibit specific localization. This localization is based on the activity of the TGF-α signaling factor Gurken (Grk) (Schüpbach, 1987; Cooperstock and Lipshitz, 2001; Roth, 2003; Van Buskirk and Schüpbach, 1999). grk mRNA and protein are transported in a dynactin-dependent manner (Januschke et al., 2002; Caceres and Nilson, 2005; Clark et al., 2007). Grk signaling between the oocyte and the adjacent somatic follicle cells, first at the posterior end of the oocyte, then shifting in dorsoanterior direction, is the founding event of anteroposterior and dorsoventral body axis determination (reviewed in

Figure 1: Oogenesis of Drosophila melanogaster. a) Graphical representation of the development of egg follicles inside of a Drosophila ovariole. The development of egg follicles is shown advancing from anterior to posterior. The scheme on the left side shows the germarium at the anterior tip of the ovariole. On the right side, an egg chamber at stage 9 is depicted displaying the ring canals connecting the cells in the follicle as well as the microtubule (MT) bundles mediating the transport from the nurse cells to the oocyte and the localization of oskar (osk). b) Diagram of cystocyte divisions from the germline stem cell to the 16-cell follicle. The pattern of ringcanals connecting the cystocytes and the process of oocyte selection are depicted. Color scheme: light blue (germline stem cells), light grey (cystocytes/nurse cells), light green (pro-oocytes), dark green (oocytes), red (ring canals), dark blue (MT), dark grey (somatic follicle cells), purple (osk). (Concept of drawing adapted from Huynh et al. 2001).
Gonzalez-Reyes et al., 1995; Anderson and Nüsslein-Volhard, 1984, Poulton and Deng, 2006). During this process the MT cytoskeleton of the oocyte undergoes several steps of repolarization (Gonzalez-Reyes et al., 1995; Theurkauf et al., 1992), which enables the transport of material from the nurse cells into the oocyte and the localized deposition of distinct mRNAs required for the formation of the anteroposterior body axis of the embryo. These factors include the posterior localization of oskar (osk) mRNA (Lehmann and Nüsslein-Volhard, 1986) and the anterior localization of bicoid mRNA (bcd) (Fröhnhöfer and Nüsslein-Volhard, 1986; Weil et al., 2006). Transport and positioning of osk and bcd are MT-dependent (Pokrywka and Stephenson, 1991; Pokrywka and Stephenson, 1995; Clark et al., 2007).

Localization of osk mRNA at the posterior pole, defines the position of the pole plasm, a cytoplasmic region containing several factors in addition to Osk protein, including the RNA helicase Vasa (Vas) (Hay et al., 1988) and nanos (nos) mRNA (Wang and Lehmann, 1991; Ephrussi et al., 1991; Lehmann and Nüsslein-Volhard, 1991). During embryogenesis the contents of the pole plasm will define where the PGCs are formed (reviewed in Hyunh and St Johnston, 2004).

1.1.2 Early embryogenesis

Based on prominent morphological characteristics, which can easily be identified in the developing embryo, the embryogenesis of Drosophila has been divided into a series of distinct developmental stages (reviewed in Foe et al., 1993; Campos-Ortega and Hartenstein, 1985). The Drosophila embryo forms a plasmodium during the first stages of embryogenesis. After fertilization, fast mitotic divisions occur, which lead to the establishment of a large plasmodium. During embryonic stage 3 (figure 2c), 5 to 10 nuclei migrate into the pole plasm located at the posterior pole of the embryo. The nuclei residing in the pole plasm form the first individual cells of the embryo by
budding out from the posterior pole. The cells formed in this process are the PGCs which proliferate and give rise to about 40 cells in later stages (Williamson and Lehmann, 1996). During this stage of development, the Osk protein (see section 1.1.1) acts as a determinant of PGC fate. Ectopic localization of Osk is sufficient to induce ectopic PGC formation in respective positions and in osk loss of function, no PGCs are formed (Ephrussi and Lehmann, 1992). The PGCs strongly differ from the somatic tissue formed at stage 5 in morphology, protein composition and gene activity (see section 1.1.1 and 1.1.3; reviewed in Strome and Lehmann, 2007). Several factors including RNAs (e.g. pole granule component (pgc)) and proteins (e.g. Osk and Vas) are found exclusively in the PGCs (see section 1.1.1 and 1.1.2; reviewed in Strome and Lehmann, 2007). PGCs are transcriptionally silent during stages 3 to 9, which is critical for their survival and the preservation of their germ cell fate (reviewed in Strome and Lehmann, 2007). The polar granules (PGs) are specialized structures in the PGCs, which are strongly enriched in maternal factors such as Osk and Vas (Lehmann and Nüsslein-Volhard, 1986; Hay et al., 1988). Components of the PG are involved in maintaining the

Figure 2: Syncytial stages of embryonic development. a) Stage 1: Pronuclear fusion and mitosis 1. b) Stage 2: Mitotic cycles 2-8, nuclei exhibiting a hollow monolayer shell distribution. c) Stage 3: Mitotic cycles 9-10, nuclei reach the pole plasm, polar budding, formation of the PGCs. d) Stage 4: Mitotic cycles 11-13, alignment of somatic nuclei at the embryonic cortex, formation of the syncytial blastoderm. Color scheme: black (nuclei), blue (stages 1 to 3; pole plasm; purple (stage 4; PGCs). (Drawings after Foe et al., 1993)
transcriptional repression in the PGCs and for the posterior morphogenesis of the embryo (Lehmann and Nüsslein-Volhard, 1986; Hay et al., 1988).

At the beginning of stage 4 (figure 2d) the majority of nuclei have reached the embryonic cortex forming the syncytial blastoderm. The translation of bcd mRNA at the anterior pole leads to the formation of a gradient of Bcd protein which acts as a transcriptional activator for anterior patterning genes in a concentration dependent way (Burz et al., 1998; Rivera-Pomar and Jäckle, 1996; Struhl et al., 1989). The Vas dependent translation of nos mRNA at the posterior pole (Wang and Lehmann, 1991) (see section 1.1.1), results in the formation of a Nos protein gradient. Nos acts as a translational repressor for the morphogen Hb, thereby inducing the derepression of posterior patterning genes (Tautz, 1988; Lehmann and Nüsslein-Volhard, 1987; Hülskamp et al., 1989; Irish et al., 1989; Struhl, 1989). The formation of these morphogen gradients is an essential step in anteroposterior axis formation. The activity of other signal cascades leads to the establishment of the dorsoventral and terminal patterning (reviewed in Moussian and Roth, 2005; LeMosy, 2003). By mitotic cycle 13, about 6000 nuclei form a monolayer directly underneath the embryonic cortex (Foe et al., 1993). The cortical cytoplasm of the embryo is cleared from yolk and LD by MT-dependent transport (Welte et al., 1998; reviewed in Jäckle and Jahn, 1998). The PGCs form a cluster of cells localized at the posterior pole atop the syncytial blastoderm.

1.1.3 Blastoderm cellularization

After the formation of the PGCs at the posterior pole is finished, the cellularization of the somatic blastoderm begins (reviewed in Lecuit, 2004). Directly after nuclear cycle 13, marking the beginning of embryonic stage 5, the formation of somatic buds occurs (figure 3a and c). These protuberances of the plasma membrane, which consist
of a arrays of villous projections (VPs) lined with F-actin, are localized atop of the spherical blastoderm nuclei (Lecuit et al., 2002) (figure 3c). During stage 5-1, the plasma membrane forms hexagonal furrows that synchronously invaginate and vertically enclose the individual cortical nuclei. The leading front of the coherent invaginating membrane structure has been termed furrow canal (FC) (Fullilove et al., 1971). FC formation is initiated and accompanied by reorganization of the actin cytoskeleton (figure 3c and d). While the cortical F-actin concentration decreases, the FC becomes densely populated by actin filaments (Warn et al., 1986) forming contractile rings (Warn et al., 1980; Miller et al., 1995). The protein composition of the FC exhibits great similarity to the contractile apparatus formed during standard cytokinesis. It includes Chickadee, Drosophila Profilin, and the motor proteins Zipper and Spaghetti Squash, the Drosophila homologs of Myosin II heavy and light chain respectively (Young et al., 1993; Karess et al., 1991). Concomitant to the formation of the FC and adjacent to it, the basal adherence junction (BAJ) is formed, which connects the membranes of neighboring cells during their cellularization (Hunter et al., 2000; Lecuit et al., 2002). Its main functional entities are the proteins Shotgun (Shg) (Oda et al., 1994, Tepass et al., 1996), the Drosophila homolog of E-Cadherin, which mediates homophilic interactions between adjacent membranes in a Ca$^{2+}$ dependent manner, α-Catenin (Oda et al., 1993) and Armadillo (Arm) (Peifer and Wieschaus, 1990), the Drosophila β-Catenin, which acts as a linker between Shg and the actin cytoskeleton. As cellularization proceeds, the BAJs are translocated in basal direction (Lecuit, 2004).

The cortical nuclei, that have maintained a spherical shape during the syncytial stages, start to elongate vertical to the embryonic surface (Lecuit, 2000; Knoblich, 2000) increasing their length about 2.5 fold (Brandt et al., 2006). When the ingressing FC reaches half of the
Figure 3: Blastoderm cellularization. Cross sections a)-e), g), i)-j). Grazing sections f), h). a) Syncytial blastoderm, stage 4 to 5-1. b) Cellular blastoderm c) Somatic budding; beginning of stage 5-1: villous projections form above cortical nuclei. d) Formation of the FC (FC): actin and myosin II, localized to the leading front of the FC, form hexagonal shaped rings (see also f)); invagination of membranes between the nuclei; nuclear elongation; formation of the basal adhesive junction (BAJ). e) and f) Slow furrow progression; Beginning of stage 5-2. g) and h) Fast furrow progression; formation of the apical spot adhesive junctions (SAJ); contraction of actin-myosin ring begins; Beginning of stage 5-3. i) Condensing of SAJs into apical adhesive junction (AAJ); contraction of actin-myosin ring leading to the pinching off of individual cells. j) End of cellularization; formation of the cellular blastoderm; resolving of BAJ. Color scheme: grey (nuclei), red (actin filaments), dark blue (MT), purple (Centrosomes), yellow (BAJ), green (SAJ and AAJ).
nuclear length embryonic stage 5-2 begins (figure 3e and f). While the inward progression of the FC commences at slow rates, the nuclei finish their elongation. When the FC reaches the basal end of the nuclei, at stage 5-3, the invagination speed increases (Lecuit et al., 2002). During this fast phase of membrane invagination (figure g-i), the contractile rings at the FC begin to contract. After the progression of the FC comes to a halt, this contraction finishes (figure 3j) leading to an incomplete pinching of individual cells which leaves a thin connection to the underlying yolk, the cytoplasmic stalk (Foe et al., 1993).

During the entire process of cellularization, the surface of the plasma membrane surrounding the embryo and separating the nuclei increases approximately 23 fold. The invagination of the membranes during cellularization is only partially achieved by stretching of preformed membrane reservoirs. Instead it is mainly brought about by integration of de novo synthesized material (Lecuit et al., 2002; Burgess et al., 1997). This process provides the force that propels the ingress of the membranes. In the onset of membrane invagination, gradually 4 membrane domains are formed, the apical-, the apicolateral-, the lateral- and the basal membrane (Lecuit and Wieschaus, 2000; Tepass et al., 2001) (figure 3i). The region, at which newly synthesized material is integrated shifts from the apical membrane during stages 5-1 and 5-2 to the apicolateral membrane during fast furrow progression at stage 5-3. The extension of the membrane thereby coincides with the generation of specialized membrane areas such as adhesive junctions (Lecuit et al., 2002). This finding indicates that the membrane, as it is formed, undergoes a polarized remodeling driven by the targeted integration of membrane material. Supporting this model, many factors and structures involved in intracellular transport are involved in cellularization.
The MT, even though not directly involved in propelling the invagination of the newly formed membranes are required for the process of cellularization. Inhibiting the formation of MTs by injection of anti-Tubulin antibodies leads to a complete inhibition of FC progression (Warn et al., 1987).

In mutants of the gene lava lamp (lva)(Sisson et al., 2000; Papoulas et al., 2005), cellularization is inhibited as well. Lva is a scaffold protein that connects Golgi particles to the dynein motor complex enabling the transport of Golgi vesicles along MTs.

Nuclear fallout (Nuf) (Rothwell et al., 1998) and the small GTPase Rab11 are localized to the recycling endosomes (RE) (Ullrich et al., 1996; Sasamura et al., 1997). nuf and rab11 mutants exhibit defects in vesicle fusion to the membrane region of the prospective FC and redistribution of actin filaments (Riggs et al., 2003). In both mutants, malformation of FC can be observed, leading to the formation of multinucleate cells indicating that the centrosome-associated REs, which are localized directly apical of the nuclei, are involved in the trafficking of membrane and cytoskeletal components during cellularization.

The activity of the formin Diaphanous (Dia) and RhoGEF2 is required for the assembly of the actin structure prior to formation of the FC. In dia and RhoGEF2 mutants, the concentration of F-actin at the FC is reduced. Additionally its morphology is altered and exhibits enlarged membrane invaginations filled with cytoplasmic blebs. In some areas the FC does not form leading to multinucleate cells (Großhans et al., 2005).

Nullo is involved in the organization of the actin cytoskeleton of the FC (Simpson et al., 1990; Postner and Wieschaus 1994; Hunter et al., 2002). Furthermore, it is required for the establishment and stabilization of the
BAJ (Hunter and Wieschaus, 2000). In *null* mutants the formation of the BAJ is compromised, leading to a partial failure of cellularization and formation of multinucleate epithelial cells (Simpson et al., 1990; Hunter et al., 2002). Serendipity-α (Sry-α), which colocalizes with Nullo (Vincent et al., 1985; Schweisguth et al., 1990; Ibnsouda et al., 1993), acts in conjunction with Nullo to maintain the cytoskeleton at the FC.

The protein Slow as molasses (Slam) is required for the proper localization of cytoplasmic myosin and the adhesion factor D-PATJ (Bhat et al., 1999; Pielage et al., 2003). In *slam* null mutants the FC and adjacent BAJ (Lecuit et al., 2002, Stein et al., 2002) are malformed. Consequentially, the cellularization is delayed and stops prematurely. This observation indicates that the Slam protein is involved in the formation of the FC and lateral BAJ (Lecuit et al., 2002), during cytokinesis. Two models have been suggested, how Slam could affect the formation of these domains. It might either act as a scaffolding protein bringing D-PATJ and other factors in contact to each other or it might be involved in directing membrane transport inferring specificity to vesicle fusions taking place in the respective membrane areas. The latter of the models is supported by the finding that Slam loss of function alters the kinetics of membrane invagination (Lecuit et al., 2002) and that Neurotactin (Nrt) (Hortsch et al., 1990), a heterophilic adhesion factor, which is localized to the apical, apicolateral and lateral membrane domains in wildtype, is mislocalized to the apical cytoplasm in *slam* loss of function. Additionally, in experiments tracing the incorporation of membranes into a fluorescently labeled plasma membrane, it was found that, while the sites of membrane incorporation are closely restricted to the apical and apicolateral plasma membrane in wildtype, this distinction does not exist in Slam lack of function (Lecuit and Wieschaus, 2000). However both postulated models for the function of Slam might apply (Lecuit et al., 2002).
During fast furrow progression, at stage 5-3, apical spot adhesive junctions (SAJ) are generated which fuse to give rise to the apical adherence junction (AAJ) and exhibit a similar composition as the BAJ. During gastrulation, this structure forms a continuous Zonula Adherens (ZA) connecting the epithelial cells (Müller and Wieschaus, 1996; Tepass, 1996). The joining of the SAJs into a continuous AAJ is in part modulated by the Nullo protein, which appears to protect adhesion factors of the BAJ of being recruited into the AAJ. When nullo is overexpressed, no AAJ is formed. Instead the individual SAJs are stabilized (Hunter and Wieschaus, 2000).

The formation of the AAJ, which develops into the ZA after completion of cellularization, is dependent on three protein complexes that define the apico-basal polarity of the forming epithelial cells (reviewed in Johnson and Wodarz, 2003). Components of the Bazooka complex, consisting of Bazooka (Baz) (Müller and Wieschaus, 1996, Knust et al., 1996). Drosophila atypical Protein Kinase C (DaPKC) (Wodarz et al., 2000) and DPar-6 (Tomancak et al., 2000), are localized to the subapical region of the blastoderm cytoplasm, apical of the ZA (Wodarz et al., 2000; Petronczki und Knoblich, 2001; Bachmann et al., 2001; Hong et al., 2001). This localization requires the presence of Arm (Müller and Wieschaus, 1996). In arm mutants the formation of adhesive structures is affected (Müller and Wieschaus, 1996) leading to the disintegration of the blastoderm epithelium during gastrulation. In loss of function of baz, markers for apical membranes and the components of the BAJ are mislocalized and the formation of the ZA is compromised leading to malformation of the blastoderm epithelium, similar to the phenotype observed in arm mutants, and to extensive cell death at later stages (Bilder et al., 2003; Müller and Wieschaus, 1996). Similar phenotypes can also be observed for loss of function mutants of other components of the complex (Wodarz et al., 2000; Petronczki and Knoblich, 2001).
The Crumbs complex consists of Crumbs (Crb) (Tepass et al., 1990; Wodarz et al., 1995; Wodarz et al., 1993; Jürgens et al., 1994), Stardust (Sdt) (Müller and Wieschaus, 1996; Wodarz et al., 2000), DLin-7 (Bachmann et al., 2004) and D-PATJ (Pielage et al., 2003). Like the Baz complex, it is localized to the subapical region of the epithelial membranes. Crb is a transmembrane protein that acts as a structural component of the ZA. Mutants affecting the crumbs complex exert adhesion phenotypes similar to the mutants of arm or constituents of the Baz complex. The requirement of Baz for Crb localization indicates that the Crb complex acts downstream of Baz (Bilder et al., 2003; Müller and Wieschaus, 1996; Müller, 2000). The third protein complex involved in the generation of apicobasal polarity is the Discs large complex, which acts downstream of the Baz and Crb complexes and encompasses the factors Discs large (Dlg) (Perrimon, 1988), Scribble (Scrib) (Bilder et al., 2000) and Lethal giant larvae (L(2)gl) (Strand et al., 1994). Similar to mutations affecting the Baz or Crb complex, the formation of the ZA is compromised in mutants of the Dlg complex (Bilder et al., 2003), however no ectopic cell death is observed in this case.

The contraction of the contractile ring during fast furrow progression is timed by a mechanism involving the factor Bottleneck (Bnk). In mutants of bnk, this contraction happens too early during stage 5-2 leading to constrictions pinching the cortical nuclei (Schejter and Wieschaus, 1993; Theurkauf, 1994).

After the invagination of the plasma membranes is finished and the FC has fully contracted, individual cells have formed and establishment of the cellular blastoderm is complete. The LDs are imported into the newly formed epithelial cells through the cytoplasmic stalks. This transport is again MT-dependent (Welte et al., 1998; reviewed in
1.1.4 PGCs in embryogenesis

When the PGCs reach the location of the prospective gonad, they differentiate into gonadal stem cells, which give rise to sperm or eggs in the imagos (reviewed in Williamson and Lehmann, 1996).

The guidance of PGC migration in vertebrates is to a great extent mediated by a simple system consisting of the chemokine SDF1 and its receptor CXCR4 (Doitsidou et al., 2002; Molyneaux et al., 2003). In Drosophila, which contains neither of these factors, the guidance of PGC migration is dependent on a variety of factors and pathways.

During embryonic stages 4 and 5, the PGCs reside at the place of their formation, at the posterior pole of the somatic blastoderm. During this phase, the PGCs are shaped spherically, exhibit a uniform layer of cortical actin and do not form pseudopodia (Jaglarz and Howard, 1995). The JAK/STAT signal transduction pathway appears to affect PGC migration, proliferation and maintenance at blastoderm stage (Li et al., 2003). Overactivation of the pathway during early embryogenesis leads to preterm transmigration at of PGCs at the cellular blastoderm stage (Binari and Perrimon, 1994; Hou et al., 1996). After blastoderm cellularization (see section 1.1.3), the embryo enters gastrulation. The portion of primary epithelium on which the PGCs reside (Turner and Mahowald, 1976) is part of the primordial midgut anlage (PMG)(figure 4a).

At stage 6, the ventral blastoderm epithelium stretches along the longitudinal axis of the embryo. This process, the germ band extension (figure 4b-d), leads to a dorso-anterior shift of the PMG. During this movement, the PGCs are passively carried along with the underlying PMG. Even though the PGCs do not actively migrate, extensive
changes in their morphology can be observed. The PGCs form pseudopodia, making contacts to the PMG and other PGCs (Jaglarz and Howard, 1995). These changes are also reflected in the cytoskeletal morphology. The cortical actin levels detected around the pseudopodia are reduced and focal accumulations of F-actin become visible in the cytoplasm (Jaglarz and Howard, 1995). In the
onset of germ band extension the PMG invaginates (figure 4c), resulting in the formation of the PMG pocket. With the convolution of the PMG, the PGCs are passively internalized and remain tightly packed in the blind end of the PMG pocket until the end of stage 8 (Skaer, 1993; Jaglarz and Howard, 1995).

