RESEARCH ARTICLE

A conserved role of the insulin-like signaling pathway in diet-dependent uric acid pathologies in *Drosophila melanogaster*

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Elevated uric acid (UA) is a key risk factor for many disorders, including metabolic syndrome, gout and kidney stones. Despite frequent occurrence of these disorders, the genetic pathways influencing UA metabolism and the association with disease remain poorly understood. In humans, elevated UA levels resulted from the loss of the urate oxidase (*Uro*) gene around 15 million years ago. Therefore, we established a *Drosophila melanogaster* model with reduced expression of the orthologous *Uro* gene to study the pathogenesis arising from elevated UA. Reduced *Uro* expression in *Drosophila* resulted in elevated UA levels, accumulation of concretions in the excretory system, and shortening of lifespan when reared on diets containing high levels of yeast extract. Furthermore, high levels of dietary purines, but not protein or sugar, were sufficient to produce the same effects of shortened lifespan and concretion formation in the *Drosophila* model. The insulin-like signaling (ILS) pathway has been shown to respond to changes in nutrient status in several species. We observed that genetic suppression of ILS genes reduced both UA levels and concretion load in flies fed high levels of yeast extract. Further support for the role of the ILS pathway in modulating UA metabolism stems from a human candidate gene study identifying SNPs in the ILS genes AKT2 and FOXO3 being associated with serum UA levels or gout. Additionally, inhibition of the NADPH oxidase (NOX) gene rescued the reduced lifespan and concretion phenotypes in *Uro* knockdown flies. Thus, components of the ILS pathway and the downstream protein NOX represent potential therapeutic targets for treating UA associated pathologies, including gout and kidney stones, as well as extending human healthspan.
Enzymatic purine degradation in humans ends with uric acid (UA). Multiple genetic and dietary factors raise UA levels above the norm, which is called hyperuricemia or hyperuricosuria when detected in the serum or urine, respectively. Clinical studies report a correlation between elevated UA and a plethora of chronic diseases including crystalopathies like UA kidney stones and gout, or metabolic and vascular disorders such as diabetes, obesity, and coronary artery disease. Here, we identified a regulatory role for the insulin-like signaling cascade affecting UA metabolism using a Drosophila melanogaster model. In the process we determined previously unrecognized potential drug targets to treat elevated UA levels and associated pathologies such as gout or UA kidney stones, with the potential additional benefit of extending human healthspan. Our work also establishes the fly as a model system to characterize the influence of genetic and dietary factors in gout or UA kidney stone development in a manner readily amenable for small-scale screening of drug interventions. The novelty of our findings, their broad impact, and relevance for multiple diseases opens up an important area of research to define mechanisms of UA accumulation.

**Introduction**

Purine homeostasis represents a conserved metabolic pathway that is sustained by multiple enzymes orchestrating de novo synthesis, salvage, and degradation of purine intermediates. Urate oxidase, which is conserved across species, catalyzes one of the last steps of purine degradation converting uric acid (UA) to allantoin. However, due to multiple point mutations in the urate oxidase gene (Uro) human ancestors lost the ability to synthesize a functional urate oxidase, thus, increasing serum and urinary UA levels [1–5]. While most mammals show serum UA levels of 1 mg/dl and lower, healthy humans generally are in the range of 4–6 mg/dl, close to the UA solubility limit of 6.8 mg/dl at physiological pH and body temperature [6, 7]. Despite a regulatory network balancing UA production and excretion, UA levels increase due to both age and a nutrient-rich diet [8]. Several genetic risk factors are associated with increased UA levels including genes of purine homeostasis, glucose metabolism, or UA transporters. Dietary risk factors are sugar-sweetened beverages, alcohol, red meat, and seafood, all found in over-abundance in the Western diet [8, 9].

In humans, elevated UA levels in the blood (hyperuricemia) or urine (hyperuricosuria) are key risk factors for crystalopathies such as gout and kidney stones, metabolic syndrome, as well as premature death and a higher all-cause mortality risk [10–16]. Alarmingly, the prevalence of hyperuricemia in the US population is over 20% [17, 18]. Different treatment options are commonly used to lower UA levels. As first-line therapy, an inhibitor of xanthine dehydrogenase such as allopurinol is prescribed to prevent the xanthine dehydrogenase mediated oxidation of xanthine to UA. Further options include the use of uricosuric drugs such as probenecid to increase renal UA excretion or administration of a recombinant urate oxidase enzyme called pegloticase degrading extracellular UA to the water-soluble allantoin [19, 20]. Common to the UA lowering therapies are the well-documented issues of adverse drug reactions, contraindications, relevant drug-drug interactions, and the need for anti-inflammatory prophylaxis [9]. Interestingly, cohort studies also link exceptional longevity and a reduced prevalence of age-related diseases with UA levels at the lower end of the human serum UA spectrum [21, 22]. In sum, the data correlate elevated UA levels with a higher prevalence of
crystalopathies as well as a shortened healthspan and lifespan. However, the molecular mechanisms by which UA mediates negative health outcomes are still a matter of debate.

