NHA1 is a cation/proton antiporter essential for the water-conserving functions of the rectal complex in Tribolium castaneum.

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More than half of all extant metazoan species on earth are insects. The evolutionary success of insects is linked with their ability to osmoregulate, suggesting that they have evolved unique physiological mechanisms to maintain water balance. In beetles (Coleoptera)—the largest group of insects—a specialized rectal (“cryptonephridial”) complex has evolved that recovers water from the rectum destined for excretion and recycles it back to the body. However, the molecular mechanisms underpinning the remarkable water-conserving functions of this system are unknown. Here, we introduce a transcriptomic resource, BeetleAtlas.org, for the exceptionally desiccation-tolerant red flour beetle *Tribolium castaneum*, and demonstrate its utility by identifying a cation/H\(^+\) antiporter (NHA1) that is enriched and functionally significant in the *Tribolium* rectal complex. NHA1 localizes exclusively to a specialized cell type, the leptophragmata, in the distal region of the Malpighian tubules associated with the rectal complex. Computational modeling and electrophysiological characterization in *Xenopus oocytes* show that NHA1 acts as an electroneutral K\(^+\)/H\(^+\) antiporter. Furthermore, genetic silencing of *Nha1* dramatically increases excretory water loss and reduces organismal survival during desiccation stress, implying that NHA1 activity is essential for maintaining systemic water balance. Finally, we show that Tiptop, a conserved transcription factor, regulates NHA1 expression in leptophragmata and controls leptophragmata maturation, illuminating the developmental mechanism that establishes the functions of this cell. Together, our work provides insights into the molecular architecture underpinning the function of one of the most powerful water-conserving mechanisms in nature, the beetle rectal complex.

**Significance**

Beetles are the largest group of animals on earth. Unique adaptations in overcoming water stress is critical to their success in arid areas, yet the mechanisms underpinning this ability are unknown. Using genetics and electrophysiological studies, we show that a cation/H\(^+\) (NHA1) transporter is exclusively localized to specialized leptophragmata cells in the Malpighian tubules associated with the rectal complex. Ion transport by NHA1 in leptophragmata underpins the movement of water from the rectum to recycle it back to the body, and is essential for maintaining systemic water balance in beetles. This work provides insights into the molecular architecture of one of the most powerful water-conservation mechanisms in biology, and provides an important clue to the evolutionary success of the beetles.

Insects are (in terms of species) the most diverse animal group on the planet, occupying the widest possible range of habitats on earth (1). However, insects—being a predominantly terrestrial group—are faced with major problems in maintaining ion and water balance as their small size and large surface-to-volume ratio make them highly sensitive to osmotic disturbances. As such, the evolutionary success of insects is intrinsically linked with their ability to defend against harmful changes in their water contents in a wide range of environments. In insects, the renal (Malpighian) tubules (MTs) and the hindgut are the principal organs responsible for regulating body fluid composition (2). Whereas the MTs secrete excess ions and water by producing a primary urine that is drained into the alimentary canal (3), the hindgut selectively reabsorbs solutes and water in proportion to the needs of the animal (4). The hindgut (in particular the rectum) is thus the major site of water conservation in insects as it provides vital feedback control of the final composition and volume of the excretory products. However, surprisingly little is known about the mechanisms that mediate the selective absorption of water and solutes by the rectum in different external conditions.

The physiological importance of the rectum in maintaining overall water balance in insects is particularly evident in species colonizing arid environments. For example, in the mealworm *Tenebrio molitor*—a species that is capable of completing its entire life cycle without access to environmental water—a specialized rectal (“cryptonephridial”) complex has evolved that enables recovery of almost all water from the rectum (5, 6). The anatomical arrangement of this complex is defined by the distal ends of the MTs (perirectal tubules, PTs) being closely applied to the rectal wall with the entire structure enclosed beneath an impermeable perinephric membrane (6, 7). This cryptonephridial condition is found in most members of Coleoptera (beetles) and in most larvae of Lepidoptera (moths and butterflies); however, the rectal complexes found in these two insect Orders differ markedly in their anatomy and function (8–10), implying that the two structures likely evolved
through convergent evolution. In *Tenebrio*, the water-conserving properties of the rectal complex are believed to rely on the active potassium chloride (KCl) transport by the PTs to generate a fluid of sufficiently high osmotic pressure to facilitate osmotically driven water removal from the feces (11, 12). In effect, the system establishes a standing gradient along the anterior–posterior axis of the complex to maximize fluid absorption in a manner analogous to the countercurrent arrangement of the vertebrate nephron in the renal medulla (13). In this way, *Tenebrio* (and its relatives) is able to extract and recycle almost all water from the rectal lumen to produce powder-dry excreta (7, 14). Remarkably, the system can even be used as a physiological mechanism for water uptake by enabling absorption of water vapor directly from the moist air (5–7, 14, 15). It has been suggested that the accumulation of KCl in the PTs is mediated by a small population of secondary cells known as “leptophragmata,” as they are the only cells that interrupt the peritrophic membrane to enable hemolymph-to-tubule movement of KCl (6, 7). However, in spite of having been intensely studied for almost a century, the cellular and molecular architecture underpinning the water-extracting functions of this extraordinary organ remains largely unknown.

