Dehydration triggers ecdysone-mediated recognition-protein priming and elevated anti-bacterial immune responses in Drosophila Malpighian tubule renal cells

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Abstract
Background: Drosophila is a powerful model for the study of factors modulating innate immunity. This study examines the effect of water-loss dehydration on innate immune responsiveness in the Drosophila renal system (Malpighian tubules; MTs), and how this leads to elevated host defense and contributes to immunosenescence.

Results: A short period of desiccation-elevated peptidoglycan recognition protein-LC (PGRP-LC) expression in MTs, increased antimicrobial peptide (AMP) gene induction, and protected animals from bacterial infection. We show that desiccation increased ecdysone synthesis in MTs, while inhibition of ecdysone synthesis or ecdysone receptor expression, specifically within MTs, prevented induction of PGRP-LC and reduced protection from bacterial infection. Additionally, aged flies are constitutively water-stressed and have elevated levels of ecdysone and PGRP-LC. Conversely, adults aged at high relative humidity show less water loss and have reduced expression of PGRP-LC and AMPs.

Conclusions: The Drosophila renal system is an important contributor to host defense and can modulate immune responses in an organ autonomous manner, responding to environmental changes such as desiccation. Desiccation primes immune responsiveness by elevating PGRP-LC expression specifically in MTs. In response to desiccation, ecdysone is produced in MTs and acts in a paracrine fashion to increase PGRP-LC expression, immune responsiveness, and improve host defense. This activity of the renal system may contribute to the immunosenescence observed in Drosophila.

Keywords: Drosophila Malpighian tubules, desiccation, ecdysone, innate immune aging, immunosenescence

Background
In Drosophila, the humoral immune response is characterized by the rapid induction of a battery of antimicrobial peptides (AMPs). These endogenous antimicrobials are expressed at nearly undetectable levels in the blood (or hemolymph) of healthy, young adult flies but are transcriptionally induced to massive levels following systemic microbial infection. This response is triggered by microbial cell walls, peptidoglycans from bacteria or beta-glucans from fungi, and is regulated by two nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB) signaling pathways, the Toll and Imd pathways [1, 2].

The Imd pathway responds to diaminopimelic acid-type peptidoglycan (DAP-type PGN) from the cell wall of Gram-negative and certain types of Gram-positive bacteria. DAP-type PGN is sensed by two receptors, both members of the peptidoglycan recognition protein (PGRPs) family of receptors. PGRP-LC is a transmembrane receptor found on the cell surface, while PGRP-LE is a cytosolic receptor [3]. DAP-type PGN ligation of either of these receptors is sufficient to activate Imd signaling and massive AMP gene induction. In particular, these receptors trigger the formation of a functional amyloid signaling platform, involving the factor Imd [4], which in turns leads to K63-ubiquitination, activation of the Drosophila kinases TGF-β activated kinase...
1 and I Kappa B Kinase, and ultimately the cleavage and nuclear translocation of the NF-κB precursor Relish [5–7]. Relish directly induces AMP gene expression.

Developmental and environmental signals modulate the activity and sensitivity of the Imd immune response. In particular, the steroid hormone 20-hydroxyecdysone (20E) controls the expression of the key PGN-sensing receptor PGRP-LC [8]. In earlier work, we established that 20E, signaling through a canonical nuclear hormone receptor pathway, is required for PGRP-LC expression both in cultured cells and in adult flies. Given the central role PGRP-LC plays in sensing systemic bacterial infections, steroid hormone regulation provides a significant degree of modulation on the ability of cells or animals to respond to bacterial infection. It is clear from earlier literature that this regulatory network provides developmental modulation to Imd responses. In particular, early third instar larvae are largely unresponsive to challenge with DAP-type PGN-containing microbes, while wandering larvae, which have received a large bolus of 20E as part of their developmental program, are highly responsive to these infections [9].

In adult flies, it has been suggested that 20E also serves as a stress-responsive steroid hormone, analogous to adrenal-produced, stress-responsive hormones (such as glucocorticoids) in mammals [10–13]. In this study, we have examined how 20E modulates the immune response following the stress caused by a brief period of desiccation. Given the physiological role for the Malpighian tubules (MTs, the insect renal system) in solute homeostasis [14], this immune responsive organ was the focus of investigation. We found that MT autonomously exhibited dehydration-induced steroid-mediated priming of the Imd pathway. This priming response required 20E-mediated upregulation of the innate immune recognition receptor PGRP-LC, thus termed recognition-protein priming. Aged flies are known to naturally become dehydrated [15, 16], and experimental rehydration ameliorated the elevated Imd activity typically observed in older animals, suggesting a link between dehydration-induced recognition-protein priming and immunosenescence. Moreover, dehydration stress-induced steroid-mediated recognition-protein priming, specifically in the MTs, provided enhanced protection to systemic bacterial infection in adult animals. These data demonstrate the important role of steroid-mediated recognition-protein priming in modulating the immune response of adult Drosophila, shaping host defense based on life history and environmental conditions.