Embryonic stage 9 marks the transition to active migration of the PGCs (figure 4d and e). They intercalate the epithelial cells of the PMG pocket and transmigrate the epithelium. Several studies have shown that this process is not exclusively mediated by the PGCs but that it also requires changes in the epithelial cells of the PMG (Jaglarz and Howard 1994; Callaini et al., 1995; Warrior, 1994).

During PGC transmigration, the PMG epithelium undergoes morphological changes, which can be observed also in the absence of PGCs (Jaglarz and Howard 1994; Callaini et al., 1995). The epithelial cells of the PMG form pseudopodia on their luminal surface, contacting the PGCs. Upon contact with the PGCs, discontinuities in the adhesive belts of neighboring PMG cells occur, permitting the transit of the PGCs (Jaglarz and Howard, 1994). The PMG epithelium therefore appears to possess specialized traits that actively promote the process of transmigration. In good agreement to this notion is the finding that the PGCs are trapped the PMG pocket of serpent (srp) and huckebein (hkb) mutants. In these mutants, the PMG epithelium is misdetermined to become hindgut epithelium (Reuter, 1994; Warrior, 1994; Jaglarz and Howard, 1994; Brönner and Jäckle, 1996; Moore et al., 1998). A similar PGC migration phenotype has been shown for mutants of the G-Protein coupled receptor Trapped in endoderm 1 (Tre1) (Kunwar et al., 2003). The downstream target of Tre1 is the Rho GTPase 1, which has been suggested to mediate cytoskeletal rearrangements occurring specifically in migrating PGCs (Kunwar et al., 2003).
Tre1 has also been hypothesized to act in conjunction with components of the Wunen signaling pathway (Kunwar et al., 2003). The timing and directionality of PGC migration during and after their transmigration of the PMG pocket is governed mainly by the Wunen pathway. The proteins Wunen (Wun) (Zhang et al., 1996) and Wunen2 (Wun2) (Starz-Gaiano et al., 2001) are lipid-phosphate phosphatases. It has been shown that both Wun proteins act as receptors for a lipid derived second messenger, which is hypothesized to act as an attractant and survival signal for PGCs and that is modified and internalized by PGCs and somatic tissue alike in a Wun dependent manner (Starz-Gaiano et al., 2001; Renault et al., 2004). Somatic tissues expressing Wun at high levels, such as the ventroanterior part of the PMG, act repellant on the PGCs by depriving their surroundings of the second messenger (Starz-Gaiano et al., 2001). Accordingly, the PGCs, migrating away from the wun expressing tissue, transmigrate the PMG in dorso-posterior direction. Somatic Loss of function of wun leads to mismigration of PGCs (Renault et al., 2004).

After crossing the PMG, the PGCs arrange dorsally of it (figure 4f), next to the visceral mesoderm (Jaglarz and Howard, 1995). During stages 10 and 11, the PGCs partition into two groups which split along the longitudinal axis of the embryo and migrate away from ventral midline in lateral directions. Again wun signaling is implicated in the directionality of PGC migration. wun2 is expressed in an ectodermal domain along the midline and lateral of the mesoderm. The PGCs, avoiding the wun2 expressing tissue migrate in lateral direction to meet the visceral mesoderm (Renault et al., 2004; Kusch and Reuter, 1999).

The protein Slam (see section 1.1.3), which is involved in cellularization, appears to have a function in PGC migration at the stages 10 to 11 as well. Mutations of the slam gene cause mismigration of PGCs after the
cells have left the PMG. The functional role of Slam in PGC guidance is unclear as its protein levels decrease below detection level shortly after blastoderm cellularization is finished (Stein et al., 2002) a fact that is interpreted by a Slam function during cellularization that affects PGC migration later in development.

Mutations of STAT92E, a major component of the Jak/STAT signal transduction pathway, cause a reduction of PGC number and mismigration of PGCs after the transmigration of the PMG at stage 10. This mismigration leads to a complete loss of gonad formation (Li et al., 2003; Brown et al., 2006).

When the PGCs have left the PMG they migrate in lateral direction and finally enter the visceral mesoderm where they contact the mesodermal somatic gonadal precursor cells (SGPs) (figure 4 f and g). The directionality of migration thereby is dependent on several factors including Columbus (Clb) (Van Doren et al., 1998) a HMG CoA Reductase, involved in isoprenoid biosynthesis and the Hedgehog signaling pathway (Hh) (Deshpande et al., 2001; Tabata and Takei, 2004). Factors which are involved in the specification of the mesoderm, Zinc finger homeodomain protein 1 (Zfh-1) (Broihier et al., 1998), the FGF-receptor Heartless (Htl) (Gisselbrecht et al., 1996), Fushi-Tarazu (ftz) (Wakimoto et al., 1984; Duncan et al., 1986) and Oddpaired (Opa) (Jürgens et al., 1984) are as well required for PGC migration (Moore et al., 1998). PGCs and SGPs form loose cell accumulations, which are shifted ventrolaterally in the course of germ band retraction during stages 12 to 13 (figure 4 h and i). Mutations affecting the development of the gonadal mesoderm like abdominal A (abdA) (Cumberledge et al., 1992; Karch et al., 1990), abdominal B (abdB) (Celniker et al., 1989), trithorax (trx) (Mazo et al., 1990), clift (cli) (Boyle et al., 1997; Bonini et al., 1993) and tinman (tin) (Bodmer et al., 1993) cause the dissociation of PGCs from the mesoderm during these
stages, resulting in a dispersed distribution of PGCs (Boyle and DiNardo, 1995; Moore et al., 1998).

During germ band retraction, the accumulations of PGCs and SGPs compact, culminating in the coalescence of the primordial gonads during stage 14 (Moore et al., 1998). In this process a compact cluster of PGCs is formed and enveloped by SGPs (figure 4j). In mutants of the transmembrane protein Fear of intimacy (Foi), the gonad coalescence remains incomplete without affecting the fate of the cells (Van Doren et al., 2003). As the mutation of the adhesion factor shg (see section 1.2.1) causes a similar phenotype in the gonads, (Jenkins et al., 2003) Foi has been suggested to act as a regulator for shg expression or function.

1.2 The aeneas gene and protein

The objective of this work was the characterization of the role of the gene aeneas (aen) (Molitor, 2002) during the processes of blastoderm cellularization and PGC migration. The ectopic induction of aen in muscle apodeme precursor cells causes the occurrence of fused muscle fibres with highly abnormal morphology (Molitor, 2002). Also, in contrast to wildtype, muscle fibres crossing the ventral midline occur. This phenotype laid the basis to the further analysis of aen in the group of Dr Gerd Vorbrüggen.

The sequence analysis of the genomic locus of aen, which is located at cytogenetic band 82A6 (Flybase, 1999; http://flybase.bio.indiana.edu/) of the 3rd chromosome, indicated that the gene contains 8 exons and codes for two transcripts which arise from differential promoters initiating the transcription of either exon 1a or 1b and collinear usage of exons 2-6 (figure 5 a and c; Molitor, 2002). The conceptual translation of the 2695 bp and 2542 bp transcripts resulted in two hypothetical protein sequences of 546 aa and 339 aa
respectively (figure 5d). In the course of this study these variants will be termed AenA and AenB. Sequence comparison of the AenA protein sequence with sequences obtained from the computational database showed, that Aen belongs to a conserved group of proteins (Molitor, 2002).
To analyze the aen gene in mutant condition, flies of the line aen\(^1\), which contains a transposable P-element \(l(3)L1233^{R80}\) in the first intron of both transcripts, were analyzed. It was found that no apparent muscle phenotypes occur in embryos originating from aen\(^1\) parents. However, the embryos exhibit defects in PGC migration (Molitor, 2002). For further analysis of the aen gene, a null mutant allele was generated by remobilization of the \(l(3)L1233^{L1233}\) P-element (Robertson et al., 1988), which led to removal of genomic sequence by imprecise excision in one of the lines generated. The line aen\(^2\) was established, which contains the deletion \(\Delta l(3)L1233^{L1233}C2\) covering 1466 bp. This deletion results in the removal of the complete second and about half of the third exon of aen (Molitor, 2002) (figure 5 b). Primary analysis of the P-insertion and the deletion mutant alone or in trans to the deficiency \(Df(3R)(XM3)\) removing the entire genomic region showed that in all cases premature transmigration of PGCs through the newly formed epithelium of the PMG occurs (Molitor, 2002).

Analysis of aen transcription by a developmental Northern Blot showed that the gene is expressed weakly throughout development but that there is strong maternal expression (Molitor, 2002). During early embryogenesis, at stages 1 to 5, high levels of aen transcripts are detectable. Visualization of the aen transcripts in whole mount \textit{in situ} stainings of embryos showed that aen transcripts are ubiquitously present during stages 1 to 4 and with stage 5 are reduced in the blastoderm with exception of the PGCs and the PMG epithelium (Molitor, 2002). At stage 9 to 10, aen transcript levels were reduced in the PGCs and instead became detectable in the domains of the invaginating midgut and, from stage 11 to 12, in the visceral mesoderm (Molitor, 2002.)
The objective of my work was to characterize the biological and molecular function of the Aen protein in order to understand the mechanisms underlying the temporal and spatial control of invasive PGC migration.

For this purpose, a detailed analysis of the PGC mismigration phenotype in aen mutants was performed. The abundance of Aen protein variants in the course of oogenesis and embryogenesis was examined. Factors interacting with the Aen protein were identified and the biological relevance of the interactions found was characterized both functionally in genetic interaction experiments as well as on the molecular level in a detailed biochemical analysis.
2 Materials and methods

2.1 Molecular Biology

2.1.1 Small scale plasmid-DNA preparation

All chemicals if not specified otherwise were obtained from Sigma-Aldrich.

Small amounts of plasmid DNA were isolated from 1.5 ml bacterial culture according to the LiCl-Boiling method (Sambrook, 1989) or using the QuiaQuick mini prep Kit (Qiagen) according to the protocol supplied by the manufacturer. The dried DNA pellet was redissolved in 40 µl TE buffer at 60°C.

2.1.2 Medium scale plasmid DNA preparation

For midi preparations of Plasmid DNA, a 150 ml bacterial culture was grown overnight. The DNA preparations were then carried out with the Qiagen Plasmid Midi Kit (Qiagen) according to the protocol supplied by the manufacturer. The DNA pellet was redissolved in 100 µl TE buffer at 60°C.

2.1.3 Large scale plasmid DNA preparation

For maxi preparations of Plasmid DNA, a 300 ml culture was grown overnight. The DNA preparations were then carried out with the Qiagen EndoFree Maxi Kit (Qiagen) according to the protocol supplied by the manufacturer. The DNA pellet was redissolved in 100 µl EndoFree EB at 60°C.
2.1.4 Extraction of genomic DNA from flies

For extraction of genomic DNA from flies (adapted from Gloor et al., 1993), a single fly was placed in a 1.5 ml reaction tube and paralyzed by cold. The fly was squished with a pipet tip and suspended in 50 µl of squishing buffer (10 mM Tris-Cl pH 8.2, 1 mM EDTA, 25 mM NaCl, 200 µg/ml Proteinase K (Sigma)). After 30 min incubation at 37°C, the Proteinase K was heat-inactivated by incubation for 2 min at 95°C.

2.1.5 Determination of DNA concentration

For the determination of the DNA concentration in aqueous solute, a Genesys 6 Spectrophotometer (Thermo Spectronic) was used for absorption measurement at 260 nm wavelength.

2.1.6 Sequencing of DNA

DNA-sequence analysis was carried out MWG-Biotech AG or by Gordon Dowe (MPI biophysical chemistry) using the Prism ready reaction kit (ABI) together with the automated sequencer ABI 377/96 (ABI). The samples were diluted to 100 ng/µl. 500ng of template, 6pmol of primer and 2.5 µl of kit solution were combined for the sequencing reaction.

2.1.7 Sequence analysis and Primer design

For the alignment and analysis of DNA sequences, the software Sequencher (Gene Codes Cooperation) was utilized. For the primer-design the software OLIGO® 4.06 Primer Analysis Software (National Biosciences) was used. Reference sequences were retrieved from NCBI (http://www.ncbi.nlm.nih.gov; Altschul et al., 1990) and Flybase (http://flybase.bio.indiana.edu; Flybase, 2003).
2.1.8 Sequence IDs of the sequences included in the phylogenetic tree of AenA

The AenA full-length protein sequence was retrieved from Flybase (http://flybase.bio.indiana.edu/). Sequences Conserved between 11 Drosophila species were identified by BLAT search (http://genome.ucsc.edu/index.html) and the software-tool Evoprinter (http://evoprinter.ninds.nih.gov). Homologs of the N-terminal conserved domain covering 294 aa from Dm-AenA residue 25 to 318 were identified by BLAST search (http://www.ncbi.nlm.nih.gov/BLAST/; Altschul et al., 1990) using the blastp algorithm. The output was restricted with a difference cutoff of 0.7. The phylogenetic tree was generated using the BLAST Tree Widget (http://www.ncbi.nlm.nih.gov/BLAST/).

NP_649466 (Drosophila melanogaster); XP_001359602 (Drosophila pseudoobscura); EAT46680 (Aedes aegypti); XP_395864 (Apis mellifera); XP_966481 (Tribolium castaneum); XP_310457 (Anopheles gambiae); NP_00103950 (Bos Taurus); XP_851637 (Canis familiaris); NP_001038564 (Danio rerio); XP_001085714 (Macaca mulatta); XP_414185 (Gallus gallus); BAB14383 (Homo sapiens); XP_511153 (Pan troglodytes); XP_001376232 (Monodelphis domestica); CAF99528 (Tetraodon nigroviridis); BAE24631 (Mus musculus); CAH89819 (Pongo pygmaeus); XP_001195086 (Strongylocentrotus purpuratus); AAH94085 (Xenopus laevis)

2.1.9 PCR reactions and conditions

For analytical PCR, the HotStar Taq Mastermix kit (Qiagen) was used according to the instruction manual. For amplification of sequences below 3 kb, the PfuTurbo DNA polymerase (Stratagene) was utilized. For Long range PCR with a designated product above 3 kb, the Expand Long Template PCR System (Roche) or the La Taq Kit (TAKARA)
were used according to the instruction manual. The annealing temperatures of were chosen 5°C below the mean annealing temperature of the designated Primers. All PCRs were carried out in a Gene Amp 9700 PCR cycler (Applied Biosystems).

2.1.10 DNA digest with restriction endonucleases

Restriction digests were carried out according to the methods described in Sambrook et al. (1989). The software EnzymeX (The Netherlands Cancer Institute) was used for the determination of buffer conditions in parallel digests. Generally, 1 µg DNA was digested in 50 µl total volume including buffer and restriction endonucleases (Fermentas, New England Biolabs) at the concentrations indicated in the respective instruction manuals. The restriction digest was carried out at 30-37°C for 2 h and subsequently stopped by heat inactivation of the restriction endonucleases at 60°C for 10 min. For buffer exchange in sequential restriction digests, the DNA samples were instead of heat inactivation purified using the QiaSpin Nucleotide Removal Kit (Qiagen) and eluted in 40-45 µl H₂O.

2.1.11 Agarose-gel electrophoresis

For analysis or purification of restriction fragments, the reactions were subjected to agarose-gel electrophoresis on 0.8 - 1.5 % gels in 0.5 x TBE buffer (Sambrook et al. 1989) including Ethidiumbromide (Roth). The stained DNA fragments visualized next to GeneRuler DNA Ladder Mix (Fermentas) with an UV-transilluminator (Raytest) at 366 nm.

2.1.12 Gel extraction of DNA

DNA fragments were isolated from agarose gel with the QiaSpin Gel Extraction Kit (Qiagen) according to the protocol supplied by the manufacturer.
2.1.13 DNA ligation

For ligation (modified from Sambrook et al. 1989), digested plasmids and inserts with compatible ends were mixed in a molar ratio of 4:1 and included in 10 µl reaction volume with T4 DNA Ligase (Fermentas) according to the manufacturers manual. Ligation was performed overnight at 14°C.

2.1.14 Gateway recombination

For gateway recombination, Entry-vectors containing the designated insert inside an rfa-recombination cassette were mixed with Destination vectors containing promoter region and fusion-tags as well as selective traits. The recombination was carried out with the Gateway recombination kit (Invitrogen) according to the manufacturers instruction.

2.1.15 Heatshock transformation of bacterial strains

For transformation, 100-200 ng of DNA were mixed with 50 µl chemocompetent E. coli (protocol adapted from Sambrook et al. 1989). After 30-45 min preincubation on ice, the cells were subjected to a heatshock by 30 s incubation at 42°C and transferred back on ice. After 1 h recovery in 300 µl LB, 50 µl bacterial suspension were plated on LB-Agar plates containing selective antibiotics.

2.1.16 Phenolization of plasmid DNA

For injection of constructs into fly embryos, the DNA needs to be of high purity. Any contamination with proteins and potentially toxic agents has to be avoided. Therefore, the acquired constructs were further processed after Midi Preparation. A phenolization was carried out. The whole DNA yielded by the Midi preparation was resolved in 300 µl final volume of H2O and transferred into a Phase Lock Gel
reaction vial (Eppendorf-Netheler-Hinz GmBH). 300 µl Phenol were added and the liquids were suspended by vortexing thoroughly. For phase separation, the suspension was centrifuged for 2’ at 12 krpm. The watery (upper) phase was transferred into a new a Phase Save reaction vial and 300 µl Phenol/Chloroform were added. The liquids were suspended by vortexing thoroughly and centrifuged. The watery phase was transferred into a new vial and twice washed with 300 µl chloroform as described above. This way, all Phenol was removed from the sample. Then the DNA was precipitated adding 30 µl of 3 M Sodium-Acetate and 750 µl Ethanol (100%, p.A.). For pelleting the DNA was centrifuged for 15’ at 12 krpm. The supernatant was discarded. Then the DNA was rinsed with 500 µl Ethanol (70%) to remove residual salt. The DNA was again centrifuged for 5’ at 12 krpm and the supernatant was discarded. The DNA pellet was dried in the SpeedVac for 5’ and resolved in 20 µl H2O LiChrosolv; Merck).

2.1.17 Preparation of injection mixture from Plasmid DNA

For injection, the DNA was combined with a helper construct that expresses the P-Element-Transposase that enables insertion of the construct into the fly genome. For this purpose, 12 µg of the DNA and 4 µg helper construct were diluted in 200 µl final volume of H2O and precipitated as described in paragraph 2.1.9. The DNA pellet was then resolved in 20 µl H2O (HPLC grade) and the concentration was spectrophotometrically determined. For injection (see paragraph 2.2.1), the DNA concentration was adjusted to 400 ng/µl.

2.2 Genetics

2.2.1 Germ line transformation

For the germ line transformation of fly embryos with P-Element-Vectors (Rubin and Spradling, 1982), the DNA was prepared as described in
section 1.1.16. The DNA mixture was then injected into early the pole plasm of pre-blastoderm embryos of the flystock $w^{sn3}$. The flies developing after this procedure were crossed back against $w^{sn3}$. Transgenic flies were selected from the F2 generation by the expression of the marker gene white indicating the integration of the P-element-plasmid into the genome. Injections of into fly embryos were carried out by Iris Plischke and Ursula Jahns-Meyer (MPI biophysical chemistry).

2.2.2 Culturing of flies

Flies were cultured according to the methods described by Ashburner (Ashburner, 1989). Generally, flystrains were grown at 18-22°C on standard Cornmeal/Agar substrate.

2.2.3 Crosses for the genetic interaction analysis between aen and slam

For the analysis of the slam single mutants, the offspring of $w^{*}; w^{do20.84}m/Cyo, ftz-lacZ$ heterozygous slam1 mutant females and $Df(2L)ED369, P(w^{+mW.Sc	ext{er}	ext-}	ext{FRT,hs3} = 3'.RS5+3.3')ED369/Cyo, hb-lacZ$ heterozygous males was analyzed. While the remaining half of maternal contribution is equally present in all embryos originating from this cross, four genotypes (denoted as slam#;+ in figure 17 and 19) occur in equal ratios:

$w^{*}; w^{do20.84}m/ Df(2L)ED369, P(w^{+mW.Sc	ext{er}	ext-}	ext{FRT,hs3} = 3'.RS5+3.3')ED369$

$w^{*}; w^{do20.84}m/Cyo, hb-lacZ$

$Df(2L)ED369, P(w^{+mW.Sc	ext{er}	ext-}	ext{FRT,hs3} = 3'.RS5+3.3')ED369/Cyo, ftz-lacZ$

$Cyo, ftz-lacZ/Cyo, hb-lacZ$
For the analysis of the slam-aen double mutants, following the same scheme, the offspring of w*;wdo^{20.84}m/Cyo,ftz-lacZ;aen\textsuperscript{2} heterozygous slam\textsuperscript{1} mutant females, bearing the homozygous aen\textsuperscript{2} mutation, and Df(2L)ED369, P\{w^{+mW.Scer\textbackslash FRT.hs3=3'.RS5+3.3'}ED369/Cyo,ftz-lacZ; aen\textsuperscript{2}\} males was analyzed, resulting in homozygous aen\textsuperscript{2} mutant offspring bearing half of the slam maternal contribution. The genotypes generated in this cross (denoted as slam#; aen\textsuperscript{2} in figure 17 and 19) are:

\[
\begin{align*}
\text{w*;wdo}^{20.84}m/ \text{Df(2L)ED369, P\{w^{+mW.Scer\textbackslash FRT.hs3=3'.RS5+3.3'}ED369;aen\textsuperscript{2}\}} \\
\text{w*;wdo}^{20.84}m/ \text{Cyo,ftz-lacZ;aen\textsuperscript{2}} \\
\text{Df(2L)ED369, P\{w^{+mW.Scer\textbackslash FRT.hs3=3'.RS5+3.3'}ED369/Cyo,ftz-lacZ;aen\textsuperscript{2}\}} \\
\text{Cyo,ftz-lacZ/Cyo,ftz-lacZ;aen\textsuperscript{2}}
\end{align*}
\]

2.3 Protein biochemistry

2.3.1 Generation of the Aen antibody

The anti-Aen antibody was generated by Eurogentec by immunization of rabbits with two synthesized peptides.