Cataloging the relative strength and specificity of gene expression across different tissues and life stages of an organism can provide valuable insights into most biological functions. Transcriptomic atlases have therefore become powerful tools in the functional genomics arsenal by enabling the annotation of physiological mechanisms and developmental processes on a gene-by-gene basis (16–20). However, despite their obvious benefits to their respective communities, the number of such transcriptomic atlases is limited. In the field of insect functional genomics, the red flour beetle *Tribolium castaneum* (a closely related species of *Tenebrio*) has emerged as a powerful model system because of its rapidly expanding transgenic toolkit (21–24) and its amenability to large-scale mutagenic studies (25, 26). The construction of an authoritative expression atlas for *Tribolium* would thus not only complement with existing postgenomic resources, but also help broaden the scope for functional analysis of one of the most ecologically (contain many devastating crop pests) and ecologically (largest group of insects) important animal groups on Earth, the beetles.

Here, we report the development of BeetleAtlas, a transcriptomic atlas covering distinct ontogenetic and tissue-specific expression profiles of *Tribolium*, and demonstrate the utility of this resource by identifying a cation/H+ antiporter (NHA1) that is essential to the water-extracting properties of the *Tribolium* rectal complex. Using a combination of bioinformatics, genetics, imaging, electrophysiology, and organ assays, we show that NHA1 localizes exclusively to the specialized leptophragmata cells in the PTs where it acts as an electroneutral cation/H+ antiporter. Genetic depletion of *Nha1* dramatically increases excretory water loss and impairs whole-animal survival during desiccation stress, suggesting that NHA1 activity is essential for maintaining systemic water balance. Finally, we show that NHA1 expression and leptophragmata maturation is regulated by a transcription factor called Tiptop, which is part of a conserved gene regulatory network that is central to the function of the rectal complex in tenebrionid beetles. Taken together, our work provides insights into the molecular architecture underlying the function of one of the most powerful water-extracting mechanisms in biology, the tenebrionid rectal complex.

**Results**

**BeetleAtlas: A Transcriptional Atlas of *Tribolium* Tissues and Life Stages.** A large fraction of any animal genome is differentially expressed in distinct cell types, tissues, and life stages. Therefore, it is vital to consider spatiotemporal changes in gene expression in order to understand cell functions. To gain a comprehensive view of the transcriptional landscapes of individual tissues and life stages of *Tribolium*, we have created BeetleAtlas (BeetleAtlas.org). This is a transcriptional atlas of gene expression based on RNA sequencing (RNAseq) that covers several embryonic stages as well as major larval and adult tissues. Specifically, it catalogs transcriptomic datasets covering 11 distinct adult tissues (head, brain, anterior midgut, posterior midgut, hindgut, Malpighian tubules, rectal complex, fat body, female/male gonads, and carcass) dissected from 7-d-old San Bernardino adult beetles; nine larval tissues (head, brain, anterior midgut, posterior midgut, hindgut, Malpighian tubules, rectal complex, fat body, and carcass) dissected from L6 larvae; and four distinct (0 to 1, 1 to 24, 24 to 36, and 36 to 72 h post egg lay) stages of embryonic development (Fig. 1A). Each sample was prepared according to standardized protocol and sequenced on the same sequencing platform in a minimum of three biological replicates and compared to matched whole-animal samples. Together, BeetleAtlas thus allows systematic comparisons of gene expression across all major tissues and ontogenetic stages.

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**Fig. 1.** BeetleAtlas is a comprehensive transcriptomic atlas of gene expression. **(A)** Adult *Tribolium* anatomy highlighting the tissues selected for microdissection and bulk RNA sequencing. **(B)** Clustered heat maps of gene expression (log2-transformed values of fragments per kilobase of transcript per million, FPKM) for select genes that are enriched in a certain tissue or life stage in *Tribolium*. The same genes are depicted across all samples demonstrating distinct transcriptional signatures for each tissue or ontogenetic stage.
of more than 16,500 genes (with over 18,500 transcripts) encoded by the Tribolium genome (27).