Results
Desiccation increases immune sensitivity in the Drosophila renal system
It was previously shown that Drosophila MTs are autonomously immune responsive with robust AMP gene induction following immune challenge in intact animals and in dissected tubules [3, 17, 18]. In MTs and other immune responsive tissues, tracheal cytotoxin (TCT), a monomeric fragment of DAP-type PGN, activates Imd signaling [3, 19]. Thus, to characterize the innate immune response in MTs, we challenged MTs excised from wild-type (wDah) females with TCT and measured AMP gene expression using NanoString nCounter. As shown here, TCT induced robust expression of multiple AMP genes in MTs dissected from 7-day-old females (Fig. 1a, Additional file 1: Figure S1).

In preliminary studies with this ex vivo MT assay, we noticed a high degree of variability in the amplitude of AMP induction. This led us to hypothesize that animals may be exposed to relatively brief periods of dehydration during handling on the CO2 anesthesia ‘fly pad,’ which could be affecting immune response amplitudes. An experimental protocol was designed to directly test this hypothesis. Without anesthesia, 7-day-old females were transferred to vials and maintained in PBS (isotonic to fly hemolymph) or in a dry vial (desiccation) for 2 h (both conditions without food). RNA was isolated from freshly dissected MTs (time 0), and from dissected MTs incubated for 4 h with or without TCT. Defensin and Cecropin-A2 (Cec-A2) mRNAs were quantified. MTs stimulated with TCT strongly induced Defensin and Cec-A2 compared to freshly dissected MTs or to mock-treated MTs (Fig. 1b). For Defensin, desiccation triggered a modest increase in expression in the mock-treated samples and amplified the induction upon TCT treatment (two-way ANOVA, $\chi^2_{(\text{desiccation x TCT})} = 3.96, p < 0.05$). For Cec-A2, the effect of desiccation trended in the same direction, but did not reach statistical significance ($\chi^2_{(\text{desiccation x TCT})} = 1.31, p = 0.31$). In a second similarly designed experiment, RNA was extracted from four biologically independent sets of MT, isolated from control (PBS) or desiccated wDah animals, and analyzed by Nanostring nCounter for the expression of multiple AMPs and other immune related genes (Heat Map in Additional file 2: Figures S2). In this analysis, 13 AMP genes were significantly upregulated either by desiccation alone or upon TCT-challenge following desiccation. Many of these AMP genes showed significant interactions between desiccation and TCT (side table in Additional file 3: Figure S3, after Bonferroni correction for multiple comparisons). Together, these data demonstrate that desiccation robustly induces AMP gene expression within the MT, and frequently increases the sensitivity to TCT.

Desiccation triggers increased expression of the innate immune recognition protein via steroid hormone signaling
In MTs, TCT is a direct agonist of both the cell surface receptor PGRP-LC and the cytosolic receptor PGRP-LE
Desiccation caused a significant increase in PGRP-LC mRNA expressed in MT, while PGRP-LE expression was unchanged (Fig. 1c). Overall, MT from adults held without water for a short duration increased expression of PGRP-LC and their capacity to induce AMP gene expression. Among its many functions [24], ecdysone modulates AMP expression in larvae, cultured cells, pupae, and adults [8, 9, 25–28]. In particular, 20E regulates PGRP-LC expression [8]. Ecdysone also induces expression and synthesis of its nuclear hormone receptor, ecdysone receptor (Ecr) [29]. Here, we found elevated Ecr mRNA levels in MTs of desiccated flies, compared to PBS-treated or conventionally reared animals (Fig. 2a). In addition, ecdysone is known to activate an Ecr-responsive reporter in adult tissues including the MT [30]. Therefore, we tested if Ecr is required for desiccation to induce PGRP-LC in MTs. Ecr RNAi expressed in stellate cells (c724-Gal4) or

Fig. 1 TCT and desiccation stress induce multiple AMP genes in Malpighian tubules (MTs). a Nanostring nCounter analysis of AMP transcripts from isolated MTs of w1118 females (7 days old) exposed to TCT for 4 h compared to unstimulated tubules. The mean of four independent biological replicates, harvested on separate days, is shown, and error bars represent standard error of the mean. Statistical significance was calculated by two-way ANOVA and Sidak’s multiple comparison test; **p < 0.01. b mRNA expression (means with SEM) for Defensin and Cecropin-A2 (Cec-A2) measured by qRT-PCR relative to control (time 0, no TCT). TCT induces Defensin (Likelihood ratio test, DF = 1, χ² = 9.32, p = 0.0023) and Cec-A2 (Likelihood ratio test, DF = 1, χ² = 6.15, p = 0.013). Generalized Linear Model estimated by maximum likelihood with TCT and desiccation as main effects and TCT x desiccation as interaction effect, N = 12; desiccation and TCT significantly interact to amplify Defensin mRNA (TCT x desiccation χ² = 3.96 p = 0.046). The desiccation-induced increase for Cec-A2 did not reach significance (TCT x desiccation χ² = 1.03, p = 0.31). c qRT-PCR analysis of PGRP-LC and PGRP-LE expression from MTs, normalized to PBS-treated control. *p < 0.05; means with SEM; N = 6; t test; PGRP-LC p = 0.038, PGRP-LE p = 0.27
principal cells (c324-Gal4) blocked the ability of desiccation to induce EcR or PGRP-LC expression in MTs (Fig. 2a, b). In a complementary fashion, ecdysone induced protein 75 (Eip75), a negative regulator of ecdysone-mediated signaling, was targeted via tissue-specific RNAi. Knockdown in stellate cells (via c724-Gal4/UAS-eip75B(RNAi)) significantly amplified the ability of desiccation to increase PGRP-LC mRNA relative to control genotype (βgenotype x desiccation = −0.018, χ² = −3.50, p < 0.01), while c324-Gal4/UAS-eip75B(RNAi) trended in the same direction, but did not reach statistical significance (Fig. 2c).