Peptide I : C+ETAQKRTPRRKGRFV

Peptide II: C+DVGGARIEAVDTTTV

The sera received from Eurogentec were subjected to affinity purification using both synthesized antigens on CNBr-activated Sepharose 4B (Amersham) according to the manufacturers instruction.

2.3.2 Collection of embryos and dissection of ovaries

For egg laying, flies were kept on apple agar plates supplemented with yeast (adapted from Rothwell and Sullivan, 2000). Embryos for Western blotting were collected in intervals from 0 to 3 h, from 3 to 10 h and from 10 to 22 h. Embryos were scraped of the agar surface and
transferred into sieves. For removal of the chorion, the embryos were incubated in a 50 % solution of custom detergent for 3 minutes at RT. The detergent solution was thoroughly washed off with isotonic embryo wash solution (0.8 % NaCl, 0.1 % Triton X-100).

Ovaries were dissected from approximately 4 days old adult females in isotonic embryo wash solution (modified from Heinrich et al., 2000; Rothwell and Sullivan, 2000). All samples were stored at –20°C.

2.3.3 Protein extraction from embryos and ovaries

About 20 µl of embryos, larvae, pupae or ovaries were transferred into a reaction vial and covered with 100 µl RIPA buffer (20 mM Tris pH 7.5; 138 mM NaCl; 0.1 % SDS; 0.5 % DOC; 1 % Triton X-100; 2 mM EDTA; 10 % Glycerol) (modified from Harrison et al., 1998). After three times freezing at –80°C and thawing at 55°C, the samples were grinded twice with a potter. The debris was spun down (2', 5 krpm) and the supernatant was transferred into a new tube.

The protein concentrations were measured against a BSA standard with the BCA Protein Assay Kit (Pierce) in a microtiter plate reader.

Alternatively, the samples were directly covered with 1:5 (V/V) SDS loading dye (Sambrook et al., 1989), grinded with a potter and heated to 95°C for 5'.

2.3.4 SDS-PAGE

SDS polyacrylamide gelelectrophoresis was performed using the Mini Protean II system (Bio-Rad). Gels and buffers were prepared according to standard procedures (Sambrook et al., 1989). PageRuler Prestained Protein Molecular Weight Marker (Fermentas) was used for Protein size estimation.
Materials and methods

2.3.5 Western blot

The Western blots were carried out according to standard procedures (Sambrook et al., 1989) using the Mini Protean II system (Bio-Rad, Munich, Germany). For blotting, Nitrocellulose membranes (Schleicher & Schuell) or PVDF membranes (Immobilon-P) were used. Signals from HRP coupled secondary antibodies were generated using the SuperSignal West Pico Chemiluminiscent Substrate Kit (Pierce). Detection of HRP signals was carried out in an Intelligent Dark Chamber II (Fujifilm Europe GmbH) or by exposition of X-Ray films (Kodak BioMax XAR Film, Kodak). After exposure, films were developed in a developing machine (X-Ray Automatic Processor QX-60, Konica).

2.3.6 Protein expression and purification

For bacterial expression in *E. coli* BL21 (DE3) (adapted from Sambrook et al., 1989), precultures were grown in LB in 50 ml Volume at 37°C over night. Expression cultures were inoculated at a ratio of 1: 20 from preculture to final Volume in LB. Expression cultures were grown until their optical density (OD) reached OD=0.5 at 600 nm wavelength. OD-measurements were done with a Genesys 6 Spectrophotometer (Thermo Spectronic). For induction of protein expression, aqueous IPTG (Sigma) solution was added to a final concentration of 0.4 mM. The cultures were incubated at 30°C until the OD reached a value of OD=1.0. Bacteria were harvested by centrifugation in a RC-5B Sorvall Refrigerated Superspeed Centrifuge (Sorvall) at 4 krpm and 4°C for 15 min. The bacterial pellets were resuspended in ice-cold PBS (130 mM NaCl, 7 mM Na₂HPO₄, 3 mM NaH₂PO₄) and again pelleted by centrifugation at 4 krpm and 4°C for 10 min. Then the pellets were resuspended in 20 ml HEMG-1 buffer per 1l culture volume (25 mM Hepes-KOH ph 7.6, 0.1 mM EDTA, 12.5 mM MgCl₂, 0.5 M NaCl, 0.1% NP40 (IGEPAL CA-630 (Sigma), 10% Glycerol), Complete Roche
Protease Inhibitor mix (Roche) and Lysozyme (Biomol) was added to 10 mg/l final concentration. After 30 min incubation at 4°C, the bacterial lysates where subjected to 2 freeze-thaw cycles and homogenized by sonification. The debris was pelleted from the lysates by centrifugation at 15 krpm and 4°C for 20 min. For affinity purification, 50 % slurry of Glutathione sepharose beads (Amersham Biosciences) was preconditioned by rinsing with HEMG-1 buffer (without protease inhibitors). 20 ml of cleared culture supernatant were added to 3 ml bed volume of 50% Glutathione sepharose beads and rotated at 4°C for 2h to allow binding. Subsequently, the resin was trice washed with HEMG-1, HEMG-2 (25 mM Hepes-KOH ph 7,6, 0,1 mM EDTA, 12,5mM MgCl₂, 0,1 M NaCl, 0,1 % NP40 (IGEPAL CA-630, Sigma), 10 % Glycerol) and HEMG-3 (25 mM Hepes-KOH ph 7,6, 0,1 mM EDTA, 12,5 mM MgCl₂, 0,1 M NaCl, 0,01 % NP40 (IGEPAL CA-630, Sigma), 10% Glycerol) respectively. The washed resin was mixed 1:1 with HEMG-3 including Protease inhibitors. For storage, the resin was aliquoted to 20 µl bedvolume (+ 20µl buffer), frozen in liquid N₂ and stored at -80°C.

2.3.7 In vitro translation

For in vitro transcription/translation, the TNT Coupled Reticulocyte Lysate System (Promega) was used according to the provided protocol to produce radiolabelled (³⁵S) proteins in 50 µl reactions. As a radioactive label, the Redivue ProMix (³⁵S) (Amersham Bioscience) was applied. Translation efficiency was assayed with a Luciferase control reaction (supplied by the manufacturer).

2.3.8 GST-Pulldown

For GST-pulldown, the in vitro translation reactions were diluted to 260 µl total volume with HEMG-1 buffer (see section 2.3.8) including protease inhibitors and centrifuged at 12 krpm and 4°C for 15 min (protocol adapted from Blanke and Jäckle, 2006). 40 µl His6-Gst-AenA
or His6-Gst bound Glutathione sepharose (50% slurry) was mixed with 50 µl of diluted translation product (1:5,2). Additionally 1 µl of Luciferase control reaction (undiluted) was added to serve as an internal negative binding control. To allow binding of in vitro translated proteins, the resin was rotated at 4°C for 2h. After 5 washing steps of 10 min at RT with 10 bed volumes of HEMG-1 buffer, 15 µl of resin was mixed with 30 µl SDS loading dye (Sambrook et al., 1989). After denaturing by incubation for 5 min at 95°C, 10 µl of each sample were loaded onto 12 % SDS-PAGE gels next to PageRuler Prestained Protein Molecular Weight Marker (Fermentas). The gels were dried overnight and exposed to Kodak BioMax XAR X-Ray films (Kodak) for 18-24 hours at –80°C.

2.3.9 Cell culture conditions

Kc167-cells (Cherbas et al., 1977) originating from Drosophila, were grown in 75cm² flasks (BD Biosciences) containing 10 ml Schneider’s Drosophila medium (Gibco Invitrogen) which was supplemented with 10% fetal calf serum (Sigma-Aldrich) and penicillin/streptomycin (0.1 mg/ml, PAA Laboratories) at 25°C.

2.3.10 Expression in cell culture

For expression of fusion proteins in cell culture, cells were seeded at a density of 5 x 10⁵ per well of a 6-well plate and allowed to settle for 1-2h at 5°C. Transfection was carried out using the Effectene Transfection Reagent (Qiagen) according to the supplied manual.

2.3.11 Co-IP

For Co-IP, transfected cells were grown at 25°C for 4 days until they reached confluency. The cells were harvested by thorough rinsing and transferred into 15ml conical bottom tubes (Sarstedt). After
For co-immunoprecipitation, 1 ml AffiPrep ProteinA Support resin (BioRad) was equilibrated by washing with PBS (protocol adapted from Harlow and Lane, 1999). For binding, NaCl was added to 4 ml anti-Myc (Iowa-Hybridoma Bank) cell supernatant to a final concentration of 3 M. The serum was added to the resin and rotated overnight at 4°C. Afterwards, the resin was 3 x rinsed and 2 x washed for 20 min with RIPA-buffer. The culture cell extracts were thawed on ice. 30 µl were saved as an input control, mixed with 30 µl of protein loading dye (Sambrook et al., 1989), denatured at 95°C for 5 min and stored at -20°C. The residual 370 µl of extract were mixed with 30 µl of preconditioned resin and incubated for 24h at 4°C. The supernatant was removed and the resin was twice rinsed and washed 4 x 30 min with RIPA. Then, 30 µl resin and 15 µl RIPA were mixed with 30 µl of Protein loading dye (Sambrook et al., 1989) and denatured at 95°C for 5 min. 10 µl of each sample were loaded onto 8 % SDS-PAGE gels next to PageRuler Prestained Protein Molecular Weight Marker (Fermentas) and subjected to Western Blot (see section 2.3.5).

2.4 Histology and Embryology

2.4.1 Formaldehyde fixation of Drosophila embryos

For fixation, embryos were collected and dechorionated as described in section 2.3.1. Collection intervals of 2-4h embryonic age were
chosen. For stainings using the anti-Vasa antibody, the embryos then were fixated on the shaker in 8ml Heptane, 0.5ml Formaldehyde (37%) and Fixation buffer (100 mM HEPES; 2 mM MgSO$_4$; 1 mM EGTA, pH 6.9) for 20' (modified from Rothwell and Sullivan, 2000). The aqueous phase was removed. The embryos were devitellinized by addition of methanol and vigorous shaking. The Heptane- and interphase were removed and the embryos were three times washed with methanol. For storage, the embryos in methanol were kept at -20°C.

2.4.2 Paraformaldehyde fixation of embryos

For stainings of epithelial morphology, embryos were fixated on the shaker in 5ml Heptane, in 4ml PEM-buffer (100 mM PIPES, 5 mM EGTA, 2 mM MgCl$_2$, pH 6.8, 0.2% Triton X-100 (TX-100)) including 5% PFA and 1 ml Picric acid (Sigma; 13% aqueous solution) for 20' (modified from Kreitmeier et al., 1995). The aqueous phase was removed. The embryos were devitellinized by addition of methanol and vigorous shaking. The Heptane- and interphase were removed and the embryos were three times washed with methanol. For storage, the embryos in methanol were kept at -20°C.

For stainings using fluorescently labeled phalloidin, fixation was carried out as described above replacing Methanol with 80% ethanol.

2.4.3 Dissection and fixation of ovaries

Ovaries were dissected as described in section 2.3.2 and fixated in 5ml PEM-PFA (100 mM PIPES, 5 mM EGTA, 2 mM MgCl$_2$, pH 6.8, 0.2% Triton X-100 (TX-100)) including 4% PFA for 20 min. Fixation was stopped by 2x rinsing with PBT (PBS, 0.1% Tween20). For storage, ovaries were dehydrated stepwise in ethanol and stored at -20°C.
2.4.4 Generation of transcript specific probes for RNA in situ detection

For labeling of RNA-probes, the RNA-Labeling and Detection Kit (Roche) was used according to the manufacturers manual. After labeling, 1 µl DNaseI (RNase-free) was added to the samples followed by 15 min incubation at 37°C. The labeled probe was purified using the RNeasy Mini Kit (QIAGEN), following the instruction manual.

2.4.5 RNA in situ detection in fixated embryos and ovaries

For RNA in situ staining (modified from Lehmann and Tautz, 1994) ovaries were dissected and fixated as described in section 2.4.3. The ovaries were washed in PBT 3 x 5 min and prefixated for 20 min in PBS including 4% formaldehyd. After further washing steps with PBT, the ovaries were transferred into the hybridization solution in a stepwise manner. Equilibration steps included 10 min in (1:1) PBT:Hype-B (50 % Formamid; 5 x SSC; 0,1 % Tween20), in 10 min Hybe-B and 10 min in (1:1) Hybe-B:HybeA (50% formamide, 5x SSC, 0.2 mg/ml sonicated salmon testis DNA, 0.1 mg/ml tRNA, 50 µg/ml heparin).

The ovaries were prehybridized in Hybe-A for 1h at 65°C. Then, the buffer was replaced by 30 µl Hybe-A including 2 µl DIG-labeled RNA-probe. The ovaries were incubated over night at 65°C in a waterbath. After hybridization, the ovaries were washed 10 min at 65°C with prewarmed Hybe-A, 2 x 15 min at 65°C with prewarmed Hype-B, 15 min at RT in PBT:Hybe-B (1:1) and 4 x 15 min at RT in PBT. For detection of the DIG-labeled RNA-Probes, the ovaries were incubated with an anti-Dig AP-conjugated antibody (1:2000, Roche Diagnostics, Mannheim) for 1h at 37°C. After 6 x 15 min washing in PBT at RT the ovaries were transferred into AP-Puffer (20 mM Tris pH 9,5; 100 mM NaCl; 50 mM MgCl2) and washed 3 x 5 min. For the development of the staining, 4,5 1 NBT and 3,5 1 BCIP were added into 1 ml AP-Puffer. The staining was stopped by 3 x 5 min and 1 x 20 min washing in PBT.
The stained ovaries were dehydrated in an ethanol dilution series (30%, 50%, 70%, 90% and 2 x 100%), mounted on a slide in Canada Balsam (Sigma-Aldrich) and covered with a coverslip.

2.4.6 Antibody staining of Drosophila embryos

About 20 µl of fixated embryos were rehydrated by three times 5’ washing with PBT including 5 % goat serum (protocol adapted from Rothwell and Sullivan, 2000). Incubation with the primary Antibody was done at 4°C over night. The primary antibody was then removed by twice washing with PBT including 2% of goat serum. Then, either fluorescence coupled or biotinylated secondary antibodies (Molecular Probes) were applied, diluted in 380 µl PBT including 10 µl Serum, and incubated on the embryos for 2 h. All following steps were carried out at RT.

For detection of the fluorescence-coupled antibodies, the embryos after incubation were washed 6 x 10’ in PBT, 2 x 10’ in PBT and 2 x 5’ in PBS. The embryos were allowed to sediment at the tip of the reaction vial, taken up, mounted on a slide with ProLong Gold antifade reagent (Molecular Probes) and covered with a coverslip and sealed with custom nail polish.

The non-bound biotinylated Antibodies were removed by 2 x 5’ washing with PBT. For the coupling of Horseradish peroxidase (HRP) to the biotinylated secondary antibodies, the ABC-Kit Vectastain (Vector Laboratories) was used according to the manufacturers instruction. For HRP detection, the embryos were transferred into 500 µl PBT including 10 µl Diamino-Benzidine (10 mg/ml in 50 mM Tris-HCl; Sigma) and 10 µl hydrogenperoxide (0,3% in PBT; Fluka). The embryos were then stepwise dehydrated (2 x 10’ in Ethanol 70% and 2 x 10’ in Ethanol 100%), mounted on a slide with Canada balsam (Sigma), covered with a coverslip and sealed with custom nailpolish.
2.4.7 Microscope / Confocal imaging

For taking Confocal images of embryos and cells, the Leica DMRXA2 Confocal Microscope (Leica Microsystems) was used. For image processing, the softwares ImageJ (National Institutes of Health) and Adobe Photoshop (Adobe Systems) were used.

2.4.8 In vivo imaging of embryonic development

For time lapse imaging of germ cell migration, 2 h collections of embryos were made. The embryos were arranged on an apple agar slice and fixed to a coverslip with embryo glue and covered with Halocarbon-Oil (Atochem) (protocol modified from germ line transfection protocol, section 2.2.1; Rubin and Spradling 1982). The coverslip was then applied on an Aluminum slide with a notch fitting the hanging drop with the embryos. Images of the developing embryos were taken from 1 to 12 hours.

For the measurement of cellularization speed, 30 min collections of embryos were treated as described above. For analysis, the Axiovert 200M (Leica) was used, which was operated with the software Openlab (Improvision). Data analysis was conducted with the software ImageJ (National Institutes of Health).

2.4.9 Cuticle preparations

12 h embryo collections were aged in order to allow for complete hatching of viable embryos. The unhatched embryos were dechorionated, mounted in Hoyers Lactate medium (30 g gum Arabic, H₂O to 50 ml, 200 g chloral hydrate 20 g Glycerol, 150 ml lactic acid) (modified from Stern and Sucena, 2000) and incubated at 60°C overnight.
### 2.5 Materials

#### 2.5.1 Bacterial strains

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#### 2.5.2 Primers

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<td>RGO148</td>
<td>5'-TCA GAC CTC CAC GCC CC-3'</td>
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<td>RGO154</td>
<td>5'-TCT GCA TAT ATG GCT TCC AGA-3'</td>
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<td>RGO155</td>
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<td>RGO157</td>
<td>5'-GAG GGT GTA TGC CTA CTT GA-3'</td>
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<td>RGO158</td>
<td>5'-CGC AGT TCC CTG GCC ATG-3'</td>
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<td>RGO159</td>
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<td>RGO188</td>
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<td>RGO189</td>
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<td>RGO200</td>
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<td>RGO201</td>
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### Materials and methods

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<td>RGO228</td>
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<td>RGO234</td>
<td>5'-GCT TGC GCC GCC CAT GTC GCC ATA CAC CCT-3'</td>
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<td>RGO237</td>
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#### 2.5.3 Constructs