An implicit requirement of such a facility is the robustness of the data in the underlying database. To formally assess the quality of our data, we identified the most highly expressed genes in each sample and generated a clustered heat map to visualize the divergence of transcriptomes between the different ontogenetic stages and larval and adult tissues (Fig. 1B). These data show a clear discrimination of the transcriptional signatures, not only between different life stages, but also between physically adjacent tissues within the same stage, such as the larval midgut and tubules or the adult rectal complex and gonads. Furthermore, we performed manual searches for genes that show a strong enrichment in a particular tissue and performed independent validation of gene expression by qRT-PCR. These results further confirm and validate the spatial expression profiles reported by BeetleAtlas across the selected genes (SI Appendix, Table S1). BeetleAtlas is thus a powerful tool for the Tribolium and insect communities, providing a critical first step by mapping gene activity to particular stages or tissues and by offering valuable information on the function of each gene product by integrating complementary large-scale RNA interference (RNAi) screening data (28).

**BeetleAtlas Identifies Candidate Genes Involved in Rectal Complex Function.** To demonstrate the utility of BeetleAtlas in providing transcriptional insights into tissue function, we aimed to characterize the molecular mechanisms underpinning the function of the rectal complex (6). Using the “Gene” lookup section of BeetleAtlas, we performed manual searches of all genes identified as core components of the insect “epithelium” (29) and looked for transcript enrichment of the Tribolium orthologs relative to that of the whole animal. In addition, we used the “Tissue” search function to perform an unbiased search for candidate transporter genes that are most highly enriched in the rectal complex. Adopting parallel hypothesis-driven (candidate genes) and hypothesis-free (gene enrichment) approaches, we identified more than 30 putative genes that are expressed at very high levels in this multiorgan system compared to the organismal average, and further revealed that most of these genes belong to a core set that is coexpressed in other insect transport epithelia (29, 30).

From this candidate list, TC013096, a gene predicted to encode a cation/H⁺-antiporter (Tcas-Nha1, hereafter just Nha1), shows the highest enrichment in the rectal complex (Fig. 2A). Indeed, mapping the tissue-specific expression pattern of Nha1 indicates that this gene is predominantly expressed in the rectal complex of both larval and adult Tribolium (Fig. 2B–D); an expression profile that we validated by qRT-PCR (SI Appendix, Table S1). Together, the transcriptional signature of Nha1 suggests that this transporter is likely to play a key role in the function of the Tribolium rectal complex.

**Nha1 Localizes to a Specialized Cell Type in the Perirectal Tubules of the Rectal Complex.** To gain insight into the physiological roles of NHA1, we next examined the structure and spatial organization of the Tribolium rectal complex using scanning electron microscopy. Like the rectal complex from the closely related mealworm Tenebrio (6, 7, 31), the Tribolium rectal complex is organized according to the cryptonephridial condition. This condition is characterized by the distal ends of the MTs being closely applied to the rectal epithelium in a sinuous pattern and enclosed within a compartment, the perinephric space that is separated from the body cavity by an impermeable perinephric membrane (Fig. 3A–C). Notably, the distal regions of the MTs—the so-called perinephric tubules (PTs)—are conspicuously different from the “free” part of the tubules as they are characterized by small dilations over which the perinephric membrane is extremely thin and in direct contact with the dilations (“boursourflors”) under which specialized small-nucleated cells known as leptophragmata are found (Fig. 3B and B'). The anatomical position of these cells is intriguing as previous studies in Tenebrio had demonstrated that the function of the rectal complex is based on the accumulation of high concentrations of KCl in the PT lumen mediated by active