Ecdysone has a myriad of activities beyond its classic role in controlling the insect life cycle. For example, 20E drives PGRP-LC expression in S2* cells [8]. Additionally, 20E fed to adult Drosophila restores normal aging to otherwise long-lived EcR mutant females, restores courtship memory defects and promotes sleep [12, 31, 32]. Here, 20E fed to females induced the expression of EcR and PGRP-LC mRNA within MTs, as well as the expression of Defensin and Cec-A2 even without external immune challenge (Fig. 2d).

Ecdysone signaling appears to drive the expression of PGRP-LC during desiccation stress, and thus prime tubules to express higher levels of AMPs (‘recognition--protein priming’). These results suggest that desiccation stress should increase 20E levels. Measured from whole animals of two genetic backgrounds (yw and wDah), 20E was elevated after 2 h of desiccation (Fig. 3a). To determine where this 20E is produced, the expression of ecdysone biosynthetic enzyme genes (Halloween cytochrome P450s) was quantified from different tissues (Fig. 3b and Additional file 4: Figure S4, all expression was normalized relative to the head sample in the food control group). Desiccation upregulated Halloween genes in several somatic tissues. The most striking effects (Fig. 3b) were observed for disembodied (dib), a cytochrome P450 with ecdysteroid 22-hydroxylase
activity; control-treated females expressed dib strongly in ovaries (via follicle cells [33]) and modestly in fat body, while little dib mRNA expression was observed in other tissues. However, upon desiccation, dib mRNA was significantly increased in muscle and MTs. Accordingly, when dib mRNA was reduced by RNAi in stellate or principal cells (Additional file 5: Figure S5), EcR and PGRP-LC expression was not induced in MTs by desiccation relative to food or PBS controls (Fig. 3c, d).

Together, these data show that desiccation controls ecdysone synthesis within MTs, which locally regulates recognition-protein priming. On the other hand, females maintained for 2 h without food (PBS controls) produced significantly less dib mRNA in the fat body relative to food controls ($p = 0.019$). This effect on fat body expression of dib appears to be due to a starvation response rather than a desiccation response. No significant differences in dib expression in ovaries were observed among treatments.

**Aged flies exhibit desiccation-induced ecdysonemediated recognition-protein priming**

Interestingly, water loss rates and AMP gene expression both increase in aged *Drosophila* [16, 34–36]. In addition to higher AMP expression at baseline, older flies also induce higher levels of AMP gene expression when challenged with live bacteria [28, 35], suggesting that...
recognition receptor expression could be affected by age. Indeed, expression of PGRP-LC as well as EcR was increased in MT from flies at age 40 days (Fig. 4a, b). Experimental dehydration did not further elevate PGRP-LC and EcR mRNAs in aged animals, in contrast to their induction in young flies. Ecdysone titers measured from aged flies were also significantly elevated relative to young females in control (food) conditions, while desiccation increased 20E in both young and old animals (Fig. 4c). These data suggest that aging is associated with recognition-protein priming mediated through elevated ecdysone.

To identify the source of ecdysone in aged flies, tissue was dissected from 40 day females exposed to control (food, PBS) and dehydration treatments. Dib mRNA was quantified relative to young females. Relative to young females, old females expressed dib in similar but muted patterns (Fig. 4d, compared to Fig. 3b). dib was still expressed in ovaries of old, food-reared females, although in old females, fat body showed lower expression under all conditions. Old females under control conditions (PBS) expressed somewhat more dib than young females in head (1.83 fold), muscle (1.71 fold), and gut (2.97 fold) (each tissue p < 0.03). Expression of dib was similarly low in the MT of young and old control females, but significantly induced by desiccation in old MTs (3.5 fold), as observed in young females, and consistent with the ecdysone titer.

Aged females may produce ecdysone because they are intrinsically water stressed. To explore this hypothesis, cohorts of yw adults were maintained in normal food conditions in three levels of relative humidity (RH), namely 20% RH, 40% RH (incubator standard condition), and 80% RH (Additional file 6: Figure S6). Water content was measured in flies sampled from each RH cohort across 7 weeks.

![Figure 4](image-url)
of age (Fig. 5a). Consistent with reports where water loss rates were directly measured in aging *Drosophila* [16], water content per fly declined with age in all cohorts ($\chi^2 = 102.9, p < 0.0001$), and at a significantly greater rate at 20% RH compared to 80% RH ($\chi^2_{(age \times humidity)} = 50.02, p < 0.025$). Lifespan was similar among the 40% and 80% RH cohorts (median life span: 44 and 46 days respectively), but significantly less at 20% RH (median life span: 40 days) (Fig. 5b). Remarkably, elevated RH sharply blunted the increased expression of *PGRP-LC*, *EcR*, and the AMP gene *Defensin* typically observed in the MT of aged adults (Fig. 5c–e). MTs from females at 20% RH strongly upregulated *PGRP-LC*, *EcR*, and *Defensin* by age 6 weeks (age as main effect, each mRNA as independent variable: $\chi^2 > 13.0, p < 0.001$). In contrast, mRNA for *PGRP-LC*, *EcR*, and *Defensin* in cohorts at 80% RH increased to an extent that was significantly less than observed at 20% RH (two-way ANOVA for age and RH, each mRNA: $\chi^2_{(age \times humidity)} > 4.3, p < 0.04$). Thus, innate immune aging is blunted in flies reared at RH that reduces intrinsic water stress, decreasing recognition protein expression and AMP gene levels. This amelioration in water loss and immunosenescence also had a modest, but significant, effect on lifespan.