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<tr>
<td>pUASP-AenA</td>
<td>maternal expression of AenA in flies</td>
<td>Jochen Deckert (this work)</td>
</tr>
<tr>
<td>pUASP-AenA-fl-eGFP</td>
<td>maternal expression of AenA-eGFP; PCR with RGO045/047 on LD30155, subcloning in pEGFP via BamHI-HindIII, cloning in pUASP via BamHI/BglI-XbaI</td>
<td>Roland Graf (this work)</td>
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<tr>
<td>pCasper4-Aeneas-gen-resc</td>
<td>genomic rescue of aen mutants; PCR with RGO043/044 on wt genomic DNA, subcloning in pTopo-XL, cloning into pCasper4 via EcoRI</td>
<td>Gerd Vorbrüggen (this work)</td>
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<tr>
<td>pCITE-shg-ICD</td>
<td><em>in vitro</em> transcription/ translation of shg-ICD; PCR with RGO085/086 on pUAST-DEcadherin, cloning into pCITE via NcoI-XhoI</td>
<td>Roland Graf (this work)</td>
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<tr>
<td>pENTR-AenA-N</td>
<td>gateway recombination of the aenA cassette; PCR with RGO053/054 on LD30155 and cloning into pENTR-D via NotI-AscI</td>
<td>Roland Graf (this work)</td>
</tr>
<tr>
<td>Construct Name</td>
<td>Purpose / Cloning Strategy</td>
<td>Reference</td>
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<tr>
<td>pENTR-AenB-N</td>
<td>Gateway recombination of the aenB cassette; PCR with RGO056/054 on RH35990 and cloning into pENTR-D via NotI-AscI</td>
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<td>pENTR-Arpl-N</td>
<td>Gateway recombination of the arpl cassette; PCR with RGO059/060 on RH04757 and cloning into pENTR-D via NotI-AscI</td>
<td>Roland Graf (this work)</td>
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<tr>
<td>pENTR-SlamNI-N</td>
<td>Gateway recombination of the slam-N-fragment cassette; PCR with RGO119/237 on LD22808 and cloning into pENTR-D via NotI-AscI</td>
<td>Roland Graf (this work)</td>
</tr>
<tr>
<td>pENTR-SlamCI-N</td>
<td>Gateway recombination of the slam-C-fragment cassette; PCR with RGO120/234 on LD22808 and cloning into pENTR-D via NotI-AscI</td>
<td>Roland Graf (this work)</td>
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<tr>
<td>pETG-TrxA-AenA</td>
<td>Expression of TrxA-AenA in BL21 (DE3); gateway recombination of pENTR-AenA-N with pETG-20A</td>
<td>Roland Graf (this work)</td>
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<td>pETG-MBP-AenA</td>
<td>Expression of MBP-AenA in BL21 (DE3); gateway recombination of pENTR-AenA-N with pETG-40A</td>
<td>Roland Graf (this work)</td>
</tr>
<tr>
<td>pETG-NusA-His6-AenA</td>
<td>Expression of NusA-His6-AenA in BL21 (DE3); gateway recombination of pENTR-AenA-N with pETG-60A</td>
<td>Roland Graf (this work)</td>
</tr>
<tr>
<td>pETG-His6-GST-AenA</td>
<td>Expression of His6-GST-AenA in BL21 (DE3); gateway recombination of pENTR-AenA-N with pETG-30A</td>
<td>Roland Graf (this work)</td>
</tr>
<tr>
<td>pETG-30A-df-ccdB</td>
<td>Expression of His6-GST in BL21 (DE3); from pETG-30A by excision of the ccdB cassette with KpnI-Xhol, blunting and religation</td>
<td>Roland Graf (this work)</td>
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<tr>
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<td>Purpose / Cloning strategy</td>
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<tr>
<td>pUbiP-10xMyc-Arp1</td>
<td>expression of 10xMyc-Arp1 in cell culture; gateway recombination of pENTR-Arp1-N with pUbiP-10xMyc-rfa</td>
<td>Roland Graf (this work)</td>
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<tr>
<td>pUbiP-10xMyc-SlamNterm</td>
<td>For expression of 10xMyc-Slam-N-fragment in cell culture; gateway recombination of pENTR-SlamN1-N with pUbiP-10xMyc-rfa</td>
<td>Roland Graf (this work)</td>
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<td>pUbiP-10xMyc-SlamCterm</td>
<td>expression of 10xMyc-Slam-C-fragment in cell culture; gateway recombination of pENTR-SlamC1-N with pUbiP-10xMyc-rfa</td>
<td>Roland Graf (this work)</td>
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<tr>
<td>pUbiP-3xFLAG-AenA</td>
<td>expression of 3xFLAG-AenA in cell culture; gateway recombination of pENTR-AenA-N with pUbiP-3xFLAG-rfa</td>
<td>Roland Graf (this work)</td>
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<td>pUbiP-3xFLAG-AenB</td>
<td>expression of 3xFLAG-AenB in cell culture; gateway recombination of pENTR-AenB-N with pUbiP-3xFLAG-rfa</td>
<td>Roland Graf (this work)</td>
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<td>pCMV TNT-SlamFrag1</td>
<td><em>in vitro</em> transcription/ translation of bp 1-600 of the slam ORF; PCR with RGO147/128 on LD22808, subcloning in pCRII-Topo and cloning into pCMV TNT via EcoRI</td>
<td>Roland Graf (this work)</td>
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<tr>
<td>pCMV TNT-SlamFrag2</td>
<td>For <em>in vitro</em> transcription/ translation of bp 303-920 of the slam ORF; PCR with RGO125/155 on LD22808, further strategy see above</td>
<td>Roland Graf (this work)</td>
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<td>Purpose / Cloning strategy</td>
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<td>pCMVTNT-SlamFrag3</td>
<td><em>in vitro</em> transcription/ translation of bp 610-1230 of the slam ORF; PCR with RGO154/132 on LD22808, further strategy see above</td>
<td>Roland Graf (this work)</td>
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<td>pCMVTNT-SlamFrag4</td>
<td><em>in vitro</em> transcription/ translation of bp 901-1532 of the slam ORF; PCR with RGO129/134 on LD22808, further strategy see above</td>
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<td>pCMVTNT-SlamFrag5</td>
<td><em>in vitro</em> transcription/ translation of bp 1140-1838 of the slam ORF; PCR with RGO227/228 on LD22808, further strategy see above</td>
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<td>pCMVTNT-SlamFrag6</td>
<td><em>in vitro</em> transcription/ translation of bp 1514-2128 of the slam ORF; PCR with RGO134/157 on LD22808, further strategy see above</td>
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<td><em>in vitro</em> transcription/ translation of bp 2410-3019 of the slam ORF; PCR with RGO158/144 on LD22808, further strategy see above</td>
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<td>reference</td>
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<td>Original vectors used for cloning or recombination</td>
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<td>pUAST</td>
<td>generation of transgenes and zygotic expression</td>
<td>Brand and Perrimon, 1993</td>
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<td>pUASp</td>
<td>generation of transgenes and maternal expression</td>
<td>Rorth, 1998</td>
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<td>pCaspeR4</td>
<td>generation of transgenes</td>
<td>Klemenz et al., 1987</td>
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<td>pEGFP-N1</td>
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<td>Clontech</td>
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<td>pETG-20A (TrxA-rfa)</td>
<td>bacterial expression of TrxA-fusion proteins</td>
<td>A. Geerlof (EMBL)</td>
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<td>pETG-40A (MBP-rfa)</td>
<td>bacterial expression of MBP-fusion proteins</td>
<td>A. Geerlof (EMBL)</td>
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<td>pETG-60A (NusA-His6-rfa)</td>
<td>bacterial expression of His6-NusA-fusion proteins</td>
<td>A. Geerlof (EMBL)</td>
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<tr>
<td>pETG.30A (His6-GST-rfa)</td>
<td>bacterial expression of His6-Gst-fusion proteins</td>
<td>A. Geerlof (EMBL)</td>
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<tr>
<td>pUbip-10xMyc-rfa</td>
<td>expression of 10xMyc-fusion proteins in culture cells</td>
<td>generous gift from A. Herzig</td>
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<tr>
<td>pUbip-3xFLAG-rfa</td>
<td>expression of 3xFLAG-fusion proteins in culture cells</td>
<td>generous gift from A. Herzig</td>
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<td>pCMV TNT</td>
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2.5.4 cDNA vectors

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<td>LD30155</td>
<td>pOT2</td>
<td>DGRC</td>
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<td>aenB</td>
<td>RH35990</td>
<td>pFLC-1</td>
<td>DGRC</td>
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<td>arp1</td>
<td>RH04757</td>
<td>pFLC-1</td>
<td>DGRC</td>
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<td>slam</td>
<td>LD22808</td>
<td>pCS+</td>
<td>Lecuit et al., 2002</td>
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<td>arp11</td>
<td>RH13978</td>
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<td>DGRC</td>
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<td>capZ</td>
<td>RE71950</td>
<td>pFLC-1</td>
<td>DGRC</td>
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</table>

2.5.5 Fly stocks

Following fly stocks were used in this study:

w^1118 (Bloomington #5905), w^*;aen^2 (Molitor, 2002), Df(3R)XM3, ru^1 th^1 st^1 kni^p-1 cu^p e^1/TM3, Sb^1 (Bloomington #1655), w^*;P[w^+mC=VASAGFP]; aen^2 (Gerd Vorbrüggen, this work), w^*;P[w^+mC=UAST-aenA]#1.1 (Gerd Vorbrüggen, this work), w^*; P[w^+mC=UAST-aenA]#1.1.aen^2 (Roland Graf, this work), w^*; P[w^+mC=UASP-aenA-fl-eGFP]#1 (Roland Graf, this work), w^*; P[w^+mC=UASP-aenA-fl-eGFP]#17 (Roland Graf, this work), w^*; UAS-Slam-HA/TM6C, AbdZ (kindly provided by Thomas Lecuit), w^*; wdo^20.84m/Cyo,ftz-lacZ (kindly provided by Ruth Lehmann), w^1118; Df(2L)ED369, Df(2L)ED369, P[w^+mW.Scer\FRT hs3=3'.RS5+3.3']ED369/Cyo, hb-lacZ (Bloomington #8903), w^*; wdo^20.84m/Cyo,ftz-lacZ; aen^2 (Roland Graf, this work), w^*; Df(2L)ED369, Df(2L)ED369, P[w^+mW.Scer\FRT hs3=3'.RS5+3.3']ED369/Cyo, hb-lacZ; aen^2 (Roland Graf, this work)
2.5.6 Antibodies used for Western blot detection and co-immunoprecipitation

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<th>working concentration</th>
<th>reference</th>
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<tr>
<td>anti-Aen</td>
<td>rabbit</td>
<td>1:1000</td>
<td>Roland Graf (this work)</td>
</tr>
<tr>
<td>anti-(β)Tubulin (E7)</td>
<td>mouse-monoclonal</td>
<td>1:50</td>
<td>Iowa-Hybridoma bank</td>
</tr>
<tr>
<td>anti-Myc-E10</td>
<td>mouse-monoclonal</td>
<td>1:10 undiluted (resin loading)</td>
<td>Iowa-Hybridoma bank</td>
</tr>
<tr>
<td>anti-Flag (M2)</td>
<td>mouse-monoclonal</td>
<td>1:1000</td>
<td>Sigma</td>
</tr>
<tr>
<td>HRP anti-rabbit</td>
<td>goat</td>
<td>1:4000</td>
<td>Pierce</td>
</tr>
<tr>
<td>HRP anti-mouse</td>
<td>goat</td>
<td>1:4000</td>
<td>Pierce</td>
</tr>
</tbody>
</table>

The anti-(β)Tubulin (E7) antibody developed by Michael Klymkowsky and the anti-c-Myc (9E10) antibody developed by J. Michael Bishop were obtained from the Developmental Studies Hybridoma Bank developed under the auspices of the NICHD and maintained by The University of Iowa, Department of Biological Sciences, Iowa City, IA 52242.

2.5.7 Antibodies used in immunohistological stainings

<table>
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<th>Working concentration</th>
<th>reference</th>
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</thead>
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<td>Roland Graf (this work)</td>
</tr>
<tr>
<td>anti-Tubulin-E7</td>
<td>mouse-monoclonal</td>
<td>1:50</td>
<td>Iowa-Hybridoma bank</td>
</tr>
<tr>
<td>anti-Lava Lamp</td>
<td>rabbit</td>
<td>1:1000</td>
<td>Sisson et al., 2000</td>
</tr>
<tr>
<td>antibody</td>
<td>origin</td>
<td>working concentration</td>
<td>reference</td>
</tr>
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<tr>
<td>anti-Rab11</td>
<td>rat</td>
<td>1:100</td>
<td>Emery et al., 2005</td>
</tr>
<tr>
<td>anti-Dynein Heavy Chain</td>
<td>mouse</td>
<td>1:100</td>
<td>McGrail and Hays, 1997</td>
</tr>
<tr>
<td>anti-HA</td>
<td>mouse</td>
<td>1:2000</td>
<td>generous gift from R. Jahn</td>
</tr>
<tr>
<td>anti-Discs lost</td>
<td>rabbit</td>
<td>1:500</td>
<td>Bhat et al., 1999</td>
</tr>
<tr>
<td>anti-Nrt (BP106)</td>
<td>mouse-monoclonal</td>
<td>1:10</td>
<td>Iowa-Hybridoma bank</td>
</tr>
<tr>
<td>Alexa Fluor 488 anti-rabbit</td>
<td>goat</td>
<td>1:400</td>
<td>Invitrogen</td>
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<tr>
<td>Alexa Flour 568 anti-rabbit</td>
<td>goat</td>
<td>1:400</td>
<td>Invitrogen</td>
</tr>
<tr>
<td>Alexa Flour 488 anti-mouse</td>
<td>goat</td>
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<td>Alexa Flour 568 anti-mouse</td>
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<td>Invitrogen</td>
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<tr>
<td>Cyanine Cy3 anti-rat</td>
<td>goat</td>
<td>1:400</td>
<td>Jackson Immunoresearch</td>
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<td>Biotinylated anti-rabbit</td>
<td>goat</td>
<td>1:400</td>
<td>Vector Laboratories</td>
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</tbody>
</table>

The anti-Nrt (BP106) antibody developed by Corey Goodman was obtained from the Developmental Studies Hybridoma Bank developed under the auspices of the NICHD and maintained by The University of Iowa, Department of Biological Sciences, Iowa City, IA 52242.

The antibody anti-Discs lost, which was formerly described to detect the protein Dlt (Bhat et al., 1999) has been shown to specifically detect D-PATJ instead (Pielage et al., 2003). In this study the anti-Discs lost antibody was used for the detection of D-PATJ.
3 Results

3.1 Conservation of the Aeneas proteins

The conservation of a protein provides important indications towards its function and structure and its evolutionary age might be indicative of its requirement.

To identify functional domains in the Aeneas (Aen) protein sequence, an analysis using the SMART protein analysis tool (http://smart.embl-heidelberg.de) was conducted, using the AenA protein sequence as input. The result showed that AenA features no domains, which have been functionally characterized.

Therefore, the conservation of Aen within the family of Drosophilids was investigated to identify regions of high conservation. This analysis encompassed known or predicted Aen protein sequences from 11 Drosophila species covering an evolutionary distance of about 40 Mio Years (Ashburner et al., 1989). The sequences were identified by BLAT search (http://genome.ucsc.edu/index.html) and compared utilizing the software-tool Evoprinter (http://evoprinter.ninds.nih.gov). The analysis showed that two regions exist in the Aen sequence that are highly conserved within the Drosophilid family (figure 6b). The conserved N-terminal region covers 294 aa from residue 25 to 318 and a carboxyterminal (C-terminal) region of 41 aa covering residue 416 to 456 of the Drosophila melanogaster AenA protein.

A detailed analysis of the regions conserved between Drosophilids revealed that Aen is a member of a highly conserved protein family.
Sequences of potential Aen homologs were retrieved by analyzing the protein sequence of the conserved N- and C-terminal Aen domains with the blastp algorithm using standard parameters (BLOSUM62 scoring matrix) (www.ncbi.nlm.nih.gov). It was found that the C-terminal region is only conserved within Drosophilids (data not shown), whereas the long N-terminal conserved region exhibits a high degree of conservation in a wide range of species. A phylogenetic tree was generated from the retrieved sequences with the BLAST Tree Widget, using the Fast Minimum Evolution method with a difference cutoff of 0.7 (figure 6a; for BLAST results see section 2.1.8). It shows that the Aen protein is conserved throughout the animal kingdom. A detailed analysis of the Aen homologs suggests that in primates at least two protein variants exist, which arise from alternative usage of an internal
exon. Other potential splice variants, representing aminoterminal (N-terminal) truncations were not included in the phylogenetic tree because the incompleteness of the transcriptome and proteome data in most organisms does not allow for their secure prediction.

The high conservation of Aen indicates that its function is required for animal life.

3.2 Characterization of the phenotype observed in aen mutant embryos

The aen phenotype was analyzed in two aen loss of function (LOF) allelic combinations, in aen\(^2\) and in aen\(^2\)/Df(3R)XM3 transheterozygous embryos (deficiency Df(3R)XM3 covering cytogenetic bands 082A03-06:082B including the aen locus). The aen\(^2\) allele was generated by imprecise remobilization a P-element insertion in the first intron of both aen transcripts (figure 5b) in the germline of the aen\(^1\) mutant (Molitor, 2002). In aen\(^2\) mutants, the remobilization resulted in a deletion of 1476 bp that removes the complete second and part of the third exon of the aen gene (figure 5b), which are included in both aen transcripts, aenA and aenB. Despite the fact that the promoter regions and the transcriptional start sites of both aen transcripts were not affected by the deletion, the removal of major parts of the open reading frames suggested that aen\(^2\) is a null allele or a strong hypomorphic allele. Furthermore, flies carrying the aen\(^2\) allele are viable and fertile, which allowed the investigation of the aen LOF phenotype in the offspring of homozygous aen\(^2\) mutant parents. Therefore, the aen mutant phenotypes presented throughout this work represent a maternal and zygotic null mutant or strong hypomorphic situation.
3.2.1  *aen* mutant embryos show a strong increase in untimely PGC migration

For the analysis 2-4 hours old *aen* mutant embryos and wildtype embryos were collected, fixed and subjected to immunohistochemical stainings using an anti-Vasa antibody, which specifically marks the primordial germ cells (PGCs) (Jaglarz and Howard, 1995). The number of PGCs that were present in ectopic positions was counted. During blastoderm stage, ectopic PGCs are found either intercalated between the forming somatic blastoderm cells or in one plane with the progressing furrow canal (FC) (figure 7b and 7b'). It was found that untimely transmigration of PGCs through the PMG can be detected at earliest at stage 5-3 during blastoderm cellularization, after the FC has passed the nuclei of the somatic blastoderm. Ectopic PGCs occur both in wildtype and in *aen* mutant embryos. Therefore, the numbers of wildtype (wt) embryos exhibiting different phenotypic intensities were counted in order to determine the phenotypic background present in the assayed system. Phenotypic intensities were characterized by the number of mismigrating PGCs. It was found that mismigration of 3 or more PGCs occurs in less than 10 % of wt embryos (figure 7c). Therefore, the phenotypic intensity of 0 to 2 ectopic PGCs per embryo was defined as a cutoff criterion for wt versus mutant phenotypes.

Subsequently, the ratio of embryos exhibiting wt and mutant phenotypes were analyzed in homozygous *aen*<sup>2</sup> mutants, in transheterozygous *aen* mutants bearing the *aen*<sup>2</sup> allele in trans to the deficiency *Df(3R)XM3* and in wt. The results of the subsequent analysis show that in wt, 92 % of the embryos show 0-2 transmigrating PGCs while only 8 % have a higher number of mismigrating PGCs (figure 7d, left). In contrast, 44 % of *aen*<sup>2</sup> mutant embryos exhibit a strong PGC transmigration phenotype while the effect is weak in 56 % (figure 7d,
A comparison of aen\(^2\) homozygous mutant embryos with the offspring of aen\(^2\)/Df[XM3] transheterozygous mutant females, crossed to aen\(^2\) homozygous males, shows that the distribution is shifted towards strong phenotypes in the transheterozygous situation. While 70% of the aen mutant embryos exhibit a strong PGC transmigration phenotype, weak in phenotypic intensities are found in 30% (figure 7d, right).

This result indicates that the aen\(^2\) allele might exhibit residual gene activity. However the deletion of 1476 bp in the aen\(^2\) allele removes
most of the ORF and reduces the transcript and protein amounts to undetectable levels (Molitor, 2002; see below) suggesting that $aen^2$ represents a null allele.

3.2.2 *In vivo* imaging of PGC migration in $aen$ mutant embryos

The morphology of the PGCs observed in the anti-Vasa stainings of $aen^2$ mutant embryos is similar to the one found in PGCs actively transmigrating the PMG at stage 10. The formerly spherical PGCs form filopodia that intercalate in between the underlying epithelial cells. The bodies of cells crossing the forming epithelium exhibit an elongated, spindle-shaped form and readopt a spherical shape once they reach the body cavity (figure 7b'). These observations give a first indication that the cells might actively migrate into the epithelium, however these stainings reflect only a static image of a dynamic process. To clarify in which way the ectopic PGCs become separated from the PGC cluster. Time-lapse movies of living $aen^2$ embryos carrying a Vasa-eGFP construct under the control of the endogenous vasa-promoter were taken (Breitwieser et al., 1996). The Vasa-eGFP fusion protein, which labels the PGCs, was used to track PGC development from their budding out at stage 3 to germ band extension at stage 6 (see stills in figure 7e).

The recorded image series shows that pole cap formation and morphology are normal in $aen^2$ mutant embryos until the beginning of stage 5-2, approximately 60 min after budding (figure 7e'). At this point, several PGCs that are positioned basally in the PGC cluster undergo a change in cell shape, elongating in apicobasal direction in respect to the underlying epithelium (figure 7e''). Thereby they start to translocate in basal direction. This movement continues for the following minutes (figure 7e'''), leading to an ectopic positioning of the PGCs inside the somatic blastoderm (figure 7e''''). By the start of gastrulation, the ectopic PGCs are no longer detected (figure 7e''''').
3.2.3 The PGC transmigration phenotype in aen mutants can be rescued by introducing a genomic fragment covering the aen locus

To show that the aen mutation causes the PGC mismigration phenotype observed, rescue experiments were performed. The genomic region of the aen gene including the flanking sequences up to the 5' and 3' neighboring genes (figure 8a) was amplified by PCR from the BAC clone BACR15G02 and cloned into the pCaspeR4 vector (Klemenz et al., 1987) for the generation of transgenic flies by pole plasm injection (Rubin and Spradling, 1982). Two independent transgenic lines carrying the genomic rescue fragment were crossed into the aen\(^2\) background. In embryos derived from these strains, the ratio of embryos with wt and mutant PGC transmigration phenotypes was examined according to the scheme described in section 3.2.1. The analysis (figure 8b) shows that aen\(^2\) embryos containing one copy
of the rescue construct exhibit lower PGC transmigration rates than \( \text{aen}^2 \) embryos. 44% of \( \text{aen}^2 \) embryos have strong PGC mismigration phenotype. In the \( \text{aen} \) genomic rescue experiment this ratio decreases to 6% using transgenic rescue line #13 and to 22% using the independent transgenic rescue line #14 (wt: 8%; figure 7c). The complete rescue of the mutant phenotype by the addition of one copy of the \( \text{aen} \) genomic rescue fragment shows that the observed PGC mismigration is most likely caused by the loss of \( \text{aen} \) activity.