![Fig. 2](https://www.pnas.org/content/120/13/e2217084120.full)
hemolymph-to-tubule transport to facilitate osmotically driven water removal from the feces (6, 7, 12). Such a model is supported by the high expression of genes encoding the different subunits of the plasma membrane V-ATPase (V-ATPase) (Fig. 2 A and SI Appendix, Fig. S1 C)—a plasma membrane transporter critical to energizing insect epithelia (32)—and by the exclusive localization of the protein to the apical brush border of the PTs in the *Tribolium* rectal complex (Fig. 3 G). As the perinephric membrane is highly impermeable except for the “blister-like” windows under which leptophragmata sit (33), the anatomical data strongly suggest that these small cells are the only sites of exchange. Consistent with this notion, backscattered electron detection—a method that carries information on differences in atomic number ($Z$) of the sample—reveals that silver staining is exclusively found at sites corresponding to the anatomical position of the leptophragmata (Fig. 3 A–F). Maximum projection confocal microscopy images of paraffin sections of the rectal complex demonstrate that NHA1 localizes predominantly to the apical brush border of principal cells in the PTs (scale bar, 40 µm). Note the cuticle of the rectal epithelium shows strong autofluorescence (arrows). (H) Maximum projection confocal microscopy images demonstrate that NHA1 is expressed in this subpopulation of LP cells along the PTs (small arrows). Zoom: note the small nucleus of the LP and the characteristic central band of F-actin spanning the apical surface of the cell (small arrows). Cross-section of PT: Subcellular localization of NHA1 demonstrates exclusive expression in the small-nucleated LP cells. Rectal complex dissected from animals injected with dsRNA targeting the *Nha1* gene (*Nha1-RNAi*) shows a dramatic reduction in immunoreactivity confirming the specificity of the antibody. (I) Identity is further confirmed by coexpression of the Tiptop (Tio) transcription factor, which is a marker of the LPs [as well as secondary cells in the "free" part of the tubule (34)].
NHA1-specific antibody. The specificity of the antibody was verified by lack of immunoreactivity in Nha1-depleted animals (SI Appendix, Fig. S1 A and B). Indeed, the expression of NHA1 in leptocephalagmata was confirmed by double staining with Tiptop (Fig. 3D), a transcription factor that marks leptocephalagmata and is involved in the differentiation of secondary cells in Tribolium and other insects (34–36). This finding suggests that the leptocephalagmata and the secondary cells of the free tubules are related. Taken together, our data indicate that the anatomical features of the Tribolium rectal complex are largely identical to those described previously for Tenebrio (6, 7), and that NHA1 is exclusively localized to the specialized leptocephalagmata cells in the P Ts of the rectal complex.

**NHA1 Acts as an Electroneutral Cation/H+ Antipporter.** The NHA transporters belong to the Cation/Proton Antiporter-2 (CPA2) subfamily of proteins, which are known to participate in a broad range of transport mechanisms across all kingdoms of life (37). Classically, the NHAs are secondary transporters that exchange sodium ions against protons, yet several CPA2 members have been shown to move different types of substrates (38, 39), suggesting that their molecular function cannot be inferred from structural similarity. To gain insight into the role of NHA1 in rectal complex physiology, we first performed sequence analysis and computational modeling of the protein. Surprisingly, we found only one Nha gene (Nha1) encoded by the Tribolium genome, which contrasts with other insect genomes in which two paralogs (Nha1 and Nha2) are present (37, 38, 40). Interestingly, a BLASTp search against Drophiila proteins reveals that Tribolium NHA1 is more closely related to Drophiila NHA1 (53% sequence identity) than to NHA2 (35% sequence identity).

Next, we performed three-dimensional structure predictions using I-TASSER (41) followed by Ramachandran plot and molecular dynamics simulations (42) to identify the putative transmembrane domains and globular structure of the protein. Consistent with the structural hallmarks of CPA2 members, the structural architecture of Tribolium NHA1 is defined by 12 transmembrane helices (both the N- and C-terminal tails are located in the cytoplasm) that collectively form a negatively charged transport funnel containing the putative ion binding and translocation domains (Fig. 4 A and B and SI Appendix, Fig. S2 A–C and Movie S1). These findings led us to perform molecular dynamics simulations in the presence of cation substrates (K+ or Na+) to help resolve the ion binding and cation selectivity of NHA1. After an initial phase of equilibrium, we observed that the structural stability of the protein was predicted to be higher (i.e., lower radius of gyration and root mean square deviation) in the presence of bound K+ relative to bound Na+ (Fig. 4 C and D), implying that K+ binding is favored over Na+ by the Tribolium NHA1 protein.

We next sought to experimentally validate our molecular dynamics simulations by cloning and heterologously expressing Tribolium NHA1 in Xenopus laevis oocytes to allow electrophysiological characterization of the protein, as described in ref. 43. Monitoring intracellular pH (pHi) in NHA1 oocytes while sequentially manipulating extracellular [Na+]i, [K+]i, and pH in the superfusing solution revealed that the oocytes rapidly responded to these changes, whereas similar responses were not observed in control oocytes injected with water (Fig. 4 E and F). These data suggest that Tribolium NHA1 is capable of recognizing both Na+ and K+ as substrates with either cation being exchanged for a H+. However, the significantly lower pHi of resting NHA1 oocytes relative to water-injected controls (Fig. 4 G) supports our in silico analysis predicting that K+ is favored over Na+ (Fig. 4 C and D). The membrane potential (V_m) of NHA1 oocytes was insensitive to extracellular pH, implying that NHA1 transport is not electrogentic (Fig. 4E and SI Appendix, Fig. S3). Taken together, our results show that in Tribolium, NHA1 is functionally coupled to...
the H⁺ gradient to mediate electroneutral exchange of K⁺ (Na⁺) for H⁺ (Fig. 4A) in a manner similar to that observed for fungal KHA members (39, 44, 45).