**Desiccation-triggered ecdysone-mediated recognition-protein priming leads to enhanced host defense**

We were also interested to examine how recognition-protein priming might alter host defense. To test if desiccation-induced recognition-protein priming affects survival following bacterial infection, we exposed two wild-type
fly strains (*wDali* and *yw*) to 2 h desiccation and then immediately infected the adults with the Gram-negative pathogen *Erwinia carotovora carotovora* 15 (*Ecc15*). Desiccated flies were significantly more susceptible to *Ecc15* infection compared to those without prior desiccation (Additional file 7: Figure S7). Given that PGRP-LC expression is elevated upon desiccation, these results were unexpected and led us to hypothesize that some time, following desiccation, may be required for PGRP-LC mRNA to produce functional receptors. Accordingly, we allowed the desiccated flies to recover upon regular food for 3 or 6 h prior to *Ecc15* challenge. Wild-type flies that were allowed to recover for 6 h significantly improved their survival relative to all other infected cohorts, including non-desiccated controls (Fig. 6a–c and Additional file 7: Figure S7). Importantly, this improved survival following septic infection required ecdysone signaling and PGRP-LC expression in the MT; knockdown of *EcR* or of PGRP-LC in MT principal cells eliminated the survival benefit of desiccation with recovery following *Ecc15* challenge (Fig. 6d, e). These results demonstrate that MTs contribute to host defense, and that recognition-protein priming, specifically in this renal organ, improves outcomes following septic bacterial challenge.

![Fig. 6](https://example.com/fig6.png)

**Fig. 6** Functional immunity enhanced by desiccation stress response requires *EcR* and PGRP-LC in Malpighian tubules (MTs): a–e Kaplan–Meier survival plots of adult females sterile pricked or infected with *Ecc15*, with or without prior desiccation (des) and either no recovery, 3 h recovery, or 6 h recovery. Genotypes: a *yw*; b +/*EcR(RNAi); c +/PGRP-LC(RNAi); d c324-gal4/UAS-EcR(RNAi); e c324-gal4/UAS-PGRP-LC(RNAi). Each survival curve is from a combined lifetable of two cohorts tested at different times, each cohort initiated with 90 females. For survival analyses (right censored log-rank test) conducted with six pair-wise comparisons within each genotype; curves with different letters differ significantly by at least *p* < 0.02 (details in Additional file 9: Table S1). f Microbial load (colony-forming units) 48 h after *Ecc15* infection of adult females across treatments by desiccation (0 or 2 h) and post-desiccation recovery (0, 3, 6 h). Genotypes as in (a–e): wildtype controls (*yw*, +/*EcR*, +/PGRP-LC) and *EcR* or PGRP-LC knock-down in principal cells (c324 > RNAi). For each genotype, three independent samples were prepared from each treatment group, in both cohort trials (displaying mean and SD of all six observations). CFU compared by ANOVA with Sidak post-hoc comparison, bars indicate differences with *p* < 0.05, see details in Additional file 10: Table S2.
Improved survival upon septic bacterial challenge can arise from tolerance to the infection as well as from immune defenses that reduce microbial load, often referred to as resistance [37]. To determine whether recognition-protein priming affects tolerance or resistance, we measured Ecc15 colony-forming units (CFUs) from flies treated as in the infection survival studies. Flies were initially inoculated with ~6.0 × 10^5 CFUs of Ecc15 (0 h, Additional file 8: Figure S8). At 48 h post infection, wild-type flies harbored less Ecc15 if they had been desiccated and allowed to recover prior to infection (Fig. 6f), consistent with their improved survival. In contrast, flies with EcR or PGRP-LC expression knock-down specifically in their MT had a higher Ecc15 load at 48 h post infection, and desiccation with recovery did not reduce infection load. Thus, recognition-protein priming in adult MTs provides functional whole animal immune resistance to septic bacterial challenge, leading to improved survival.

**Discussion**

Water-loss dehydration triggers recognition-protein priming within adult *Drosophila* MTs. This response is mediated by the local synthesis and signaling of ecdysone within MTs, which induces expression of the microbial recognition protein *PGRP-LC*. Elevated *PGRP-LC* potentiates the expression of AMP mRNA in tubules. Increased AMP gene expression is expected to produce more mature AMPs, although the study of the peptides is hampered by a lack of effective antibodies. Our analysis of survival after bacterial challenge with desiccation and recovery suggests that recognition-protein priming in the MT provides functional antimicrobial resistance to the entire animal. Moreover, the load of infecting microbe is reduced in wild-type flies able to mount recognition-protein priming, but not in flies where *PGRP-LC* or EcR expression is reduced specifically in MTs, highlighting a previously unknown role for this organ in whole animal antimicrobial defense.