### 3.2.4 The PGC transmigration phenotype in \( \text{aen} \) mutants can be rescued by Gal4-UAS mediated expression of \( \text{aenA} \)

In order to rescue the \( \text{aen} \) mutant phenotype using an independent technique, the expression of \( \text{aen} \) gene products was induced in \( \text{aen} \) mutant embryos utilizing the Gal4-UAS expression system (Brand and Perrimon, 1993). This system consists of the yeast Gal4-transcriptional activator that, via binding to its DNA target sites (UAS, Upstream Activation Sequence), induces the expression of any cDNA cloned 3' of a minimal promoter with the UAS sites. It is a highly established experimental tool to induce the expression of a target gene in *Drosophila* cells or tissues. *In vivo*, the temporal and morphological domain of the expression can be chosen by crossing transgenic flies carrying a construct bearing the target gene controlled by a UAS containing promoter region to other fly lines that contain Gal4-driver constructs. These constructs bear the Gal4 gene under the control of a promoter originating from a *Drosophila* gene. The activity of this endogenous promoter indirectly induces the transcriptional activation of the target gene by activating the transcription of the Gal4 protein (Brand and Perrimon 1993).

For this purpose, the \( \text{aenA} \) ORF was cloned into the pUAST vector (Brand and Perrimon, 1993) bearing the minimal promoter with the Gal4 responsive UAS sites. The vector was then utilized for the
generation of transgenic flies by pole plasm injection (Rubin and Spradling, 1982). Analogous to the genomic rescue, the transgenic lines carrying the UAS-AenA construct and flies carrying the Nullo-Gal4 driver construct were crossed into the \( \text{aen}^2 \) background. To induce the expression of \( \text{aenA} \) in the somatic blastoderm epithelium underlying the PGCs, the \( \text{aen}^2 \) mutants bearing the UAS-\( \text{aenA} \) and Nullo-Gal4 constructs respectively were crossed. The offspring bearing both the UAS and Gal4 constructs was examined following the scheme described in section 3.2.1.

The analysis (figure 8b) shows that the PGC transmigration rate in \( \text{aen}^2 \) embryos, in which \( \text{aenA} \) is expressed under the control of the Nullo-Gal4 driver construct, is lowered as compared to \( \text{aen}^2 \) embryos. While 44% of \( \text{aen}^2 \) embryos exhibit a strong PGC mismigration phenotype, this ratio decreases to 34% in embryos expressing \( \text{aenA} \). Even though the rescue efficiency observed in this approach is lower as in the genomic rescue, this result indicates that the activity of the \( \text{aenA} \) gene product in the somatic cells underlying the PGCs is sufficient for rescuing the PGC mismigration observed in \( \text{aen}^2 \) mutants.

### 3.3 Expression of \( \text{aen} \)

Northern Blot analysis had shown that \( \text{aen} \) transcripts are present throughout all stages of \( \text{Drosophila} \) development. Furthermore, the transcripts are highly abundant in the female abdomen and during early embryogenesis (Molitor, 2002). A strong ubiquitous signal of \( \text{aen} \) transcripts is detectable at stage 1 of embryogenesis (section 1.2; Molitor 2002). By stage 3, when the PGCs are formed at the posterior pole, \( \text{aen} \) transcripts are included into their cytoplasm. During stage 5-6 the levels of \( \text{aen} \) transcript are reduced in the somatic parts of the embryo presumably by specific degradation, whereas the transcripts in the PGCs remain until stage 9. Based on the high transcript levels present in the early embryonic stages and the strong \( \text{aen} \) expression in
females as revealed by the Northern Blot (Molitor, 2002), aen transcripts are presumably imported into the oocyte as a maternal contribution.

3.3.1 aen transcripts are restricted to the germline during oogenesis

To analyze whether the female specific high transcript levels in the Northern Blot reflect only a strong maternal contribution, or whether aen is expressed earlier in oogenesis or in the somatic parts of the egg follicles as well, an in situ hybridization on dissected ovaries was
performed using a digoxigenin labeled RNA antisense probe detecting both aen transcripts. The RNA in situ staining reveals that aen transcripts are highly abundant in nurse cells and the oocyte after stage 8 (figure 9a). This finding provides a strong indication that aen transcripts are imported from the nurse cells into the oocyte as a maternal load. It also shows that aen gene products are not expressed above the detection level of the RNA in situ during early stages of oogenesis. Additionally aen transcripts are not detected in the somatic follicle cells, surrounding the germline. Together with the fact that homozygous mutant aen² females are viable and fertile it appears highly unlikely, that aen has a major function during oogenesis.

3.3.2 The Aen-antibody specifically detects both Aen protein variants on Western blots

It remained to be tested whether both gene products (figure 5c) are transcribed and whether they are equally present at all developmental stages. The two hypothetical transcripts share a nearly identical size of 2695 bp (aenA) and 2542 bp (aenB) excluding the poly-A-tail and can therefore not be clearly distinguished on a developmental Northern-Blot (Molitor, 2002). The aenA transcript features 232 bases not shared with aenB which in turn has 42 specific bases. The high degree of sequence-overlap does not allow for the generation of transcript-variant specific probes.

To overcome this predicament, it was investigated whether the protein variants inferred from conceptual translation of the two postulated transcripts, which differ significantly in size (section 1.2; figure 5d) are translated in vivo. This analysis was achieved by Western Blot analysis using a specific antibody. Based on the known cDNA sequence two sequences were chosen for peptide synthesis and the immunization of rabbits. The first peptide is located at the native C-
terminus of both protein variants and the second in a region with a high degree of antigenicity, overlapping between AenA and AenB (figure 9b). This antigen selection was made to ensure that both protein variants are detected by the antiserum produced.

Protein extracts of embryos of wt and aen\(^2\) were used in Western Blots (figure 9c). The antibody detection resulted in a main band at an apparent molecular weight of approximately 70 kDa and two weak bands with an apparent molecular weight of around 55 and 40 kDa respectively. The comparison of protein extracts from wt and aen\(^2\) mutant shows that the 70 kDa and 55 kDa bands are exclusively detected in the wt extract, while the 40 kDa band is equally present in both extracts. This finding indicates, that the 70 and 55 kDa bands represent Aen specific signals.

The molecular weight of AenA as deduced from conceptual translation is 59.7 kDa, however analysis of the sequence with the online tool Protein Calculator v3.3 (http://www.scripps.edu/cgi-bin/cdputnam/protcalc3) reveals that AenA bears a PI of 9.7, which would result in 8.8 positive charges at pH 7.5, the pH of the separation gel. The sequence analysis of AenB, with a predicted molecular weight of 37.3 kDa, suggests a PI of 10.3 resulting in 11.7 positive charges at a pH of 7.5. These additional charges might account for a substantial change in the running behavior of AenA and AenB causing the deviation from the predicted molecular weights to the observed molecular weights of 70 and 55 kDa.

The third band with an apparent molecular weight of 40 kDa detectable both in wt and mutant extracts cannot be explained by an partial translation in the aen\(^2\) mutant. The sequence analysis of the aen gene shows that the largest ORF possibly transcribed from the aen\(^2\) allele has a length of only 414 bp resulting in a peptide of 13.8 kDa molecular weight. Therefore, this signal is caused by antibody
crossreactivity with a protein unrelated to AenA and AenB and does not represent an Aen-specific signal.

In the aen\(^2\) mutant no Aen specific protein is detected. This finding strongly indicates that the aen\(^2\) mutation is in fact a null allele and that the reduction of the PGC transmigration phenotype in homozygous aen\(^2\) mutant embryos as compared to aen\(^2\)/Df[XM3] transheterozygous aen mutant embryos (see section 3.2.1) is due to the acquisition of modifications in the genetic background of the aen\(^2\) strain.

3.3.3 AenA and AenB are differentially translated in the course of development

To characterize the developmental translation profile of AenA and AenB a developmental Western Blot of tissues from wildtype flies was performed. The materials analyzed were protein extracts from dissected female ovaries, embryo collections of 0-3 h, 3-10 h and 10-22 h in age. The 0-3 h embryo collection covers embryonic stages 1 to 5 from egg laying including the stages of PGC formation as well as blastoderm cellularization. The 3-10 h embryo collection covers the embryonic stages 6 to 13 from the beginning of gastrulation to the end of germband retraction and includes the resident and migratory phase of the PGCs as well as the alignment of the PGCs with the somatic gonadal precursor cells (SGPs). The 10-22 h embryo collection covers the embryonic stages 14 to 17 from the dorsal closure to embryonic hatching and includes the coalescence of the embryonic gonad consisting of PGCs and SGPs.

The Western Blot (figure 9d) shows that AenA is translated at high levels during oogenesis. This translation is restricted to the late stages of egg-follicle maturation, as inferred from in situ analysis (figure 9a). The AenA levels exhibit a slight reduction in early embryogenesis, increasing
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again after 3h. This reduction is probably caused by the lower cell number in blastoderm embryos as compared to later developmental stages. In contrast to AenA, AenB is detectable at high levels only during oogenesis, its concentration rapidly decreasing thereafter.

3.4 The subcellular localization of Aen

In order to investigate the molecular function of the Aen proteins, it was investigated whether they are specifically localized to subcellular compartments or the cytoskeleton.

3.4.1 The anti-Aen antibody specifically detects Aen in fixed tissue

The anti-Aen antibody was tested for applicability in histological stainings of fixed whole mount Drosophila tissues. First, the specificity of the newly generated anti-Aen antiserum was tested. It was administered to a mixture containing aen\(^2\) mutant embryos and wt embryos expressing Histone2AvD-GFP (Clarkson and Saint, 1999) as a marker to detect specifically the wt embryos. The embryos of both genotypes were combined before dechorionization, fixed and stained jointly. The stained embryos (figure 10a) were analyzed by confocal microscopy. Wt embryos exhibiting Histone2AvD-GFP signal showed a very strong antibody staining in the cytoplasmic space whereas the aen\(^2\) embryos without the Histone2AvD-GFP signal showed only an extremely weak staining. This result showed that the peptide antiserum is highly specific for Aen and applicable for immunohistological stainings of fixed material.

Based on to the weak expression of AenB during embryogenesis in Western Blots, the signal obtained in embryos stained with the anti-Aen antibody reflects mainly the localization of the AenA protein.

Consequential to the PGC mismigration phenotype first observed during cellularization, the subcellular localization of Aen was
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investigated at embryonic stages 4 and 5. Aen was co-stained with Alexa488-conjugated Phalloidin to visualize the actin cytoskeleton, which is strongly enriched to the cell cortex and the FC. Therefore, the actin cytoskeleton serves as a good marker for the overall morphology of the embryo and for the visualization of the progression of blastoderm cellularization (figures 10 and 11; see also introduction section 1.1.3; fig. 3). Aen can be detected in the PGCs as well as in the somatic parts of the embryo. In the PGCs the Aen signal exhibits strong focal enrichments (figure 10 c, arrowhead). This subcellular enrichment remains static in the course of development. In the somatic part of the embryo Aen localization is more dynamic. In pre-blastoderm stages, Aen exhibits ubiquitous cytoplasmic localization but is excluded from

Figure 10: Whole mount antibody staining with the anti-Aen antibody. Immunohistochemical detection of Aen using the anti-Aen antibody in aeni² embryos and embryos expressing Histone2AvD-GFP: double immuno-histochemical detection of Aen/actin in wt embryos using the anti-Aen antibody and Alexa488-conjugated Phalloidin. a) Test of the specificity of the anti-Aen antibody in whole mount antibody staining of embryos. On the right side, a wt embryo marked with Histone2AvD-GFP (here depicted in blue) is visible, which exhibits strong antibody staining (green). An aeni² embryo is located on the left side (outline). The aeni² mutant embryo shows only a very weak staining, indicating that the antibody stains very specifically the Aen proteins. b and c) Aen localization during stage 5-3. Left column: Aen; middle column: Actin; right column: overlay. Aen (green), Actin (red). b-b") Cross section of a stage 5-3 embryo. Posterior pole marked with an asterisk. Aen is ubiquitously present in the cytoplasmic regions of the somatic blastoderm and the PGCs but not detectable in the yolk body cavity. c-c") Cross section of the posterior pole with PGCs (asterisks mark nuclei). Forming FC (arrow). The Aen signal exhibits strong focal enrichments in the PGCs (arrowhead).
Figure 11: The subcellular localization of Aen during blastoderm cellularization. Double immunohistochemical detection of Aen/actin using the anti-Aen antibody and Alexa488-conjugated Phalloidin. Left column: Aen; middle column: actin; right column: overlay. Aen (green), actin [red]. a-a"

Cross sections: Stage 3. a) Aen exhibits a cytoplasmic localization and is excluded from yolk particles (asterisk). a"

High concentrations of cortical actin. b-b"

Stage 5-1: Invagination of the FC (arrow). b) Aen enrichments (white arrowhead) basally of the cortical nuclei (asterisk). b"

The actin cytoskeleton is redistributed from the cortex to the FC. c-c"

Stage 5-2. c) Aen enrichments (arrowhead) also lateral and apical of the nuclei. c"

Concentration of cortical actin is reduced. d-d"

Stage 5-3. d) Aen is cytoplasmically distributed with enrichments in the lateral cytoplasm adjacent to the nuclei and the subapical domain. d"

The dashed lines illustrate the depths of the grazing sections. e-e"

Grazing sections of a stage 5-3 epithelium. e-e"

Apical cytoplasm. Aen localization in the apical cytoplasm exhibits focal enrichments (arrowhead). Actin and Aen exhibit no pronounced overlap in distribution. f-f"

Half of invagination depth. f) Focal enrichments of Aen in the lateral cytoplasm (arrowhead). f"

Actin weakly detected alongside the lateral membranes. g-g"

FC. g) Highest Aen concentration in the cytoplasmic stalks (asterisk). Concentration generally lower than in the apical and lateral cytoplasm. g"

Actin is organized in contractile rings (arrowhead).
the yolk and does not show an overlap with cortical actin (figure 11a). At the beginning of stage 5, Aen is found in a cytoplasmic region surrounding the nuclear monolayer with enrichments directly apical of the individual nuclei (figure 11b, c). During stage 5-2, the FC is surrounded by Aen even though no direct overlap between the Aen and Actin signal can be detected. From this stage on, Aen enrichments are also found apical of the nuclei. This change in localization continues during the fast phase of cellularization (stage 5-3) resulting in the enrichment of Aen in the subapical domain and the lateral cytoplasm (figure 11d). Grazing sections of a stage 5-3 epithelium (figure 11e-g) show, that the Aen localization in the apical cytoplasm is irregular and exhibits multiple focal enrichments (figure 11e), while it is more evenly distributed in the region of the cytoplasmic stalk (figure 11g), adjacent to the FC. The distribution of Aen signal suggests that two pools of the protein exist. One is evenly cytoplasmically distributed, whereas the other co-localizes with subcellular compartments detectable as small granular structures.

In order identify the structures that are enriched with Aen, co-stainings with different markers for structures known to show a similar subcellular distribution and morphology were conducted.

3.4.2 Aen shows a partial overlap with the recycling endosome marker Rab11

The recycling endosome (RE) is localized in the apical cytoplasm of the somatic blastoderm, directly adjacent to the centrioles of the cortical nuclei. It is a membranous compartment involved in endocytosis and membrane traffic. Its function has been shown to be required for the cellularization of the somatic blastoderm (section 1.1.3; Lecuit et al., 2002; Pelissier et al., 2003). Rab11 is specifically enriched in the RE (Sasamura et al., 1997) but also in small particles, which co-localize also with Golgi markers (Pelissier et al., 2003).
Confocal analysis of stage 5 embryos co-stained with antibodies against Aen and Rab11 shows that no large Rab11 positive structures are detectable in the PGCs. In the somatic blastoderm however, the Rab11 antibody marks large structures, characteristic for the RE, directly apical of the cortical nuclei and small particles located in the apical cytoplasm. The overlay of the Rab11 and Aen signals obtained in the co-staining shows differential results for the RE and the small Rab11 marked vesicles. Whereas the presumptive REs do not show a high degree of overlap with the Aen signal, the small particles are in most cases Aen and Rab11 positive (figure 12 a-c). However, many Aen positive structures, especially in the PGCs and the apical portion of the subapical domain do not overlap with Rab11.

The finding that Aen only co-localizes with the small Rab11 positive structures indicates that Aen has no function in the RE itself but might be involved in aspects of membrane traffic during cellularization. The reported presence of Golgi markers in the small Rab11 vesicles (Pelessier et al., 2003) would be in accordance with a potential role of Aen in the secretory pathway downstream of the Golgi apparatus.

3.4.3 Aen shows a partial overlap with the Golgi apparatus marker Lava lamp

Based on the observation of Aen signal in small Rab11 positive structures, we tested whether a similar overlap could be detected also in structures containing the protein Lava lamp (Lva), which is associated with the Golgi apparatus and derived particles (see section 1.1.3; Sisson et al., 2000; Papoulas et al., 2005). As both, the Aen and the Lva antibody were raised in rabbit, the expression of an AenA-eGFP fusion protein with the Gal4-UAS system (see section 3.2.5) was chosen for co-localization studies.
Figure 12: Co-localization of Aen with Recycling Endosome and Golgi. Double immunohistochemical detection of Aen/Rab11 using anti-Aen and anti-Rab11 antibodies. Immunohistochemical detection of Lva in embryos expressing Aen-eGFP using the anti-Lva antibody. a, b, d, e) Cross sections of a stage 5-2 epithelium. Dorsal is up, FC (arrow), a-c) Co-localization of Aen with the Recycling Endosome (RE), RE marked with anti-Rab11. Left column: Aen; middle column: Rab11; right column: overlay, Aen (green), Rab11 (red), a-a") Posterior pole of a stage 5-2 embryo with PGCs (asterisk marks nucleus). While the Aen signal exhibits strong focal enrichments, no Rab11 containing particles are detected. (b-b") Somatic blastoderm at stage 5-2. The Rab11 signal exhibits discrete focal enrichments directly apical of the nuclei (arrowhead). These focal enrichments partially overlap with the structures detected with the anti-Aen antibody. The dashed lines illustrate the depth of the grazing sections. (c-c") Grazing section of the apical cytoplasm. The localization of Aen and Rab11 shows clear differences in many of the large Rab11 positive structures (white arrowhead). Small Rab11 positive structures (black arrowhead) often show Aen signal, d-f) Co-localization of Aen with the Golgi Apparatus. Golgi marked with anti-Iva lamp (Iva). Left column: AenA-eGFP; middle column: Lva; right column: overlay, AenA-eGFP (green), Lva (red), [d-d"] Posterior pole of a stage 5-3 embryo with PGCs (asterisk marks nucleus). The strong AenA positive focal enrichments in the PGCs show no overlap with the Lva signals. (e-e") Somatic blastoderm at stage 5-2. The Lva signal exhibits strong focal enrichments in the basal and lateral cytoplasm. More dispersed signals are detected in the apical cytoplasm. While no enrichment of AenA is detected in the large Lva positive structures (white arrowhead), the two signals overlap in the apical cytoplasm (black arrowhead). [f-f"] In grazing sections of the basolateral cytoplasm an overlap of AenA and Lva positive structures is visible (white arrowhead).
Flies containing the UAS construct coding for AenA-eGFP were crossed to a V3-Gal4 driver-line (generous gift from D. St.Johnston) which induces maternal transcription in the F1 females. The offspring of these females that bear a maternal contribution of AenA-eGFP were then fixed and subjected to a staining with the anti-Lva antibody.

The maternal expression of the AenA-eGFP fusion construct results in a relatively weak eGFP signal (figure 12 d-f). Nevertheless, the AenA-eGFP localization appears to be identical to the localization of endogenous Aen under the chosen conditions. The staining with anti-Lva antibody reveals that differences exist between the localizations of Aen and Lva. The large AenA-eGFP-positive areas detected in the PGCs do not overlap with the Lva-signal. In the somatic blastoderm, Lva is detected in distinct structures in the basal cytoplasm where the AenA-eGFP signal does not exhibit structures of a similar morphology. However in the lateral cytoplasm, an overlap between the Aen and the Lva signal is detected. The Lva detection in these cytoplasmic areas shows smaller areas of signal accumulation, which in many cases are also Aen positive. In the apical cytoplasm, the Lva signal is more dispersed and shows a distribution similar to the AenA-eGFP signal. In summary, Aen is not specifically localized or enriched in the Golgi apparatus. Nevertheless it seems to co-localize with a subset of smaller structures containing the Golgi marker Lva.