Internal Water Abundance Modulates NHA1 Activity in Leptophragmata Cells. The potent activation of the rectal complex observed in Tenebrio larvae deprived of water (6) implies that the transport machinery of the complex is regulated in response to changes in hemolymph osmotic pressure. We therefore asked whether NHA1 activity is altered in animals exposed to conditions known to affect internal water abundance (34, 46). Quantifying NHA1 expression in the rectal complex revealed that both transcript and protein levels were consistently decreased in animals exposed to conditions that promote fluid retention (water, relative humidity, RH 90%), and significantly increased in beetles exposed to severe desiccation (RH 5%), compared to control animals (Fig. 5 A and B). Furthermore, artificial activation (DH37 injection) and genetic deactivation (Um8 knockdown) of a potent diuretic pathway known to regulate organismal water levels (34) induced a robust increase and a significant decrease in Nha1 expression, respectively (Fig. 5 C and D). These observations suggest that NHA1 abundance is regulated in response to internal changes in hemolymph concentration as part of homeostatic mechanisms that modulate the reabsorptive capacity of the rectal complex to maintain ion and water balance. Such a role is consistent with the observation that the concentration of the perirectal fluid in animals exposed to desiccation increases disproportionately relative to the hemolymph, presumably to allow more effective reabsorption of water from the rectal lumen (6).

Silencing Nha1 Expression Increases Sensitivity to Desiccation by Inducing Excessive Water Loss. We next sought to explore the functional significance of NHA1 in maintaining systemic water balance in vivo by selectively down-regulating Nha1 expression using RNAi. Given that the rate of water loss is a crucial factor in determining the tolerance to desiccation in Tribolium and other insects (34, 47, 48), we hypothesize that Nha1 depletion might lead to increased excretory fluid loss and thus impair the ability of Tribolium to survive dry conditions. Consistent with this hypothesis, Nha1-silenced animals showed an increased sensitivity to desiccation relative to control-injected animals, with a median survival of 3.3 d as compared to 5.8 d (Fig. 6A); RNAi efficacy was verified by qRT-PCR and immunocytochemistry showing >95% knockdown and a complete loss of detectable NHA1 expression (SI Appendix, Fig. S1 A and B). This reduction in median survival is likely explained by an impaired ability to conserve water, since Nha1 knockdown beetles consistently showed an increased rate of organismal water loss and a concomitant increase in hemolymph osmotic pressure (Fig. 6 B and C). These effects were retained, albeit to a diminished extent, in animals exposed to high humidity implying that NHA1 function is essential to maintain systemic water balance across a wide range of conditions of fluid stress (SI Appendix, Fig. S4 A–C). To test if the observed sensitivity to desiccation in Nha1-silenced beetles can be explained by increased excretory water loss, we examined the fecal output profiles of control and knockdown animals using an established in vivo excretion assay based on a dye-laced food source (34, 49, 50). These results showed that Nha1 depletion resulted in increased defection rate and excretory fluid loss as revealed by the production of more abundant, circular, larger, and less concentrated excreta as compared to control-injected animals (Fig. 6 D–H). Indeed, the deposits produced by Nha1-knockdown animals were often visibly associated with a striking increase in excess fluid as manifest by a “halo” surrounding the excreta, which was never observed in control-injected beetles (Fig. 6I). To test whether this increased excretory water loss was causally linked to defects in the reabsorptive capacity of the rectal complex, we adapted and optimized an ex vivo method based on isolating the entire alimentary canal under paraffin oil to quantify fluid reabsorption by the system (51); the preparations remained viable for several hours (SI Appendix, Fig. S5). These experiments revealed that Nha1 knockdown almost completely abolished fluid reabsorption by the rectal complex (Fig. 6J), thus demonstrating that this is, at least partly, the site responsible for the observed reduction in fluid retention. Taken together, our data suggest that loss of NHA1 function in leptophragmata cells dramatically impairs water reabsorption by the rectal complex, which affects systemic water balance and reduces the ability of adult beetles to survive desiccating conditions.

Tiptop-Induced NHA1 Expression Underlies Water Vapor Absorption by the Rectal Complex. The powerful water-extracting properties of the rectal complex are not only related to the reabsorption of water from the feces, but can also be coupled to the remarkable ability to absorb water vapor directly from moist air (5–7, 14). Given that this mechanism depends critically on generating sufficiently low water activities in the rectal lumen to allow condensation of water from the atmosphere (6), we hypothesized that Nha1-knockdown would impair the ability
to perform water vapor absorption. As predicted, we found that Nha1-deficient animals consistently lost body water when deprived of food at high humidity, whereas control animals retained, or even gradually increased, body water during identical exposures (Fig. 7 A and B). These data confirm that Tribolium is able to absorb water at high relative humidities in a manner similar to that observed for Tenebrio (7, 14, 31)—albeit with a reduced capacity—and that this process depends critically on NHA1.