Recognition-protein priming induced by water-loss dehydration may be an adaptive response if dehydration coincides with infection. Observations on this topic are scarce. Virulence in the waxmoth *Galleria mellonella* was increased by infection with *Beauveria bassiana* that exogenously expressed diuretic hormone, which stimulates host water loss [38]. In *Drosophila*, septic infection might increase if water-loss dehydration reduces barrier protection of gut or cuticle [39, 40]. Alternatively, Imd signaling primed by water-loss dehydration could provide novel immune protective or tolerance functions. For instance, Relish (NF-kB) regulates expression of cathepsin-L required for remodeling fat body in the moth *Helicoverpa armigera*, and Har-Relish is modulated by ecdysone signaling in this moth [41].

In *Drosophila*, recognition-protein priming is mediated by ecdysone produced within the MT. Water-loss dehydration induces MT to express ecdysone biosynthetic genes and targeted reduction of dib within MT stellate or principal cells prevented induction of *PGRP-LC* during water-loss dehydration. How MTs regulate dib and other Halloween gene expression in response to water loss is unknown. Desiccation can reduce *Drosophila* neuropeptide capa-1 (homolog of human Neuromedin U) and increase short neuropeptide F and tachykinin in protocerebral neurosecretory cells [42–44]. Capa-1 controls intracellular Ca^{2+} and nitric oxide/guanosine 3’,5’-cyclic monophosphate signaling in MTs [42]. Further, tachykinin induces insulin production and insulin signaling within MTs [45]. Notably, insulin signaling has the capacity to stimulate ecdysone production in the prothoracic gland of developing larvae [46–48]. It remains to be determined whether similar neuroendocrine factors control Halloween gene expression and recognition-protein priming in MTs.

Female *Drosophila* use cholesterol to produce ecdysone in egg follicles. In the fat body, this ecdysone is converted by 20-hydroxylase CYP314A1 (*shade*) to 20E, which activates nuclear hormone receptor EcR and stimulates yolk production [49, 50]. In the ovary, ecdysone also regulates germline stem cells, somatic cysts, and border cells [33]. Other adult tissues in female insects do not produce ecdysone de novo, although some may complete its final synthesis, including the step involving dib [51]. Ecdysone synthesis also occurs in adult males, where its production may take place in accessory glands [52–54]. We find that muscle, gut, and MT, express dib when adults are dehydrated; these tissues may thus produce ecdysone during physiological stress, or at least complete its synthesis. Consistent with this idea, Rauschenbach et al. [10] and Terashima et al. [11] noted that nutrient restriction and heat increased 20E levels in adult *Drosophila*.

Ecdysone has many potential somatic targets within adult *Drosophila* [24, 30]. Notably, 20E can modulate fly aging. Males and females heterozygous for mutant EcR are long-lived, and resistant to the oxidizing agent paraquat as well as to heat and starvation despite being surprisingly fertile [31]. Tricoire et al. [55] subsequently found that mild repression of EcR also extended male lifespan, while EcR inactivation decreased female lifespan except when egg production was genetically suppressed. Besides control of egg production, ecdysone in adults has stress- and aging-related functions, although where and how these work remain poorly understood.

Here, we studied *Drosophila* innate immune aging in the context of desiccation stress. Water loss may be a typical feature of fly aging. Aged *Drosophila* experience elevated water loss and are less resistant to desiccation stress [56, 57], susceptibilities that may be caused by accelerated cuticle water transpiration [16]. Cuticle lipids regulate barrier transpiration in adult insects and this
layer may deteriorate or change composition during aging [15, 58, 59]. We propose that recognition-protein priming and increased AMP response in the aged arise in part because old flies experience high water loss. Consistent with this idea, cohorts maintained at high RH repressed the typical age-related increase in PGRP-LC, Defensin, and EcR expression. Conversely, flies at low RH were somewhat shorter lived, and previous work suggests that chronic innate immune activation reduces life expectancy in an NF-xB-dependent manner [60].

The production of AMPs is costly, impacting physiology and fecundity [61], and is tightly regulated. AMP induction is robust but transient to minimize the deleterious effects of immune activation. Likewise, the Imd pathway, and AMP production, is regulated by ecdysone in the context of development and, as shown here, by stress. This steroid-mediated regulation likely optimizes the AMP response such that the benefits of this mechanism of host defense at least equal these costs. Steroid modulation of AMP induction during development may coordinate this feature of innate immunity with nutrient demands and availability, which change during different growth phases. Alternatively, developmental steroid regulation of AMP production may be an adaptation to the selective pressures associated with the different microbial pathogens encountered with different developmental stages. Furthermore, here we demonstrate that the adult flies use ecdysone to prime innate immune induction of AMPs in the renal organ in response to water-stress imbalance. Remarkably, the resulting elevated expression of AMPs from these renal tissues appears to protect adult animals from systemic, septic infection.