3.4.4 Aen partially co-localizes with the Microtubules

Based on the observation that Aen is found in particles potentially associated with the secretory pathway and that it is additionally present at high concentrations in the lateral cytoplasm and the subapical domain, we tested whether Aen positive structures co-localize with the Microtubule (MT) network. During the formation of the polarized epithelial cells of the blastoderm, MT have an important function in the transport of vesicles and organelles such as the lipid
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droplets (LD) (Welte et al., 1998; reviewed in Jäckle and Jahn, 1998). MT bundles extend throughout the cytoplasm vertically to the epithelial surface with the minus-ends directed to the cortex and the plus-ends directed to the yolk (reviewed in Welte, 2004). To investigate if Aen co-localizes with MT, antibody stainings using anti-Aen and anti-(β)Tubulin were carried out with embryo collections of early stages (figure 13). The MT bundles appear as thick filamentous structures stretching from the apical moiety of the cytoplasm to the

Figure 13: Localization of Tubulin and Aen. Double immunohistochemical detection of Aen/β Tubulin using anti-Aen and anti-β Tubulin antibodies. Left column: Aen; middle column: β Tubulin; right column: overlay. Aen (green), β Tubulin (red). a-c) Cross sections of Stage 5-2. a-a') Overview. The asterisk marks the dorsoposterior side of the embryo. b-b') Posterior pole with PGC (asterisk marks nucleus). b) The PGCs exhibit strong Aen signals, often focally enriched, b') Adjacent to the PGC nucleus the centriole is detected as a singular Tubulin rich structure. c-c') Somatic blastoderm. Aen and Tub exhibit strong signals in the apical cytoplasm, extending in between the nuclei towards the FC. c') Enrichments of Aen and Tubulin partially overlap in the apical and lateral cytoplasm (arrowheads). No enrichments of Aen and Tubulin can be found in the FC (arrow). The dashed lines illustrate the approximate depth of the graizing sections. d-e) Grazing sections. d-d') apical cytoplasm of the lateral blastoderm. d') The cytoplasmic distribution of Aen shows local enrichments. d') Also Tubulin is unevenly distributed. Focal enrichments of Tubulin thereby might represent centrioles. e-e') The Aen and Tubulin signals exhibit an overlap in part of the focal enrichments detected (arrowheads). In addition structures with high concentrations of Aeneas but no enrichment in Tubulin occur, e-e') Lateral cytoplasm. e) The cytoplasmic distribution of Aen shows local enrichments (arrowhead). e') Vertical tubulin bundles in the cytoplasm and lining the nuclei appear as punctate structures in the graizing section, e'). The Aen and Tub signals exhibit an overlap in part of these punctae.
cellularization front. The cortical part of the MT network appears as focally condensed structures in the apical cytoplasm. The grazing section (figure 13 e) shows that MT bundles are arranged directly adjacent to the nuclei.

The overlay of the (β)Tubulin and Aen signals obtained in the co-staining shows that the Aen signal partially co-localizes with the Tubulin signal. In cross sections the Aen signal is detected next to the nuclei and the FC, this localization to a great extend reflects the Tubulin localization in filamentous structures found next to the nuclei and the cellularization front. Furthermore, the structures in the apical region of the cells show a partial overlap between Aen and (β)Tubulin. The Aen signal thereby extends further towards the embryonic cortex. In optical sections parallel to the surface of the embryo, the (β)Tubulin and Aen signals show a partial overlap in the apical cytoplasm while in the lateral cytoplasm small, punctate enrichments of Aen are detected that overlap or are directly adjacent to the Tubulin bundles. These findings are in agreement with the observation that Aen co-localizes with particles that are potentially part of the secretory pathway.

3.5 Biochemical interaction studies

As a means to gain further insight into the molecular function of the Aen protein, its ability to bind other, characterized factors was investigated. The basis for candidate selection was the analysis of the aen phenotype, the subcellular localization of the Aen protein and published yeast two-hybrid data from a genome-wide screen (Giot et al., 2003).

The analysis of the aen phenotype had shown that the earliest timepoint of PGC mismigration was at embryonic stage 5-2, when the somatic blastoderm cellularizes. The coincidence of the PGC
mismigration and the cellularization of the somatic PMG cells raised the question which cellular process depends on aen activity: the transmigration activity of the PGCs or the cellularization process preventing the PGC mismigration normally at this stage of development. In the literature only one factor, Slow as molasses (Slam) is described as being required both for cellularization of the somatic cells and for the migration of PGCs in later stages (see section 1.1.4; Stein et al., 2002; Lecuit et al., 2002). Furthermore, the Slam protein is expressed exclusively at the time of cellularization. Therefore, slam appeared to be a good candidate for an interaction partner of Aen.

Shotgun (Shg; Drosophila E-Cadherin; see section 1.1.3; Oda et al., 1994; Tepass et al., 1996) is an important component of the basal and apical adhesive junctions formed during blastoderm cellularization. Mutations of shg affect the adhesion between the somatic blastoderm cells (Tepass et al., 1996). Therefore Shg was tested for biochemical interaction with Aen.

In addition the factor Actin related protein 1 (Arp1) (Fryrberg et al., 1994) was tested, which had been identified as a potential interactor of Aen in a genome wide yeast two-hybrid approach (Giot et al., 2003). Arp1 is a component of the dynactin complex mediating MT based transport.

3.5.1 In vitro interaction studies of Aen with Arp1, Slam and Shg

The method chosen for initial analysis was a pulldown-approach utilizing tagged-Aen protein to bind in vitro translated target proteins labeled with $^{35}$S-cysteine and -methionine. Several constructs coding for Aen fusion-proteins with solubility promoting tags (reviewed in Esposito and Chatterjee, 2006) were generated. The bacterial expression levels and solubility of Maltose Binding Protein-AenA, Thioredoxin-AenA, N-utilizing substance-AenA and Glutathione-S-
Transferase-AenA (GST-AenA) were tested. In these tests only the GST-AenA fusion exhibited expression rates and solubility sufficient for medium scale expression and purification.
Results

The Slam protein as well as the protein Luciferase, used as an internal negative control in this experiment, were generated in a combined \textit{in vitro} transcription/translation reaction (section 2.3.8) and applied on a resin loaded with GST-AenA as well as mock resin loaded with GST without fusion portion. After extensive washing, the proteins bound to the resin were subjected to SDS-PAGE and exposed to image-plates. The analysis (figure 14a) showed that the Slam protein binds to the resin preloaded with Gst-AenA while it does not bind to the mock-resin preloaded with GST. In contrast, Luciferase used as a control shows no interaction with the GST-AenA loaded and the mock resin. This result gave a first indication that a specific interaction might exist between Slam and AenA.

Based on the cytoplasmic localization of Aen an interaction of Aen with the extracellular and transmembrane domains of Shg could be excluded. Therefore, only the intracellular domain of Shg was tested for a potential interaction with AenA. In order to express the cytoplasmic tail, the sequences encoding the extracellular and transmembrane region were removed by PCR and an artificial start codon was introduced into the cDNA. The \textit{in vitro} translation resulted in a protein with the expected size of 16 kDa as deduced from the aminoacid sequence of the cytoplasmic tail of Shg. Even though the protein amount obtained \textit{in vitro} was comparably low, the results show that no direct interaction exists between the intracellular domain of Shg and GST-AenA (figure 14a).

To investigate if Arp1 is able to bind to purified recombinant AenA, it was labeled and subjected to the GST-pulldown assay as described above. The result of the analysis showed that Arp1 binds to GST-AenA loaded resin and does not bind to the mock-resin (figure 14a). To further characterize the interaction between AenA and Arp1, the proteins CapZ and Arp11 were tested in the GST-Pulldown analysis as
well. Both proteins are subunits of the dynactin complex and direct binding partners of the Arp1 protein (reviewed in Schroer 2004, Imai et al., 2006). The results of the pulldown assay show, that neither CapZ nor Arp11 exhibit specific binding affinity to AenA-GST. These findings, which are in good agreement with the results of the yeast two-hybrid approach (Giot et al., 2003), indicate that AenA might be associated to Arp1 as a part of the dynactin complex.

In summary, the GST-pulldown experiments revealed an interaction of GST-AenA with in vitro translated Slam and Arp1, but no interaction with CapZ, Arp11, the cytoplasmic tail of Shotgun or the control protein Luciferase. To confirm the obtained interactions of AenA with Slam and Arp1, independent protein-protein interaction experiments were performed.

3.5.2 AenA and AenB interact with Arp1 and Slam in co-immunoprecipitation experiments

The in vitro data provided initial evidence for the interaction between Aen and Arp1. In order to achieve an independent control of these experimental results, co-immunoprecipitation experiments were conducted. To allow for an efficient immunoprecipitation and detection of the proteins in the subsequent western blot, AenA, Slam and Arp1 were differentially tagged. Vectors were generated for the expression of 3xFlag-Aen and 10xMyc-Arp1, tagged at their N-termini, under the control of an Actin promoter. Drosophila Kc167-cells (Cherbas et al., 1977) were transfected and harvested after 4 days. An antibody against the Myc-Tag was used for immunoprecipitation; Western Blot detection of co-precipitated proteins was carried out using an anti-Flag antibody.

The results of the Co-IP (figure 14b) show that the Flag-AenA protein is co-precipitated with the Myc-Arp1 protein. Thus the interaction of Aen
with Arp1 could be shown by GST-pulldown experiments and in co-immunoprecipitation and are in good agreement with the results of the yeast two-hybrid experiments (Giot et al., 2003).

To restrict the region of Aen required for the interaction with Arp1, the interaction of Arp1 with the N-terminally shortened Aen variant AenB was analyzed in co-immunoprecipitation experiments as well (figure 14b). Since AenB interacts with Arp1 in a comparable strength the interaction domain could be restricted to the region between aa 207 and 546 of AenA.

The interaction between Aen and Slam, which had been shown in the in vitro pulldown assay, was likewise tested by means of co-immunoprecipitation of tagged version of full length AenA and Slam. The expression strength of a 10xMyc-fusion of full length Slam, a protein of 1291 aa, was found insufficient for co-immunoprecipitation experiments. Therefore, two constructs coding for truncated Slam-versions representing the N-terminal 834 aa and the C-terminal 493 aa of the protein both tagged with a 10xMyc-tag were generated (figure 14d).

The results of these experiments are displayed in figure 14. AenA shows interaction with the N- and C-terminal part of Slam, indicating that either the Slam protein contains two regions interacting with AenA or that the interacting domain extends over the region at the boundary between the two Slam fragments expressed.

To restrict the interacting region of Aen the AenB protein was tested for binding to the N- and C-terminal truncated Slam variants as described for AenA. The experiment revealed an identical interaction of AenB with two Slam variants as shown for AenA, indicating that the Slam interacting region of Aen must be located in the overlapping
Results

Sequence between both Aen variants, mediating the interaction with at least two regions of the Slam protein.

3.5.3 Mapping of the Aen interacting domains of Slam

The co-immunoprecipitation experiments with Slam, AenA and AenB had revealed that both the N- and the C-terminal regions of Slam interact with Aen. To restrict these domains further, a series of 11 overlapping Slam fragments were cloned (figure 15a), in vitro translated and tested for interaction with AenA using the GST-pulldown approach described in section 3.5.1. The fragments had an average length of about 200 aa covering the entire sequence of the Slam protein. Such a systematic approach was required as no protein domains are identified in the Slam protein and no data are available that help to refer the biological functions of Slam to specific regions of the protein (Stein et al., 2002; Lecuit et al., 2002; SMART). The
systematic approach reveals that two regions of Slam interact with GST-AenA in vitro. A strong interaction is detected in a region located in the C-terminal half covering the amino acid domain ranging from aa 705 to aa 1106, represented by the fragments 7, 8 and 9, and a weaker one in the N-terminal half a region ranging from aa 300 to aa 612, represented by the fragments 4 and 5. So both the co-immunoprecipitation experiments and the GST-pulldown experiment identified two domains of Slam, which interact with Aen.

3.6 Co-localization of Aen with its interaction partners

The biochemical interaction assays had shown, that Aen binds Arp1 and Slam both in in vitro Gst-pulldown assays and in vivo as shown by co-immunoprecipitation experiments after transient transfection of cultured cells. To determine whether the interactions of Aen with Arp1 and Slam could occur in vivo in the PGCs or the somatic blastoderm and could therefore be of biological relevance, a series of double stainings was conducted. A co-localization of Aen with its potential interacting partners would provide an important indication for the biological relevance of the interactions.

3.6.1 Aen co-localizes with Dynein Heavy Chain

Since no specific antiserum against Drosophila Arp87 is available, the subcellular localization of Arp1 was determined indirectly. For this purpose, an antibody detecting Dynein Heavy Chain (Dhc), the motor protein which acts in conjunction with the dynactin complex (section 1.1.1; reviewed in Schroer 2004) was applied in double stainings with anti-Aen. A high degree of overlap between Aen and Dhc exists in the somatic blastoderm (figure 16a-d). However, in the PGCs no accumulations of Dhc analogous to the strong focal enrichments of Aen are observed. In cross sections of somatic cells of a stage 5-1 embryo, the enrichments of the Aen staining basally of the nuclei (see
also figure 11) are also observed for Dhc (figure 16a, arrowhead). At stage 5-3, the focal enrichments in the apical cytoplasm and the basal cytoplasmic distributions show a strong overlap of Aen and Dhc (figure 16c). Minor differences in localization are visible in the region of the FC where the Dhc concentration appears to be elevated in respect to Aen. The co-localization of Aen and Dhc provides a strong indication that Aen and Arp1, which belongs to the same protein complex as Dhc, co-localize as well.

3.6.2 Aen co-localizes partially with Slam

Based on the fact that both Aen and Slam antibodies were generated in rabbits, it was necessary to use a Slam-HA fusion protein (Lecuit et al., 2002) that was expressed in early embryos using the maternal V3-GAL4 driver line (section 3.2.5). The Slam fusion protein is localized identical to the endogenous Slam protein (Lecuit et al., 2002, data not shown).

The signals obtained in the antibody staining of fixed embryos show that Aen and Slam-HA, even though exhibiting differences in their apicobasal distribution, overlap in structures with high Slam-HA content. The Slam-HA protein is strongly enriched in the FC and the basolateral membrane adjacent to it and shows only a minor cytoplasmic signal. In contrast, the highest Aen concentrations are detected in the apical cytoplasm. Optical sections of the basolateral membrane and the FC, that runs parallel to the embryonic surface, show that both proteins, Aen and Slam-HA, exhibit overlapping enrichments in the lateral cytoplasm, adjacent to the basolateral membrane and the FC. Aen is not excluded from the Slam-HA positive structure marking the leading front of the FC, however the highest concentrations of Aen are found in the cytoplasmic stalks in regions directly adjacent to areas with strong Slam signal.
Figure 16: Colocalization of Aen with interacting partners. Double immunohistochemical detection of Aen/Dynein heavy chain (Dhc) and Aen/Slam-HA using anti-Aen, anti-Dhc and anti-HA antibodies. a-d) Co-localization of Aen with Dhc, a main component of the dynein-dynactin transport complex. Left column: Aen; middle column: Dhc; right column: overlay. Aen (green), Dhc (red). a-a') Cross section of a stage 5-1 epithelium. Forming FC (arrow). The enrichment basally of the nuclei detected in the Aen staining is also observed for Dhc. In addition, Dhc exhibits also apical enrichments. b-b') Posterior pole of a stage 5-2 embryo with PGCs (asterisk marks nucleus). While the Aen signal exhibits strong focal enrichments, the Dhc signal in the PGCs is only very weak. Only a minority of the Aen enrichments bears a corresponding Dhc signal. c-c') Somatic blastoderm at stage 5-3. The focal enrichments in the apical cytoplasm and the basal cytoplasmic distributions show a strong overlap of Aen and Dhc. Minor differences in localization are visible in the region of the FC (arrow) where the Dhc concentration appears to be elevated in respect to Aen. The dashed lines illustrate the depth of the grazing sections. d-d') Grazing section of the lateral cytoplasm. The localization of Aen and Dhc exhibits a high degree of overlap. e-g) Co-localization of Aen with Slam-HA. Left column: Aen; middle column: Slam-HA; right column: overlay. Aen (green), Slam-HA (red). e)
The differences in apicobasal distribution of Aen and Slam-HA do not support the results of the biochemical interaction studies, however the partial overlap found in the basolateral membrane and the FC indicates that a subpopulation of Aen can interact with Slam in vivo.

3.7 Genetic interaction studies of aen with slam

To investigate whether the biochemical interaction and co-localization of Aen and Slam reflect a biologically relevant interaction, changes in cellularization velocity of aen mutant embryos were assayed. The genetic interaction of aen and slam during the cellularization of the somatic blastoderm cells was tested by analysis of the blastoderm morphology. As a third assay, the occurrence of cuticle phenotypes later in development was analyzed.

3.7.1 Genetic interaction of aen and slam during cellularization

Earlier studies of slam (Stein et al., 2002) had shown that zygotic null mutant embryos that still bear half of the normal maternal contribution of slam mRNA exhibit only mild developmental defects characterized by PGC mismigration from stage 10 onwards (see section 1.1.4). When additionally the maternal contribution of slam mRNA was removed, it was found that the resulting maternal-zygotic null mutant embryos exhibited strong defects in blastoderm cellularization during stage 5, resulting in a preterm stop of FC invagination and extensive cell death in the incompletely cellularized epithelium. It was found that the localization of the proteins D-PATJ (section 1.1.3; Bhat et al., 1999; Pielage et al., 2003), a component of the FC, and Neurotactin (Nrt; section 1.1.3; Hortsch et al., 1990), an adhesion factor localized to the
Results

Apicolateral membranes, is altered in the affected epithelia (Stein et al., 2002; Lecuit et al., 2002). While D-PATJ positive membrane domains broadened and fuzzied, Nrt, in addition to its enrichment in the apicolateral membranes, is found ectopically enriched in the apical cytoplasm (Lecuit et al., 2002). This localization defect might be the consequence of an impairment of vesicle-integration into the apicolateral domain of the plasma membrane (PM) (Lecuit et al., 2002).

To characterize a potential genetic interaction between aen and slam it was analyzed if the loss of aen activity increases the strength of the zygotic slam mutant phenotype. The rationale of this experiment was that the absence of Aen would reduce the remaining zygotic Slam activity thereby leading to the occurrence of cellularization defects even in the presence of maternal slam mRNA (figure 18) if the potential interaction between Aen and Slam is important for slam activity. For this purpose, the aen\textsuperscript{2} mutation was crossed into the genetic background of both the slam\textsuperscript{1} mutation and the slam deficiency (Df(2L)Exel6016, covering cytogenetic bands 26C1-D1). Crossing of these fly stocks resulted in offspring, bearing half of the maternal contribution (for details see Materials and Methods section 2.2.3). As a control, the cellularization in slam single mutant embryos (for details see Materials and Methods section 2.2.3), and aen\textsuperscript{2} embryos was investigated. For the analysis, embryos of wt, the aen\textsuperscript{2} and slam single mutants and slam-aen double mutants were fixed and stained using antibodies detecting the D-PATJ and Nrt, which both are good markers for the morphology of the forming epithelium and in addition might exhibit localization defects as shown in the maternal zygotic slam mutant (Stein et al., 2002; Lecuit et al., 2002).
The analysis of the single mutants gave the following picture: In slam single mutant embryos, 5.4% of the embryos show strong defects in blastoderm cellularization, characterized by a local reduction of the invagination depth of the FC, covering one of the poles and...
extending towards the median of the embryo. A clear majority of 94.6% shows no such defects.

In homozygous aen\(^2\) mutant embryos, the penetrance of cellularization defects is elevated in comparison to slam/deficiency mutants and wt. 23% of the embryos exhibit a partial failure of blastoderm cellularization (figure 17c). Interestingly, a partial mislocalization of Nrt to the apical cytoplasm, similar to the defect reported for maternal-zygotic slam mutants, can be observed in some cases. With a low penetrance of less than 5% the formation of multinucleate cells is observed (17c'). Also severe disruption of the palisadal morphology of the somatic epithelium is observed with a low penetrance. In these cases, Nrt mislocalization in punctate enrichments in the lateral membranes occurs (figure 17c''). However, blastoderm cellularization was unaffected in 77% of the embryos.

After the characterization of the single mutants, the phenotype of the slam-aen double mutant was analyzed. It was expected that a biologically relevant interaction between the two factors would result in a strong induction of the cellularization- or PGC migration phenotypes observed in the single mutants.

It was found that the majority of offspring originating from slam heterozygous mothers exhibited a variety of cellularization defects. With almost full penetrance, a filling of the micropyle, a small appendix at the anterior pole of the vitellin membrane surrounding the embryo (figure 2a), with cytoplasm was observed. Also nuclei were frequently found to invade the micropyle. Strong cellularization defects which included asynchrony of cellularization were observed in 45% of the embryos. In these embryos, a local reduction of invagination depth of the FC occurs. Furthermore, the incomplete formation of the FC and the formation of multinucleate cells were found some in regions of the embryos, resulting in a local complete
block of the cellularization. In most embryos analyzed, the strongest defects were found at the embryonic poles.

These results indicate that the phenotype found in slam single mutants is strongly enhanced in conjunction with the aen\textsuperscript{2} mutation, adopting the characteristics of the phenotype found in the maternal/zygotic slam null mutants (Stein et al., 2002).