We next explored the gene regulatory networks that govern Nha1 expression and leptophragmata physiology. The finding that Tio—a transcription factor involved in secondary cell differentiation in many insects (34–36)—is coexpressed with NHA1 in the leptophragmata of the PTs (Fig. 3 I), suggests that Tio controls Nha1 expression in leptophragmata. Accordingly, we selectively down-regulated Tcas-Tio (TC012322; hereafter just Tio) expression during early development and subsequently probed for NHA1 expression in the rectal complex of adult beetles. The results revealed that Tio knockdown caused a complete loss of NHA1 expression, and further induced overt defects in the cytoarchitecture of the leptophragmata as evidenced by the failed...
formation of blister-like windows in the perinephric membrane (Fig. 7C). To test whether these defects affected whole-animal physiology, we silenced Tio expression in larvae of Tenebrio (the larger size makes them more amenable to gravimetric studies) and assessed their ability to absorb water vapor (6). As observed in Tribolium, knocking down Tio in T. molitor caused an almost complete loss of NHA1 expression as well as clear morphological changes, such as a loss of the central band of F-actin, to the leptophragmata of the rectal complex (Fig. 3D). Furthermore, silencing Tio expression consistently impaired their ability to absorb water as the body water contents of knockdown animals were significantly lower than those of mock-injected controls (Fig. 7D). Taken together, our study identifies Tio as a key regulator of NHA1 expression and leptophragmata differentiation, which is essential to the function of the rectal complex and to the maintenance of systemic water balance in Tribolium (Fig. 7F).

**Discussion**

**The Transport Functions of the Specialized Leptophragmata Underpin the Water-Conserving Functions of the Rectal Complex.** Classically, absorption of water by Tenebrioid beetles when exposed to subsaturated air was believed to occur across the cuticle, but it has since emerged that this process is entirely performed by the modified rectal complex (6, 7, 15, 31). In this work, we demonstrate that NHA1 in the specialized leptophragmata cells of the PTx is essential for the water-conserving functions of the rectal complex as well as the homeostatic control of water balance in Tribolium (Fig. 7F). How might we explain these physiological effects? Our combined analysis of the NHA1 protein suggests that, while it is capable of handling both Na⁺ and K⁺ movement, the main transport function of NHA1 is likely to mediate K⁺/H⁺ exchange. If this transport modality is replicated in intact animals, these data strongly imply that NHA1 is functionally coupled to the active uptake of K⁺ by the complex, and thus to generating the osmotic forces necessary to facilitate water removal from the feces. Consistent with this idea, Nha1 depletion results in impaired fluid reabsorption by the rectal complex, increased excretion, and reduced organismal water levels. Moreover, both NHA1 transcript and protein are up-regulated in response to cues related to internal water stress. However, a difficulty presents itself when trying to reconcile these observed physiological effects with the fine structure of the rectal complex. Our work and previous studies (6, 7, 12) suggest that the main route of active KCl transport into the system is through the specialized leptophragmata, but these cells do not possess the anatomical hallmarks of active ion-transporting cells. The leptophragmata lack basal infoldings, have a reduced brush border, and contain few mitochondria (31, 52). It would therefore seem unlikely that these cells are solely responsible for mediating the high KCl concentrations measured in the PT lumen (6). By contrast, the larger principal cells possess...
extensive basal infoldings, contain an extensive brush border full of mitochondria (7, 31), and are characterized by high expression of the plasma membrane V-ATPase (Fig. 3G). We therefore speculate that the H⁺ electrochemical gradient generated by the V-ATPase in the principal cells could drive the secondary active transport of K⁺ via NHA1 in the leptocephara, with Cl⁻ following passively (7) to produce the necessary osmotic gradients. Such a model is supported by the recent observation that luminal-directed K⁺ secretion is mediated by the secondary cells of the “free” parts of the MTs in Tribolium (34), and is further analogous to that proposed for Na⁺ uptake by NHA2 in the secondary cells of MTs from the mosquito Anopheles gambiæ (53). However, Nha1 is not expressed in the “free” parts of the tubule in Tribolium (BeetleAtlas.org), and Urn8R—a hormone receptor diagnostic of the secondary cell identity—was not detectably expressed in the leptocephara cells (34), implying that the PTs of the rectal complex operate via a two-cell type model that is distinct from that of the free tubules (34, 35). Mapping the exact routes and mechanisms with which water and ions flows through this multiorgan system remains an exciting prospect for the future. This could be addressed by single-cell RNAseq analysis of the rectal complex to obtain cell-specific insights into the molecular machinery that underlies the actions of the PT and rectal epithelia to help understand one of the most powerful water-conserving systems in biology.