**Conclusions**

In this study, we examined how water-loss dehydration increases innate immune responsiveness in *Drosophila* MTs, and how this response improves outcomes following infection and contributes to elevated innate immune gene expression with age. In this model, the stress of water-loss dehydration increased systemic levels of 20E, which is responsible for the pattern recognition-protein priming observed in MTs. While it is not possible to measure 20E levels in a tissue-specific manner, genetic manipulations of both the 20E synthesis gene *dib* and the *EcR*, demonstrate that the tubules are autonomously responding to water-loss dehydration through the production of ecdysone and subsequent sensing of this hormone signal. Interestingly, the data presented here further argue that MTs serve a critical function in anti-bacterial immunity, wherein recognition-protein priming in MTs is required for increased host defense observed after desiccation and recovery, and loss of recognition-protein expression specifically in MTs caused a marked sensitivity to infection. How MTs contribute to immunity and host defense will be the focus of future studies.

In addition, aged flies also display symptoms consistent with water-loss dehydration. Aged flies, reared at 20%, 40%, or 80% RH, were dehydrated, consistent with earlier reports [58]. Aged flies also displayed increased systemic 20E levels and increased expression of the innate immune recognition protein PGRP-LC and effectors. However, adults aged at high relative humidity show less water loss, and have reduced expression of PGRP-LC, AMPs, and EcR in their tubules. Our study suggests that the steroid hormone ecdysone in old flies is induced in response to age-related water imbalance, and this facilitates immunosenescence in *Drosophila* renal MTs.

**Methods**

**Drosophila strains**

Stocks were reared and maintained at 25 °C, 40% RH and 12-h light/dark cycles. Food media contained cornmeal (5.2%), sugar (11.0%), autolyzed yeast (2.5%; SAF brand) and agar (0.79%) (w/v in 100 mL water) with 0.2% Tegosept (methyl 4-hydroxybenzoate, Sigma, St. Louis, MO, USA) as an antifungal agent. MT-specific Gal4-driver stocks c724 (stellate cell) and c324 (principal cell) were provided by Julian Dow (University of Glasgow); second chromosomes containing these transgenes were crossed and balanced into a genetic background marked with *yw*, producing *yw; c724-Gal4/Cyo and yw; c324-Gal4/Cyo*. RNAi to reduce PGRP-LC, EcR, *Eip75B*, and *dib* was respectively provided from Vienna Stock Center as strains P[UAS-PGRP-LC*KK101636 RNAi*]-VIE-260B, P[UAS-EcR*GD1428 RNAi*]-VIE-260B, P[UAS-Eip75B*KK108982 RNAi*]-VIE-260B, and P[UAS-dib*KK106954 RNAi*]-VIE-260B. Genotypes (F1 offspring) were generated from crosses of these UAS-RNAi stocks with MT Gal4-drivers or by crosses with *yw*. *wDah* was provided by the laboratory of Linda Partridge (University College London).

**20E treatment**

20E was fed to *wDah* females aged 7–10 days old by transferring adults from standard laboratory diet to 1 g fly instant food media with 2 mL of water (Genesee Scientific, San Diego, CA, USA) for 24 h containing vehicle control (20 μL ethanol) or vehicle with 20 μL 1 mM 20E (Sigma cat H-5142).

**Desiccation protocol**

Prior to conducting a trial, adult flies were placed in vials with standard food at a density of approximately 15 per vial without anesthesia. Vials were maintained overnight at 25 °C. To initiate each trial, flies from vials were...
flipped without anesthesia into fresh vials that contained standard food (food control), tissue soaked with 0.5 mL 0.5× PBS solution (PBS control), or nothing (desiccation). MTs were dissected after 2 h of treatment from flies within 5 minutes, immediately transferred to Trizol, and processed for RT-qPCR. Three replicate biological samples of 15 MT (sampled from across five vials) were generated per treatment.

**MT stimulation with TCT**
Females (*wDah*) aged 7 to 10 days post eclosion, were processed through the desiccation protocol (with PBS as the only control). Dissected MTs were transferred for 4 h to S2 cell culture medium with TCT (0.01 mM dissolved in water) or without TCT. Expression of *defensin* and *Cec-A2* was measured by qRT-PCR from RNA extracted from these tubules immediately upon dissection (0 h), and at 4 h with or without TCT. Three replicate biological samples were produced for each measure set.

**NanoString nCounter expression analysis**
The mRNA content of dissected MTs was analyzed by NanoString methodology according to published procedures [62]. Total RNA (100 ng per sample) was hybridized to the target-specific codeset overnight at 65 °C. The codeset used in Additional file 1: Figure S1 contained probes against a panel of 45 genes encoding AMPs, Imd pathway components, and four loading controls (*nucleostemin, alpha tubulin, Rp49*, and *Faf2*), while that in Additional file 2: Figure S2 utilized an expanded codeset of 113 immunity-related genes and three loading controls (*nucleostemin, alpha tubulin, and Faf2*). The hybridized reactions were loaded onto the NanoString Prep station, which removes excess reporter, binds the reporter to the cartridge surface, and stretches the probes for scanning. Subsequently, the cartridges were loaded onto the NanoString Digital Analyzer and scanned. The nCounter data were normalized in two steps. In the first, data was normalized to positive spiked-in controls, as per the manufacturer's instructions (NanoString Technologies). In the second, the housekeeping genes were used to normalize overall mRNA levels. For Fig. 1a, AMP gene expression was further normalized amongst the four biological replicates by setting the highest value for each gene to 100% and plotted with standard error of the mean. For heat maps in Additional file 1: Figure S1 and Additional file 2: Figure S2, only genes with expression greater than 50 normalized counts in one or more conditions are shown, as lower counts are below the reliable detection threshold for the nCounter.