3.7.2 Cellularization kinetics in aen mutants

The analysis of slam (Lecuit et al., 2002) had shown that the kinetics of FC invagination is altered by RNAi-mediated slam knockdown. The membrane invagination speed was found reduced by half in slam knockdown as compared to wt. As the biochemical and genetic interaction and co-localization studies suggests a functional cooperation of Aen and Slam during cellularization, the speed of membrane insertion was investigated in aen mutants during late cellularization phase. For this purpose, embryos were dechorionated and mounted with halocarbon oil. Recordings were taken with phase contrast microscopy, which allows the visualization of the highly light diffracting FC in cross sections of the developing epithelium. The vitellin membrane, which marks the outer boundary of the embryo, was chosen as a reference structure.

The recordings of the fast phase of FC invagination in aen mutant embryos show that the kinetics of membrane invagination is not significantly altered in aen mutants (figure 18a and b). This finding might be a result of the low penetrance of cellularization defects in aen\textsuperscript{2} mutant embryos.
3.7.3 Cuticle phenotypes in single and double mutants of aen and slam

In previous functional studies of factors involved in the establishment of cell polarity and adhesion, the frequent occurrence of cuticle malformation had been observed (Wieschaus, 1984; Müller and Wieschaus, 1996; Knust et al., 1996). The cuticle is the exoskeleton of Drosophila larval and adult stages. The larval cuticle, which is secreted by epidermal cells during late embryogenesis from late stage 15 onwards, fully encloses the embryo and exhibits a characteristic structured form that reflects the identity of the embryonic segments by the pattern of the denticle belts, anterior structures, the head skeleton including the mouthparts, as well as posterior structures with the filzkörper (reviewed in Martinez Arias, 1993). Cuticle preparations have been applied for the analysis of mutant phenotypes in body axis formation and segmentation (e.g. Fröhnhofer and Nüsslein-Volhard 1986). However, cuticle preparations were also used to identify defective cellularization processes. Mutants affecting membrane...
polarity or adhesion (e.g. bazooka (Müller and Wieschaus, 1996, Knust et al., 1996) and crumbs (Tepass et al., 1990; Wodarz et al., 1995; Wodarz et al., 1993; Jürgens et al., 1994); see also section 1.1.3), result in extensive cell death and the disruption of the primary epithelium shortly after cellularization. Therefore, only parts of the surviving tissues secrete cuticle later in development. These cuticles often exhibit partial reductions in the structures described above, contain holes (bazooka (Wieschaus, 1984)) or are completely incoherent and unstructured (crumbs, stardust (Tepass et al., 1990; Müller and Wieschaus, 1996)). These examples show that strong defects in cellularization often are distinctly reflected in the cuticle morphology.

To quantify penetrance of the cellularization phenotypes observed in the slam and aen\(^2\) single mutants as well as in the slam-aen double mutants, the cuticle phenotypes found in these three genotypes were examined and quantified.

For the analysis three categories of cuticle phenotypes were defined. The first category was characterized by the lack of cuticle, which could either reflect extreme cellularization phenotypes resulting in the formation of no epidermal cells whatsoever or the death of formed tissues prior to the secretion of the cuticle. Unfortunately these phenotypes could not be distinguished from unfertilized eggs. Therefore this phenotypic class was excluded from the consecutive statistical analysis. Embryos that exhibited a cuticle lacking all three signs of defined morphology such as denticle belts, headskeleton and filzkörper were grouped in the second category. The third category encompassed embryos that showed either a wildtype cuticle or an incomplete cuticle bearing at least one of the morphologically defined traits.

The analysis of the cuticle preparations yielded the following results (see figure 18). In slam single mutants 90.5 % of unhatched embryos
with cuticle exhibited signs of morphological differentiation. In most of the cases all three morphological criteria, head skeleton, filzkörper and denticles were observed. Only 9.5 % of the embryos showed a completely undifferentiated cuticle.

In aen\(^2\) single mutants a similar observation was made. 90.0 % of the embryos showed a differentiated cuticle while an undifferentiated cuticle was only found in 10.0 % of the cases.

In double mutants the rate of embryos with an undifferentiated cuticle was much higher than in the collections of the two single mutant genotypes: 47.8 % of the embryos belonged to this morphological category. 52.2 % of the embryos showed partially or fully differentiated cuticle morphology. In this group, holes in the cuticle and partial reductions of the head skeleton were frequently observed.

These results show, that aen and slam single and double mutants exhibit cuticle malformations characteristic for cellularization defects. Furthermore, a dramatic increase in phenotypic strength induced by the loss of aen gene activity in slam mutants strongly suggests that a functional interaction exists between the two factors.
4 Discussion

In the present study, the role of the recently identified factor Aeneas (Aen) was characterized as a modulator of primordial germ cell (PGC) migration (section 3.2; Molitor, 2002). Furthermore, Aen is required for the establishment and maintenance of the epithelial integrity during cellularization of Drosophila melanogaster (section 3.7.1). The Aen protein represents a new family of proteins with homologs in a broad range of organisms (section 3.1; figure 6). In aen mutant embryos untimely transmigration of the PGCs occurs with a high frequency during the fast phase of cellularization of the somatic cells of the blastoderm embryo. Aen is expressed both in PGCs as well as in the somatic cells in which it specifically co-localizes with structures of the secretory pathway during cellularization (section 3.4; figures 12 and 13). The investigation of the influence of the aen mutation on the PGCs migration and the cellularization of the underlying epithelium revealed that a functional interplay exists between PGC migration and cytokinesis of the somatic blastoderm. Furthermore, I have identified two factors that interact with the Aen protein, which represent a functional link of Aen activity and the observed cellularization defects. The interaction of Aen with these two factors is shown by two independent biochemical techniques, GST-pulldown assays and co-immunoprecipitation (section 3.5; figure 14). Furthermore, the functional significance of the interaction could be shown for one of the interacting factors by a genetic interaction of the two mutants in vivo (section 3.7; figures 17 and 19). The results obtained in the work presented here indicate that aen loss of function (LOF) affects subcellular protein transport mediated by the Dynactin complex in the somatic blastoderm cells. This transport is involved in
the polarized insertion of membranes in the plasma membrane (PM) during the cellularization of the somatic epithelium (Lecuit and Wieschaus, 2000; Sisson et al., 2000; Papoulas et al., 2005). aen LOF results in a partial malformation of the somatic blastoderm (section 3.7.1; figure 17) which might reduce its resistance to the transmigration of PGCs. Furthermore, biochemical interaction and functional genetic analysis reveal that Slow as molasses (Slam) represents one of the Aen interacting activities required for cellularization of the somatic epithelium (sections 3.5 and 3.7).

4.1 Correlation between aen expression, phenotype and localization

4.1.1 aen expression

The expression and localization data presented in this work show that aen expression initiates in the nurse cells of the ovariole after oogenesis stage 8 and that aen expression is restricted to the germline during oogenesis. This finding shows that aen is neither required during early oogenesis nor in the somatic parts of egg follicles (section 3.3.1; figure 9). After transport of the aen mRNAs into the oocyte as a maternal contribution, aen transcripts can be detected ubiquitously distributed until embryonic stage 4 (Molitor, 2002). At stage 5, aen transcripts are degraded in the somatic blastoderm but not the PGCs. Weak ubiquitous zygotic expression is detectable during the following stages in in situ hybridisations and by RT-PCR (Molitor, 2002).

The developmental western blot analysis shows, that Aen protein is first detectable during oogenesis as shown for the transcripts (section 3.3.3; figure 9). In contrast to the transcript localization during early embryogenesis however, Aen protein is not restricted to the PGCs as revealed by whole mount antibody stainings (section 3.4; figures 10 and 11). Instead, Aen can be found ubiquitously during early embryogenesis and also at later stages both in the somatic cells and
in the PGCs. While the specific stabilisation of the aen transcripts in the PGCs is indicative of a PGC specific function, the ubiquitous localization of the Aen protein suggests a general role in all cells.

4.1.2 aen phenotype

The aen maternal and zygotic LOF mutant phenotype is characterized by a premature transmigration of PGCs through the underlying somatic blastoderm during embryonic stages 5 to 9 (section 3.2; Molitor, 2002). This phenotype is clearly caused by the loss of aen activity as it can be rescued by introducing a genomic rescue fragment. Furthermore, weak cellularization defects are observed. In extreme cases the cellularization fails in a region close to the poles (section 3.7.1; figure 17c). In those cases, a mislocalization of Neurotactin (Nrt) (Hortsch et al., 1990) can be observed close to the cell cortex in the apical cytoplasm. Later defects have not been observed in aen LOF mutants. The aen mutation is viable, allowing the keeping of mutant flies as a homozygous stock.

4.1.3 Subcellular localization of Aen in the somatic blastoderm

The analysis of the subcellular localization in the somatic blastoderm has shown that Aen, in addition to its cytoplasmic distribution, is localized to small granular structures that could represent vesicles involved in transport (section 3.4; figure 11). To approve this model co-staining experiments with markers known to be involved in the vesicle transport required for cellularization were performed.

These experiments revealed that Aen co-localizes partially with the Microtubules (MT) (figure 13 d-f) and with Rab11 (figure 12 a-c), a marker for vesicles involved vesicle transport and for the recycling endosome (RE) (Ullrich et al., 1996; Sasamura et al., 1997; Pelissier et al., 2003). However, the co-staining of Aen and Rab11 is specific for
small particles localized to the apical cytoplasm and is not observed
the RE directly (figure 12 a-c). As these vesicles co-express markers of
the Golgi apparatus as well (Pelissier et al., 2003), co-localization with
Lava lamp (Lva) was investigated, which is required for intracellular
transport of Golgi vesicles during blastoderm cellularization (Sisson et
al., 2000; Papoulas et al., 2005). However, similar to the situation with
Rab11, the co-localization is restricted to small structures in the lateral
and apical cytoplasm and cannot be detected in the larger Golgi
particles basal of the nuclei (figure 12 d-f). Taken together, Aen co-
localizes with small vesicular structures that are Rab11 or Lva positive
and show at least a partial overlap with the MT network indicating that
Aen is positioned on vesicles of the secretory pathway (reviewed in
Strickland and Burgess, 2004) presumably downstream of the Golgi
apparatus.

4.1.4 Functional correlation of aen with rab11 mutants

Rab11 controls trafficking through the RE in mammals and Drosophila
(Dollar et al., 2002; Ulrich et al., 1996). Injection of a dominant negative
protein variant inhibits membrane invagination at the slow and fast
phase of cellularization (Pelissier et al., 2003). In Drosophila,
hypomorphic mutations of Rab11 cause furrow canal (FC) malformation leading to the formation of multinucleate cells (see
section 1.1.3; Pelissier et al., 2003). In wildtype, Rab11 co-localizes with
Nrt in vesicular structures, but not in the RE itself (Pelissier et al., 2003).
However, in mutants, which affect vesicle budding from the RE,
accumulations of Nrt in this organelle can be found (Pelissier et al.,
2003). These results indicate that the transport of Nrt-positive vesicles
from the RE is dependent on the function of Rab11.

The inhibition of membrane invagination found in Rab11 LOF
resembles the cellularization defects in strongly affected blastoderm
regions found in aen LOF mutants (figure 18). Furthermore, Nrt is found
ectopically enriched in the apical cytoplasm in aen mutant embryos (figure 18; Pelissier et al., 2003), a phenotype similar to the rab11 mutant (Pelissier et al., 2003). Despite these similarities, no indications exist in recent publications that Rab11 function influences PGC migration in Drosophila (Pelissier et al., 2003, Riggs et al., 2003).

The co-localization of Aen with Rab11 and the similarity in aen and rab11 LOF phenotypes suggest that Aen function is related to intracellular transport. This hypothesis is further supported by the similarities between Aen and Nrt localization. Nrt mislocalization in aen mutants indicates that Aen function might be involved in transport of vesicles towards the newly forming membrane. However, a direct and exclusive functional interaction of Aen proteins with Rab11 is unlikely due to the differences observed in FC formation and PGC migration.

4.1.5 Functional correlation of aen with the lva LOF

Lva is required for intracellular transport of Golgi vesicles during blastoderm cellularization (Sisson et al., 2000; Papoulas et al., 2005). Its functional inhibition causes phenotypes similar to the ones observed in Rab11 (Papoulas et al., 2005). Injection of antibodies against Lva or single Lva protein domains causes a reduction in membrane invagination speed and discontinuous FC formation (Sisson et al., 2000). A similar effect can be caused by injection of brefeldinA, which inhibits the transport of Golgi vesicles, and colcemid, which depolymerises MT (Sisson et al., 2000; Lecuit and Wieschaus, 2000). The hypothesis of the functional interconnection between MT based intracellular transport of Golgi vesicles and blastoderm cellularization has been substantiated by the finding that Lva binds the Dynactin complex, which acts as an adjunct of motor protein complexes (Foe et al., 1993; Lecuit and Wieschaus, 2000; Sisson et al., 2000; Papoulas et al., 2005). In good agreement with these findings is the mislocalization of Lva and Nrt caused by depolymerization of MT, which leads to the
accumulation of Lva- and Nrt- positive structures close to the location of the highest Golgi density, in the basal cytoplasm (Lecuit and Wieschaus, 2000; Sisson et al., 2000). The cellularization phenotypes found in aen mutants exhibit similarities with the effects of anti-Lva, brefeldin or colcemid injection. However, no multinucleate cells have been observed in aen mutants, which is in contrast to the observations made after injection of dominant negative Lva (figure 19; Sisson et al., 2000). Furthermore, Nrt is enriched in the apical cytoplasm in aen mutants, similar to rab11 mutants and to embryos with depolymerised MT (Pelissier et al., 2003; Lecuit and Wieschaus, 2000). In contrast, Lva LOF probably results in an Nrt accumulation next to the Golgi in the basal cytoplasm (see figure 18; Sisson et al., 2000; Lecuit and Wieschaus, 2000). Similar to Rab11, the studies of Lva function do not indicate its involvement in PGC migration; however, this process might not have been observed due to the experimental setup (Sisson et al., 2000; Papoulas et al., 2005).

Taken together, the co-localization of Aen with MT and with Rab11 and Lva in small vesicles in the apical cytoplasm and the similarity in the LOF phenotypes suggest that aen function is related to intracellular transport. The apical enrichment of Nrt in aen mutants suggests, that Aen does not directly influence the transport from the Golgi to the apical cytoplasm. This finding does not support the hypothesis of a direct interaction and functional relation between the Aen proteins and Lva.

4.1.6 Subcellular localization of Aen in the PGCs

The localization of Aen protein in the PGCs differs considerably from the somatic tissues. In PGCs Aen is enriched in large granular structures that are neither Golgi nor RE and show no visible overlap with the MT. So the involvement of Aen proteins in intracellular transport in the PGCs is not probable. Based on their morphology, Aen-positive
structures observed in these cells could represent polar granules (PGs) (see section 1.1.2; Lehmann and Nüsslein-Volhard, 1986; Hay et al., 1988).

However, characterized factors involved in PG assembly and function, including oskar (osk) and vasa (vas), are deposited specifically at the posterior pole of the oocyte already during oogenesis (see section 1.1.1; Lehmann and Nüsslein-Volhard, 1986; Hay et al., 1988, 1990). During early embryogenesis the mRNAs of these factors as well as the respective proteins are localized to the pole plasm at the posterior pole of the embryo (see section 1.1.2; Lehmann and Nüsslein-Volhard, 1986; Hay et al., 1988, 1990). In contrast, aen transcripts are ubiquitously localized in the oocyte and during embryonic stages 1-4. During stage 5, aen transcripts are restricted to the posterior pole including the PGCs (section 1.3; figure 9; Molitor 2002) but Aen protein is ubiquitously detectable during early embryogenesis (figure 10). Furthermore, there are clear functional differences. Components of the PG are involved several critical developmental processes during early embryogenesis (section 1.1.2; Lehmann and Nüsslein-Volhard, 1986). In mutants of osk, the assembly of the pole plasm is abolished and no PGCs are formed (Lehmann and Nüsslein-Volhard, 1986). In mutants of vas posterior morphogenesis is affected (Hay et al., 1988). In mutants of polar granule component (pgc), the transcriptional repression in the PGCs is abolished resulting in dedifferentiation and loss of PGCs (Martinho et al., 2004; Deshpande et al., 2004). In aen mutants, PGC formation and posterior morphogenesis are not affected (PGC formation: figure 7; posterior morphogenesis: as inferred from cuticle morphology, data not shown). Additionally aen mutants are fertile, which strongly indicates that PGC maintenance is unaffected (section 3.2.1). Conversely, mutants affecting PG factors in which PGCs are still formed show no indication of PGC mismigration.
phenotypes during blastoderm stages (reviewed in Molyneaux and Wylie, 2004).

The lack of overlap between the mutant phenotypes of aen and PG components indicates that the functional role of Aen is unrelated to the functions of PG components. However, aen mutants exhibit phenotypes that affect PGC development. Therefore, a PGC autonomous function of aen might nevertheless exist.

4.1.7 Tissue specific requirement of Aen

The premature transmigration of PGCs through the underlying epithelial sheet in aen mutant embryos could be explained by a cell autonomous function of Aen in the PGCs as its transcript is stabilized and the protein is deposited in high amounts in this tissue. Such a PGC autonomous role of Aen could be proven directly by a PGC specific rescue of the aen mutant phenotype using the GAL4-UAS system. However, during early embryogenesis this experimental approach bears several principal difficulties, which cannot be overcome. After their formation, the PGCs are transcriptionally silent, rendering GAL4-UAS induced PGC-specific zygotic expression impossible up to stage 9-10 (section 1.1.2, reviewed in Williamson and Lehmann, 1996; Van Doren et al., 1998). Maternal GAL4-UAS mediated expression would be sufficient to include aen transcripts into the PGCs, similar to the wildtype situation (Molitor, 2002). However no PGC specificity of aen transcripts could be accomplished with this approach. Several maternal transcripts, including nanos (nos), are restricted to the PGCs by active degradation in the somatic tissues (Wang and Lehmann, 1991; Ephrussi et al., 1991; Lehmann and Nüsslein-Volhard, 1991, Bashirullah et al., 1999). This restriction can be induced in other genes by fusion to the nos-3'-UTR coding region (Bashirullah et al., 1999). However, this system is not efficient enough to suppress the translation
in posterior somatic cells completely and does therefore not allow for a clear distinction of PGC specific effects (Bashirullah et al., 1999).

An experimental alternative would be the transplantation of wt PGCs on aen mutant embryos. In this experiment, PGCs are extracted from fluorescein labelled donor embryos of stage 3-4 and injected into the periembryonic space at the posterior pole of a recipient embryo (Jaglarz and Howard, 1994). This technique has been utilized to investigate PGC migratory behaviour during stage 10 (Jaglarz and Howard, 1994; Jaglarz and Howard, 1995, Kunwar et al., 2003), however several problems make this solution inapplicable for the investigation of the aen phenotype. The retrieval and transplantation is a strong disturbance of the PGC surface morphology. The narrow time window between PGC injection and phenotypic observation would not allow for an appropriate recovery phase. Accordingly, non-specific effects of this physical agitation, which might influence the experimental result, could not be excluded. These experimental limitations impair the direct investigation of PGC autonomous effects of Aen.

An indirect solution for this problem is offered by the possibility of a specific zygotic rescue of the aen mutant phenotype. If Aen would be required cell autonomously in the PGCs, the expression of Aen in the somatic blastoderm should have no effect on PGC mismigration. A specific somatic expression could be obtained using the Nullo-GAL4 driver line that enables specific expression in somatic cells of the blastoderm embryo during cellularization (Kunwar et al., 2003). Since this soma specific expression of Aen results in a partial rescue of the PGC transmigration phenotype in aen^2 mutants (section 3.1.4) an exclusive requirement of aen in the PGCs can be ruled out.

Even though this finding does not directly disprove the hypothesis of an additional PGC autonomous function of Aen, it strongly suggests
that the cause of PGC mismigration is to be found in somatic blastoderm.

Despite the finding that Aen is required in the somatic blastoderm and that it partially co-localizes with components of the secretory pathway, the integration of Aen in a network of interaction partners with characterized functions was necessary for its functional characterization.

### 4.2 The interaction between Arp1 and Aen

The first indication towards potential interacting partners of Aen was gained from a genome wide yeast two-hybrid (Y2H) approach. In this high throughput screen, full-length cDNAs of annotated *Drosophila* genes were utilized to generate an interaction map of the whole *Drosophila* proteome (Giot et al., 2003). Based on the technical limitations of a high throughput screen, a critical evaluation of the identified candidates on the basis of their expression profiles and subcellular localizations is required. Among the several factors that showed interaction with Aen, the interaction with the highest confidence was found between Aen and Arp1 (Fryrberg et al., 1994).