Exploring the Transcriptional Landscapes of Tribolium. Transcriptomic atlastes are powerful tools in molecular genetics as they offer a detailed spatio-temporal view of gene expression that provides valuable clues to each gene’s physiological function (16, 20, 54, 55). Here, we introduce BeetleAtlas; a transcriptomic resource that provides a comprehensive view of the genetic signatures that underpin the functions of distinct tissues or life stages in the genetic model organism T. castaneum (Fig. 1 A and B and SI Appendix, Table S1). Besides being a useful addition to the rapidly expanding toolbox for Tribolium research, BeetleAtlas will broaden the focus of the beetle community. Tribolium is often adopted for studies in evolutionary development; but by cataloguing gene expression across different life stages, BeetleAtlas can help inform a much wider range of biological questions. This is significant because, although systemic RNAi is robust at all stages of development (24, 56), this technique precludes tissue-specific gene interference, and so the tissue(s) that contribute to a given phenotype remain unknown. Our online resource can thus provide a useful filter with which to identify the major tissue(s) in which candidate genes are most abundantly expressed and therefore most conveniently studied. Such an approach might be particularly useful for the post hoc analysis of data derived from large-scale functional screens (25, 28, 57). To further encourage this utility, BeetleAtlas is fully linked to existing databases (iBeetleBase (28) and FlyAtlas (16)) by a common vocabulary, enabling access to this utility, BeetleAtlas thus allows nontechnical users to make a wide range of prepared queries to the underlying TriboliumDB. For this study, the BeetleAtlas web application was interrogated for gene orthologs of known ion channels and transporters (“Gene” lookup function) as well as for genes enriched in the rectal complex relative to the whole-animal signal (“Tissue” enrichment function), with all candidate genes prioritized according to enrichment. For further information, see SI Appendix, Materials and Methods.

Scanning Electron Microscopy (SEM). SEM analysis of the rectal complex was performed according to a modified protocol described in ref. 58. In brief, rectal complexes were dissected under Schneider’s medium and briefly exposed to a AgNO₃ solution (30 s) as described in ref. 33 before being fixed in 2.5% glutaraldehyde in 0.1M cacodylate buffer (pH 7.4) for 90 min as in ref. 49. The tissue was then desiccated and coated with platinum (70 s B12 nm thickness) and examined with a Zeiss Sigma variable-pressure scanning electron microscope (Carl Zeiss, Oberkochen, Germany) using secondary electron and back-scatter electron detection methods to sequentially visualize both the topology and element weight distribution (atomic number, Z) of the samples. For further information, see SI Appendix, Materials and Methods.

Antibody Generation and Immunolocalization of Target Proteins. To generate anti-NHA1 and anti-VHA55 specific antibodies, we analyzed the amino acid (aa) sequence of the proteins to identify the optimal immunizing peptide region according to a previously described method (63). These antibodies were subsequently used for immunocytchemistry (58) to visualize immunofluorescence in different tissues. Where necessary, immunofluorescence levels were quantified using the FIJI software package from images acquired using identical microscope settings as described in ref. 34. For further information, see SI Appendix, Materials and Methods.

Computational Modeling and NHA1 Structure–Function Predictions. The three-dimensional tertiary structure of NHA1 was predicted by using I-TASSER followed by Ramachandran Plot Assessment (RAMPAGE) to perform structural refinement of the model. Finally, we performed dynamic stability estimations of NHA1 in Tribolium (60), yet most GAL4 drivers may not be entirely specific to particular life stages of individual tissues. Genes with tissue-specific or ontogenetically restricted roles can be easily identified using BeetleAtlas (e.g., SI Appendix, Table S1), and used to make GAL4 driver lines with tissue- or stage-specific activities. Conversely, genes with ubiquitous expression or with persistent roles throughout development can also be found. In sum, we predict that this online resource will have a major impact on the insect functional genomics community by providing an extensive catalog of gene expression across different tissues and life stages in Tribolium, which will promote both an evolutionary and an ontogenetic and tissue-centered view of gene function.