**Quantitative RT–PCR**
Total RNA was extracted from dissected tissue (MT, head, muscle, gut, fat body, ovary) in Trizol reagent (Invitrogen, Carlsbad, CA, USA) and treated with DNase (Ambion). DNase-treated total RNA was quantified with a NanoDrop ND-1000. cDNA was synthesized using iScript cDNA Synthesis (Bio-Rad) and measured on an ABI prism 7300 Sequence Detection System (Applied Biosystems, Carlsbad, CA, USA). For tissue-specific analysis of AMP, PGRP-LC, EcR, or Halloween genes, three independent biological replicates for each tissue and each treatment group were analyzed. Each replicate included RNA extracted from 10 flies for head, muscle, gut, fat body, and ovary, or 15–20 for MT. Three technical replicates were used for each biological replicate. mRNA abundance of each gene was normalized relative to ribosomal protein L32 (*rp49*) by comparative CT.

**Enzyme immunoassay (EIA) of 20E**
Female flies (25–30 flies/sample) were homogenized in 250 µL of ice-cold 100% methanol and centrifuged for 15 min at 18,000 g at 4 °C. Supernatants were transferred to 6 × 50 mm borosilicate glass tubes; precipitates were suspended in 200 µL of aqueous 75% methanol and kept on ice for 30 min (adapted from [63]). Samples were centrifuged and dried with a SpeedVac centrifuge. The precipitates were dissolved in EIA buffer at 4 °C and each individual sample was analyzed in a technical triplicate by competitive EIA using 20E EIA antiserum (Cayman Chemical: 482202) and 20E AChE Tracer (Cayman Chemical: 482200). Calibration curves were generated from commercial 20E (Sigma, H5142).

**RH cohorts**
Newly enclosed females (*yw*) were mated for 2 days and separated into replicate 1 L demography cages at 125 flies per cage in an environmentally controlled room set at 40% RH, 25 °C, and 12 h light: 12 h dark cycles. Three independent cages (plus supplemental cages for dissection samples) were placed in 60 L clear plastic, tight-lidded storage boxes that maintained RH at 20%, 40%, and 80%. To produce 20% RH, boxes contained a solution of saturated potassium acetate (Fisher Science Cat # P171–500) and solid desiccant (Drierite, W.A. Hammond Co.). To produce 40% RH, the box was left ajar, permitting cages to equilibrate at 40% RH as maintained in the environmental room. To produce 80% RH, boxes contained a solution of saturated ammonium sulfate (Sigma-Aldrich Cat # A4418-500G). Boxes were managed every second day at room temperature and room RH, at which time dead flies were removed and counted. Saturated solutions (and Drierite) were replenished weekly. HOBO data loggers (Onset Computer Corp) continuously monitored RH within boxes; after
maintenance, all boxes rapidly re-equilibrated to RH targets (Additional file 5: Figure S5).

Longevity analysis was conducted with JMP statistical software (SAS Institute, Cary, NC, USA). Data from three replicate cages were combined to produce genotype and treatment cohort life tables. Mortality distributions were compared by Log-Rank and proportional hazard analyses. Flies to measure mRNA were maintained in parallel demography cages and sampled at 1, 2, 4, and 6 weeks. MTs were dissected directly from non-anesthetized adults and subjected to RT-qPCR (15 females per biological replicate, three replicates per time point per humidity condition).

Water content was determined from samples in a replicate trial with cohorts at 20%, 40%, and 80% RH. Females were sampled at 1, 2, 4, 6, and 7 weeks from each RH cohort. Point estimates for water content were each generated as the mean among five groups of 20 females; these females were sampled from 8 demography cages, each with 125 females for every RH condition. For each group, females were weighed (Mettler Toledo semi-micron balance), dried overnight at 65 °C, and reweighed. Age, RH treatment, and main-effect interactions were analyzed by generalized linear model likelihood ratio test.

Infection challenge survival
Adult survival to a controlled infection challenge after desiccation treatment was determined using 100 to 120 flies per group (Additional file 5: Figure S5) or 180 flies per group (Fig. 6) (among two replicate vials). Females at 3 to 7 days old were exposed to food or PBS-only as control, or to 2 h desiccation. Following these treatments, either immediately or after 3 and 6 h recovery on standard food, flies were pricked with a clean microsurgery needle (control) or with a needle dipped into a concentrated pellet of (Ecc15) [64, 65]. Surviving flies were subsequently transferred daily to fresh vials and recorded as alive or dead for 8 days. Survival is represented as Kaplan–Meier plots with all pairwise differences evaluated by log-rank test with significance adjusted for multiple comparisons ($p < 0.01$).

CFU counts
Bacterial load was determined in flies at time 0 and 48 h after infection with Ecc15. Individual flies were homogenized in 200 μL of PBS. Homogenates were diluted in series and plated on LB-Ampicillin plates, incubated overnight at 37 °C, and scored for CFU count.