In the light of the previous observations, which had established a possible link between Aen function and intracellular transport, Arp1 appeared to be a promising candidate for further analysis. Therefore the interaction between AenA and Arp1 was tested by means of GST-Pulldown and Co-IP experiments. The results of both experimental designs approve the Y2H interaction of Aen with Arp1 (section 3.5.1; figure 14 a and b).
4.2.1 Function of Arp1

Arp1 is a major constituent of the Dynactin complex, forming a short filament at the base of the complex (Schafer et al., 1994; Bingham and Schroer, 1999; Imai et al., 2006). Dynactin is involved in multiple transport processes including dynamic functions, like Golgi dynamics and fast axonal transport, and static functions, including the organization of mitotic spindles, chromosome segregation and nuclear positioning (reviewed in Karki and Holzbaur, 1999; Schroer, 2004; Sharp et al., 2000; Whited et al., 2004; Karess, 2005; Kim et al., 2007). In the static functions of Dynactin, its direct interaction with MT via the Glued subunit plays an essential role (Kim et al., 2007). Dynactin has been shown to be present at the attachment site of MT to the PM and to kinetochores during mitosis and has been hypothesized to organize signaling molecules by tethering them to the MT (Geiser et al., 1997; Schroer 2004). Dynactin interacts with the motor protein complexes Dynein and Kinesin2 and is involved in their function (reviewed in Schroer, 2004; Deakon et al., 2003). The Arp1 filament has been shown to bind proteins localized at vesicle surfaces, including β III Spectrin, a component of Golgi membranes, and COPII, which is involved in the vesicle transport from ER to Golgi (Beck et al., 1994; Holleran et al., 2001; Watson et al., 2005; COPII function reviewed in Sato et al., 2007). Therefore it has been suggested that the Dynactin complex acts as a cargo receptor for motor complexes, mainly Dynein. In good agreement with this finding is the co-localization of Dynactin with Golgi associated vesicles (Habermann et al., 2001). In contrast, recent work by Haghnia et al. (2007) suggests that the complex is nonessential for membrane-recruitment of motor proteins.
4.2.2 Phenotypes of Dynactin LOF

Dynactin LOF phenotypes in Drosophila are observed in a variety of developmental processes, including oogenesis, spermatid growth, larval motility and eye formation (section 1.1.1; Haghnia et al., 2007; Gosh-Roy et al., 2005; Duncan and Warrior, 2002; Fan and Ready, 1997; Reddy et al., 1997; Fan, 2004). The requirement of Dynactin during oogenesis manifests in a failure of oocyte differentiation in Arp1 mutant ovarioles, similar to the phenotypes observed in bicaudalD (bicD) and egalitarian (egl) mutants (section 1.1.1; Mach and Lehmann, 1997; Haghnia et al., 2007). When Dynactin function is impaired during late oogenesis, the deposition of morphogens during anteroposterior axis formation in the oocyte is affected (Januschke et al., 2002).

Due to the experimental constraints of early embryogenesis in Drosophila, the function of Dynactin during blastoderm cellularization has not been investigated directly yet. However, a recent study shows, that inhibition of the Dynein motor complex during blastoderm cellularization causes effects, closely resembling the phenotypes induced by lva LOF or MT depolymerization (section 1.1.3; section 4.1.4; Papoulas et al., 2005). Embryos from transheterozygous dynein heavy chain (dhc) hypomorphic mutant mothers develop until mitosis 14 but then fail to cellularize (Papoulas et al., 2005). The localization of Lva-positive structures in these embryos is, in contrast to wt restricted to the basal side of the nuclei (Papoulas et al., 2005). Furthermore, the injection of specific antibodies against Dhc results in a strong reduction of membrane invagination speed during both the slow and fast phase of cellularization (Papoulas et al., 2005). On the basis of the extensive characterization of the cooperative function of Dynactin and Dynein (reviewed in Karki and Holzbaur, 1999; King et al., 2003), these phenotypes reflect the phenotypes that would be caused by a
direct inhibition of the Dynactin complex function with a high probability.

4.2.3 Functional correlation between Dynactin and Aen

The comparison of aen phenotypes with the phenotypes described for dynactin LOF clearly shows that many processes that require Dynactin activity are unaffected in aen mutants. This applies to the Dynactin function in axis formation. The presence of Aen proteins during late oogenesis might indicate a requirement of Aen function in Dynactin-dependent morphogen deposition (figure 9; Januschke et al., 2002). However, the axis formation is not affected in aen mutants, as shown by the unipolar positioning of PGCs and the absence of characteristic morphological defects later in development. Phenotypes in postembryonic stages attributed to dynactin LOF, like larval posterior paralysis or rough eye phenotypes in adults (Haghnia et al., 2007; Karki and Holzbaur, 1999), have not been observed in aen mutants. These observations show, that Aen is no constitutive functional adjunct of the Dynactin complex. Conversely, no PGC migration defects attributed to dynactin LOF have been described in literature (reviewed in Karki and Holzbaur, 1999; Papoulas et al., 2005)

During cellularization, however, the almost complete co-localization of Aen and the Dynactin associated motor protein Dhc suggests a functional interaction between these factors (figure 16). The similarity between phenotypes induced by dhc LOF and aen LOF, both characterized by a partial failure of blastoderm cellularization (Papoulas et al., 2005), supports this hypothesis along with the biochemical data (see also section 4.1.3; figure 14). These data, in connection with the co-localization of Aen proteins with components of the secretory pathway (figure 12), strongly suggests that Aen is involved in intracellular transport during cellularization. The ectopic accumulations of Nrt in the apical cytoplasm of affected regions of
aen mutants indicate that Nrt is still transported in apical direction suggesting that the membrane transport is affected downstream of the Golgi (Sisson et al., 2000; Lecuit and Wieschaus, 2000; Papoulas et al., 2005). The co-localization of Aen and Rab11 shows that Aen-positive vesicles reach the RE and are normally processed as indicated by the absence of ectopic Nrt enrichments directly in the RE (section 1.1.3; figure 18; section 4.1.3; Pelissier et al., 2003). Thus the effect of the aen mutation most probably affects a process that functionally is arranged in between the budding of secretory vesicles from the RE and their fusion to the PM.

In order to further narrow down the functional role of Aen in secretion, further candidates for a direct interaction with the Aen proteins needed to be identified.

4.3 The interaction between Slam and Aen

The identification of Slam as a potential interaction partner of Aen was due to its described involvement in cellularization and PGC migration (Lecuit et al., 2002; Stein et al., 2002). Similar to Arp1, the binding of Slam to AenA was tested by means of GST-Pulldown and Co-IP. It was found, that the two proteins interact and exhibit a partial co-localization, which covers the FC and the basolateral membrane.

The Slam protein has been shown to be involved in the formation of the FC and the basal adherence junction (BAJ) (Lecuit et al., 2002). It is required to recruit other components of the FC and the contractile ring to the forming furrow (section 1.1.3, Lecuit et al., 2002). Slam has been suggested to mediate the deposition of yet uncharacterized factors that affect PGC migration at stage 10 after their transmigration through the primordial midgut (PMG) (section 1.1.4; Stein et al., 2002). The phenotypes of amorphic Slam LOF alleles include a malformation of the FC, which is not cleared of villous projections prior to
invagination (Lecuit et al., 2002). Invagination of the FC is delayed and progresses at about half the velocity of wt during the fast phase of cellularization (Lecuit et al., 2002). In good agreement with its localization to the FC and to the basolateral membrane, is the finding that slam LOF affects the formation of the BAJ and the contractile ring (Lecuit et al., 2002). The localization of several factors including Armadillo, the PDZ-protein D-PATJ, the motor protein Myosin-II is dependent on Slam (Lecuit et al., 2002). Also the transport of Nrt has been found to be affected in slam LOF, leading to its ectopic localization in the apical cytoplasm (Lecuit et al., 2002). Therefore, it has been hypothesized that the integration of vesicles to the exocytic domains of the PM is affected in Slam mutants (Lecuit et al., 2002). By comparison with amorphic alleles, the phenotypes of hypomorphic alleles occur in later developmental stages. In these embryos, the migration of PGCs is affected (Stein et al., 2002). The defects observed are characterized by an inhibition of the transit of PGCs from the PMG to the mesoderm (section 1.1.4; Stein et al., 2002).

The phenotypes described for maternal-zygotic, hypomorphic Slam LOF and amorphic slam LOF mutants show cellularization defects that are characterized by a delay of membrane invagination resulting in a failure of blastoderm cellularization (Lecuit et al., 2002). Furthermore, Nrt is localized in the apical cytoplasm instead of the PM. Although no delay in cellularization was detectable in aen mutants, Nrt is mislocalized similar to slam mutant embryos. However, the PGC migration defects observed in zygotic, hypomorphic slam mutant embryos (Stein et al., 2002) do not resemble those observed in aen mutant embryos. No premature PGC transmigration during stages 5 to 8 is observed in hypomorphic slam mutants. Conversely, no abnormal migration of PGCs after stage 9
occurs in aen mutant embryos. Therefore the processes causing PGC mismigration in slam mutants are most probably independent of aen.

To show whether Aen and Slam interact on a functional level during blastoderm cellularization, genetic interaction experiments were done utilizing a hypomorphic mutation (Stein et al., 2002) in presence of a maternal contribution. The results of these experiments show that phenotypes observed in slam-aen double mutants are strongly increased in frequency and intensity as compared to slam and aen single mutants (sections 3.7.1 and 3.7.3; figures 17 and 19). This phenotypic increase is not due to the addition of independent slam related and aen related defects, but clearly exceeds the added effects of the single mutants.

The increase phenotypic strength could be interpreted as a qualitative shift from a hypomorphic to an amorphic phenotype, which can no longer be rescued by the maternal contribution (Lecuit et al., 2002; Stein et al., 2002). This shift might be explained by the decrease of Slam protein concentration at the place, where it is normally localized and functionally required. This would mean that the concentration in hypomorphic mutants is high enough to maintain Slam function. If the concentration of Slam is further decreased by removing the maternal contribution or as a consequence of aen LOF, the concentration sinks below a critical threshold and cellularization defects occur. Such a model has been offered by Stein et al (2002) to explain the phenotypes induced by removal of the maternal contribution in the hypomorphc mutants. Alternatively, the phenotypic shift is not caused in dependence of Slam concentration, but represents a decrease of functional efficiency, which could be due to the missing of Aen as a functional adjunct.
4.3.1 Functional models for Aen Slam interaction

1) Slam transport: Aen acts as a specific cargo receptor, which recruits Slam to the Dynein-Dynactin transport complex. Upon binding Aen, Slam is transported to the sites of the prospective FC.

In addition to the biochemical and genetic interaction studies, this hypothesis is supported by the close co-localization of Aen with the transport complexes and secretory vesicles on one and its association with Slam adjacent of the FC and the basolateral membrane on the other side. The strong enhancement of the zygotic slam phenotypes found in the double mutants could be explained with this model. In the slam single mutant, low levels of maternal supplied Slam are sufficient to maintain cellularization. If the efficiency of transport of Slam to the prospective FC is decreased because of the lack of Aen as a specific cargo adaptor, the Slam concentration at the FC is reduced beyond a critical threshold and cellularization fails.

2) Slam mediated membrane recruitment to the FC: In addition to the recruitment of cargo to the motor complexes the intersection of Aen with Slam could enable specific membrane targeting.

Slam might mediate specific recruitment of Aen-decorated cargo to the basolateral membrane and enable subsequent membrane fusion. This targeting of transport might be involved in the establishment of adhesive structures and membrane polarity. This hypothesis would be in agreement with the subcellular localization of the factors involved and might also explain the phenotypes observed in slam-aen double mutants. Furthermore, this essential function of the Slam-Aen interaction would explain, why Aen is not required in all Dynactin complex functions. However, the interaction of Aen and Slam cannot
be essential for the proper localization, as the aen phenotype is not identical to the slam maternal zygotic phenotype. Therefore, Aen seems be required for membrane fusion but not essential for the targeting of membranes.

3) Constitutive Aen Slam interaction at the FC: Aen might cooperate with Slam in the establishment of a membrane-targeting complex that promotes polarized membrane insertion at the basolateral membrane.

Aen might provide a physical link between the BAJ and the MT cytoskeleton by connecting the Dynactin complex to Slam. This function might support BAJ integrity and also facilitate the polarized transport of secretory and transcytotic vesicles. This hypothesis is in partial agreement with the phenotypes observed in aen-slam double mutants. However a direct involvement of Dynactin in adherence junction assembly and maintenance during embryogenesis has not been shown yet. Furthermore, this hypothesis would not explain the large pools of cytoplasmic and vesicle associated Aen proteins (figure 10; figure 11; figure 12).

These functional models however do not include an explanation for the PGC mismigration observed either in aen or in slam mutants. The most obvious explanation for the premature transmigration of PGCs through the PMG would be a weakening of the epithelial integrity.

The analysis of aen and slam mutants reveals some phenotypes that are in good agreement with a defective epithelial integrity. The phenotypes observed in amorphic slam mutants include the malformation of adhesive structures and affect membrane polarity (Lecuit et al., 2002). Moreover, the adhesion factor Nrt is mislocalized (Hortsch et al., 1990). Similarly, in a subset of aen mutant embryos mislocalization of Nrt can be observed as well (figure 7). In addition,
the analysis of the cuticle structures of aen-slam double mutant larvae shows a dramatic loss of cuticle structure (figure 18). This phenotype is highly reminiscent of phenotypes observed in mutants of bazooka (baz), a factor required for the establishment of epithelial polarity (section 1.1.4; Müller and Wieschaus, 1996; Knust et al., 1996). This indicates that the epithelial integrity is strongly affected in aen-slam double mutant embryos suggesting that both factors are cooperatively involved in the establishment of the epithelial barrier function. Therefore, it would be expected that the integrity of the epithelium is also affected in the hypomorphic slam mutant as well as in the aen mutant. The mislocalization of Nrt in both mutants and the untimely PGC transmigration in aen mutants support this hypothesis. So, based on the idea that untimely PGC transmigration is caused by a reduction in epithelial integrity, PGC migration could be used as an independent assay to screen for defective epithelial integrity. However, the lack of premature transmigration in slam single mutants shows that untimely PGC transmigration might only occur if the integrity is dramatically weakened – as in the double mutant - or alternatively, that the PGC transmigration in aen mutants is not monocausally induced by the reduction of epithelial integrity but rather reveals an additional function of aen. Therefore Aen might be involved in the transport of signaling factors required for the modulation of PGC migration.

1.5.2. Signaling pathways potentially involved in premature PGC migration in aen mutants

In the wt situation PGC migration occurs after embryonic stage 9 when the close adherence of the PGCs is weakened (Jaglarz and Howard, 1994). In addition, epithelial transmigration is controlled by the cells of the PMG, as a transformation of these cells into hindgut cells in huckebein mutants blocks of transmigration completely
(Jaglarz and Howard, 1994; Reuter 1994; Moore et al., 1998). The importance of the PMG cells is further shown by transplantation and in vitro experiments (Jaglarz and Howard, 1994; Jaglarz and Howard, 1995). When PGCs are explanted from stage 5 embryos and kept in primary cell culture, they start to migrate. When instead they are transplanted into embryos, they do not migrate through the PMG epithelium until stage 9 begins (Jaglarz and Howard, 1995). By isochronic and heterochronic transplantation experiments, where the age of donor and recipient embryo were either matched or leading or lagging by up to 3 h, it was shown that the time of PGC transmigration is not determined by the age of the PGCs, but rather by the stage of the midgut (Jaglarz and Howard, 1994). Based on these results transmigration of PGCs is controlled by signaling from PMG cells.

1.5.2. JAK/STAT signaling in PGC migration

The only signaling cascade, which has been shown to influence PGC migration during stage 5, is the Janus kinase/signal transducer and activator of transcription (JAK/STAT) signaling pathway (reviewed in Hou et al., 2002, Brown et al., 2006). In the PGCs, the JAK/STAT system functions downstream of the receptor tyrosine kinase (RTK) Torso (Tor) (Li et al., 2003; reviewed in Duffy and Perrimon, 1994) and has been shown to influence both early PGC division and migration.

Components of the JAK/STAT signaling pathway, including hopscotch, encoding the Drosophila Jak, and Stat92E (Hou et al., 1996; Li et al., 2003) exhibit a similar localization than Aen (Li et al., 2003). Transcripts are maternally deposited and ubiquitous during early embryogenesis followed by a stabilization of the mRNA in the PGCs after stage 3 (Li et al., 2003). The phenotypes observed in LOF and gain of function (GOF) variants of components of the JAK/STAT pathway (Li et al., 2003; Brown et al., 2006) suggest that the signaling pathway is involved in several aspects of PGC development. In Stat92E mutants, the formation of
filopodia during active migration is compromised (Jaglarz and Howard, 1995; Brown et al., 2006) and consecutively, PGC migration at stage 10-11 is delayed or inhibited (Li et al., 2003; Brown et al., 2006).

Striking phenotypes are also observed in the GOF situation. The analysis of a hopscotch (hop) GOF allele, which leads to overactivation of the pathway (Binari and Perrimon, 1994) showed an overproliferation of the PGCs. Furthermore, the PGCs show an increased migratory behavior resulting in their translocation from the posterior pole. However, in contrast to the aen phenotype, the PGCs do not only transmigrate but translocate on the embryonic surface, leading to dispersed PGC clusters that extend into anterior direction. However, later at stage 8-9 the PGCs show premature migration through the PMG epithelium (Li et al., 2003) resulting in PGCs in ectopic locations during later stages. Furthermore, the ectopically localized PGCs appear to be stabilized in hop GOF (Li et al., 2003), while in wt they are eliminated (Coffman et al., 2002).

Despite the occurrence of premature PGC transmigration phenotypes in both aen mutant and hop GOF, a direct functional interaction of aen with the JAK/STAT pathway appears unlikely. The phenotypes caused by STAT overactivation demonstrate a general increase in the migratory behavior of PGCs resulting in a migration along the embryonic surface, while only a subset of PGCs transmigrates the blastoderm epithelium (Li et al., 2003). In aen mutants, migration along the embryonic surface has not been observed. This observation is counterindicative of a general increase in migratory potential of PGCs in aen mutants. Moreover, a genomewide RNAi screen for so far uncharacterized components of the JAK/STAT pathway (Müller et al., 2005) has given no indication of a direct implication of Aen in JAK/STAT signaling. However a potential mutual influence of Aen and JAK/STAT signaling remains an interesting option.
4.3.2 Tre1/Wun signaling in PGC migration

The G-protein coupled receptor Trapped in endoderm-1 (Tre1) in a signaling factor involved in the initiation of PGC migration through the PMG (Kunwar et al., 2003). In tre1 mutant embryos the PGCs remain trapped in the PMG pocket at stage 10 (Kunwar et al., 2003). The inability to migrate through the PMG does not reflect a general effect on PGC migration as escaping PGCs retain the ability to migrate towards the somatic gonadal precursor cells (SGPs) and become incorporated into the gonads (Kunwar et al., 2003). Tre1 functions cell autonomously in the PGCs as shown by transplantation and GAL4-UAS-mediated rescue experiments. tre1 mRNA is like aen maternally contributed and stabilized in the PGCs. However, no antibody is available to show if the Tre1 protein is PGC specific (Kunwar et al., 2003).

These data indicate that an attractive signal from the PMG is required to allow PGC transmigration at stage 9. Based on the tre1 mutant phenotype a potential explanation for the PGC migration observed in aen mutants could be a premature signal from the PMG cells to the PGCs at late stage 5 instead of stage 9 in wildtype. The ectopic secretion of an attractive signal would be contradictory to the reduced secretion efficiency indicated by Nrt mislocalization in aen single- and slam-aen double mutants.

Another possibility would be that Aen is acting as a negative regulator of tre1 signaling in the PGCs. However, based on the partial rescue of the transmigration phenotype in aen mutants by specific expression in the somatic cells a PGC autonomous effect can be ruled out.

Wunen (Wun) and Wunen2 (Wun2) are phospholipid phosphatases (section 1.1.4; Zhang et al., 1996; Starz-Gaiano et al., 2001; Renault et al., 2004) and have been hypothesized to act in concert with Tre1 in
influencing PGC migration during stages 9 to 11. Somatic tissues expressing Wun and Wun2 have been suggested to repel PGCs by depleting their environment from an otherwise ubiquitous attractive signal (Renault et al., 2004). However, neither do wun and wun2 mutants exhibit premature PGC mismigration (Zhang et al., 1996; Starz-Gaiano et al., 2001) nor does PGC mismigration occur in aen mutants at stages 9 to 11. This clearly suggests that aen phenotypes are independent of Wun signaling.

4.4 Outlook

In Drosophila, several factors have been identified that participate in the regulatory and physical interaction between migratory cells and tissue. However, the understanding of the functional network required for initiation and permission of tissue invasion is still incomplete (reviewed in Kunwar et al., 2006). In this work, it was shown that Aen participates in the transport of de novo synthesized factors to the PM during the cellularization of the somatic blastoderm. aen mutants exhibit premature PGC transmigration phenotypes. These phenotypes might reflect the mislocalization of factors, which otherwise prevent PGC transmigration until embryonic stage 9.

Therefore Aen might assist in the identification of novel signaling factors involved in the regulation of PGC transmigration. For this purpose, further interaction partners of Aen could be identified by pulldown from embryonic extracts and subsequent mass-spectrometrical analysis. Additionally the homozygous aen\(^2\) mutation could be used as a sensitized genetic background for EMS mutagenesis to identify mutations inducing infertility as a consequence of complete premature PGC transmigration.

These experiments might contribute to the identification of molecular programs that control cell migration and tissue invasion.
5 Literature


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