Materials and Methods

Animal Husbandry. Tribolium castaneum (San Bernardino strain) stocks were maintained on organic whole-wheat flour supplemented with 5% (w/w) yeast powder (Tribolium medium) at 30 °C at a constant 50% relative humidity (RH) and 12:12 light–dark cycles as in ref. 61. For further details, see SI Appendix, Materials and Methods.

Tissue Dissection and RNA Extraction. Tissues were dissected and total RNA was extracted from embryos, nonseated 6th instar larvae or 1-wk-old mature adults under a freshly prepared mixture of Schneider’s medium (Invitrogen, CA, US) and Tribolium saline (1:1, v/v) as described in SI Appendix, Materials and Methods.

RNA-seq Analyses and Database Construction. Total RNA libraries were sequenced on a BGISEQ-500 using paired-end chemistry, and subsequent bioinformatic analyses were performed using the Tuxedo pipeline (62). These datasets were used to populate a database, TriboliumDB, that underlies a web application, BeetleAtlas, which is publicly available at www.BeetleAtlas.org. The web application employs a Java servlet to generate web pages and communicate with the TriboliumDB database, and separate smaller servlets for subsidiary functions. It contains a documentation section with full details and version dates. As a web application, BeetleAtlas thus allows nontechnical users to make a wide range of prepared queries to the underlying TriboliumDB. For this study, the BeetleAtlas web application was interrogated for gene orthologs of known ion channels and transporters (“Gene” lookup function) as well as for genes enriched in the rectal complex relative to the whole-animal signal (“Tissue” enrichment function), with all candidate genes prioritized according to enrichment. For further information, see SI Appendix, Materials and Methods.

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the presence of either Na⁺ or K⁺ ions, by submitting the protein for 10 ns molecular dynamics simulation using AMBER18 on the Computerome 2.0 high-performance computing cluster as described in SI Appendix, Materials and Methods.

Molecular Cloning of Nha1. cDNA of Nha1 was synthesized from total RNA extracted from adult T. castaneum rectal complexes using the High-Capacity cDNA Reverse Transcription Kit with RNaseFree Inhibitor (ThermoFisher, MA, USA), and the coding region of the gene was amplified using OS® Hot Start High-Fidelity 2× Master Mix (New England Biolabs, MA, USA) using Nha1-specific primers (SI Appendix, Table S2). The PCR products were subsequently cloned into pGEMHEX X. laevis oocyte expression vector using In-Fusion® HD cloning kit (TaKaRa Bio Inc, Kusatsu, JP) and the final sequence validated (Eurofins, Luxembourg, LU).

pH Measurements in X. laevis Oocytes. Tribolium Nha1 was cloned and heterologously expressed in X. laevis oocytes to characterize the molecular function of the protein as described in detail in SI Appendix, Materials and Methods.

Gene Expression Analysis. Validation of RNAi-mediated gene knockdown (performed 3 d post dsRNA injection) and environmentally induced changes in gene expression (SI Appendix, Table S2) was assessed by quantitative real-time PCR (qPCR) as described in SI Appendix, Materials and Methods.

Production of dsRNA and RNAi-Mediated Knockdown. dsRNA synthesis and knockdown of target gene expression by RNAi was carried out according to a protocol described in SI Appendix, Materials and Methods. Gene-specific primers that were tagged with 17 promoter sequences at both the 3' and 5' ends are listed in Table S2, SI Appendix.

Environmental Stress Exposure. Beetles were subjected to different environmental stressors as described in SI Appendix, Materials and Methods.

Desiccation Tolerance. Desiccation tolerance of control and knockdown animals exposed to different relative humidities was assessed as described in SI Appendix, Materials and Methods.

Hemolymph Collection and Quantification. Hemolymph was collected according to a modified protocol (34, 49, 64) from animals exposed to different environmental stress exposures. See SI Appendix, Materials and Methods for further information.

Quantification of Water Content. Gravimetric estimates of total body water were made by measuring wet and dry body weight of each animal after a given environmental exposure in control and Nha1 knockdown animals; see SI Appendix, Materials and Methods for further details.

Defecation Behavior. The effects of manipulating Nha1 expression on in vivo whole-animal excretory behavior was performed as described in (34) with further details provided in SI Appendix, Materials and Methods.

Ex Vivo Fluid Reabsorption Assay. The water reabsorption rate from the rectal complex ex vivo was assessed using a modified protocol (51) and as described in SI Appendix, Materials and Methods.

Water Vapor Absorption Assay. The ability to extract water vapor directly from the atmosphere was quantified gravimetrically as described in (5). Further details are provided in SI Appendix, Materials and Methods.

Statistics. Data analysis was performed for each experimental condition using relevant methods as described in SI Appendix, Materials and Methods.

Data, Materials, and Software Availability. All study data are included in the article and/or supporting information.

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