Additional files

**Additional file 1:** Figure S1. Heat map of differentially expressed innate immune transcripts in Malpighian tubules (MTs). Nanostring nCounter analysis of AMP transcripts from isolated MTs of wDah females (7 days old) exposed to TCT for 4 h compared to unstimulated tubules. The mean of four independent biological replicates, harvested on separate days, is shown by heat map. Scale as indicated in side bar. (PDF 797 kb)

**Additional file 2:** Figure S2. Heat map summarizing innate immune gene expression profiles in desiccated Malpighian tubules (MTs). MTs of 7-day-old wDah flies exposed to 2 h desiccation or PBS control treatment were excised, and then either RNA was immediately isolated, or were treated for 4 h with TCT or mock treated, and then RNA was isolated. These RNA samples were used to measure immune-related gene expression by NanoString nCounter. Values shown by heat map represent the mean of four independent experiments. Scale as indicated in side bar. (PDF 847 kb)

**Additional file 3:** Figure S3. Desiccation stress upregulated and amplified TCT-induced expression of 13 AMP genes in Malpighian tubules. Analysis of individual AMP gene expression (as indicated), data from nCounter experiments displayed by heat map in Additional file 2: Figure S2. Values represent the mean of four biologically independent replicates, and error bars are standard error of the mean. Statistical analysis was performed by two-way ANOVA and Sidak’s test for pair-wise comparison of desiccation to PBS treatment (*$p < 0.05$, **$p < 0.01$, ***$p < 0.001$, ****$p < 0.0001$). Side table displays statistical metrics after Bonferroni correction for multiple comparisons for the interaction of desiccation and TCT treatments from the same ANOVA analyses. (PDF 631 kb)

**Additional file 4:** Figure S4. mRNA expression of Halloween genes. Halloween genes (spook, phantom, shadow, and shade) were measured in adult tissues from young (7 days) and old (40 days) females. All values were normalized relative to head samples from young adults in food control group. The mean of three independent biological replicates is shown. (PDF 2424 kb)

**Additional file 5:** Figure S5. RNA on disembarbed (db) in Malpighian tubules (MTs). Reduced db mRNA in MTs when db RNAi is driven in stellite (c324 > db RNAi) or principal (c324 > db RNAi) cells relative to control (yw/T1/db RNAi). Results shown represent the mean and SEM of three independent replicates. (PDF 897 kb)

**Additional file 6:** Figure S6. Relative humidity (RH) and temperature recorded from demography chambers under different humidity conditions. Spikes indicate when chambers were opened to access cages and show rapid homeostasis of the humidity control system. Blue tracings show realized RH, black tracing represents temperature. (PDF 3450 kb)

**Additional file 7:** Figure S7. Impact of desiccation followed by recovery on the survival of wild-type flies infected with Erwinia carotovora carotovora 15 (Ecc15). Kaplan–Meier survival of yw and wDah adult females challenged with Ecc15. Uninfected flies were desiccated or not for 2 h and survival was monitored for 8 days. Other cohorts were challenged with Ecc15 infection with or without a prior 2 h desiccation treatment and recovery of 0, 3, or 6 h prior, as indicated. Plots represent the survival kinetics of 100–120 flies, combined from two separate trials, for each genotype; cohorts with significantly different mortality are grouped with different letters (all cases, log-rank test $p < 0.05$, see side tables). (PDF 1741 kb)

**Additional file 8:** Figure S8. Initial bacterial loads in Erwinia carotovora carotovora 15 (Ecc15)-infected flies. Adult load (colony-forming units) of Ecc15 at 0 h post infection as a function of desiccation (0 or 2 h) and post-desiccation recovery (0, 3, 6 h), among three control genotypes (yw, +EcR, +PGPR-LC) and genotypes where EcR or PGPR-LC were knocked down by RNAi in principal cells (c324 > RNAi), no significant differences were observed at 0 h. Results represent the mean of six assays and the error bars show standard deviation. (PDF 1024 kb)

**Additional file 9:** Table S1. Statistical analysis for the effect of desiccation on survival to Ecc15 infection with or without recovery treatment, supporting Fig. 6a–e. (DOXX 26 kb)

**Additional file 10:** Table S2. Statistical analysis for the effect of desiccation on bacterial growth after Erwinia carotovora carotovora 15 infection with or without recovery treatment, supporting Fig. 6f. (DOXX 15 kb)
Abbreviations
20E: 20-hydroxyecdysone; AMP: antimicrobial peptide; Cec-A2: Cecropin-A2; CFU: colony-forming unit; DAP-type PGN: diaminopimelic acid type-peptidoglycan; dib: disembedded; Ecc15: Ermrina carotovora carotovora 15; Ecr: Ecdysone receptor; Epo75: ecdysone induced protein 75; MTS: Malpighian tubules; NF-κB: nuclear factor kappa-light-chain-enhancer of activated B cells; PGRP: Peptidoglycan recognition protein; RH: relative humidity; TCT: tracheal cytotoxin

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Availability of data and materials
All data generated or analyzed during this study are included in this published article and its supplementary information files. All material used, such as Drosophila strains, are available from public stock centers and other repositories.

Authors’ contributions
W2 designed, performed, and analyzed experiments. FR designed, performed, analyzed experiments, and contributed to the drafting and revising of the manuscript. AH performed and analyzed some of the experiments presented in Fig. 1. PK designed and performed RH controlled experiments. WG provided critical reagents. NS and MT together conceived of this study, contributed to experimental design and analysis and manuscript preparation, and supervised all aspects of this study. All authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

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