_Drosophila melanogaster_ provides an excellent model system to study both crystalopathies and lifespan [23–27]. Crystalopathies in flies can be induced either by feeding lithogenic agents such as ethylene glycol [28–32] or via genetic inactivation of xanthine dehydrogenase [33]. The resulting crystals usually accumulate in the Malpighian tubules, the invertebrate homolog of the human kidney convoluted tubules, and therefore recapitulate the clinical pictures of calcium oxalate urolithiasis and xanthinuria, respectively [34]. In addition, aging studies employing _Drosophila_ take advantage of its short lifespan and fast development. Many reports highlight the relevance of nutrient-sensing pathways such as the insulin-like signaling (ILS) cascade in extending the organism lifespan [35–37]. Yet, there is currently no _Drosophila_ model addressing the impact of elevated UA levels on crystalopathies and lifespan. Therefore, we designed a fly model to study the effects of elevated UA levels by recapitulating both aspects encountered in hominoid evolution: depletion of urate oxidase activity combined with changing dietary habits. Our model showed a diet-dependent accumulation of UA and shortening of lifespan. Using this model, we identified the evolutionarily conserved ILS pathway and the downstream factors FoxO and NADPH oxidase to reduce UA and its associated pathologies which might provide new targets for UA lowering therapies.

**Results and discussion**

**Uro knockdown shortens lifespan in a diet-dependent manner in Drosophila melanogaster**

Unlike humans, _Drosophila melanogaster_ uses the enzyme urate oxidase to convert UA into allantoin (Fig 1A). Therefore, we “humanized” _Drosophila_ to recapitulate the lack of functional urate oxidase making UA the end-product of purine catabolism. We used the heterologous, ligand (RU486)-induced gene switch system to generate spatially and temporally precise RNAi-mediated knockdown of _Uro_ in _Drosophila_ [38]. Thus, flies carrying the ubiquitous _daughterless_ gene switch _GAL4_ driver (DaGS) were crossed to flies with one of the two different UAS- _Uro_-RNAi responder transgenes (_Uro_-RNAi #1 or #2). 2–3 day-old progeny (DaGS > _Uro_-RNAi) of such crosses were then fed diets supplemented with the Gal4 activating ligand RU486 (+ RU) to induce the _Uro_ knockdown or the corresponding vehicle control (- RU).

Compared to the isogenic control siblings, flies consuming the + RU diet showed for both RNAi constructs a 70% knockdown of the _Uro_ mRNA (Fig 1B). To mimic the change in dietary habits of Western civilization flies were fed diets with different concentrations of yeast extract. In comparison to their isogenic control siblings, i.e. DaGS > _Uro_-RNAi #1 flies reared on a diet without RU486, the knockdown of _Uro_, i.e. DaGS > _Uro_-RNAi #1 flies fed a diet supplemented with RU486, did not affect the median or total lifespan of flies consuming a low yeast (LY) diet containing 0.5% yeast extract (Fig 1C and 1D, S1 Fig). However, when _Uro_ knockdown flies were fed a high yeast (HY) diet containing 5% yeast extract their lifespan was significantly shorter compared to the control siblings receiving the same food, but without RU486 supplementation (Fig 1C and 1D). We also compared the median lifespan of DaGS > _Uro_-RNAi #1 flies on different yeast concentrations, showing that the impact of urate oxidase depletion on survival becomes apparent at a dietary yeast concentration of 3% and above (S1 Fig.). At 3% yeast concentration the median lifespan of control flies (DaGS > _Uro_-RNAi #1 flies reared without RU486) was 48 days, whereas the median lifespan of their _Uro_ knockdown siblings (DaGS > _Uro_-RNAi #1 flies reared with RU486) was 31.5 days, a reduction of 34%. Similarly, at 4% yeast concentration the median lifespans of control and knockdown flies were 45 and 27 days, respectively, a decrease of 40%. At 5% yeast concentration the difference in
median lifespan was even more pronounced comparing 39.4 days for control flies and 20.5 days for the Uro knockdown flies, which represented a shortening of 48%. The latter value represented the plateau, given that a higher dietary yeast concentration of 8% did not further augment the difference in median lifespan between control (38 days) and Uro knockdown (20 days) flies (S1 Fig). In other words, we observed an inverse correlation between the dietary yeast content and the median lifespan with the Uro knockdown flies being more susceptible to higher yeast concentrations than controls.

To confirm our findings and demonstrate tissue-specificity as well as independence of the lifespan effect from the RU486 ligand, we used the ubiquitous (Da-GALA4) or Malpighian tubule-specific (c42-GAL4, Uro-GAL4) driver lines not relying on RU486 for activation. Considering that Malpighian tubules are the main tissue expressing Uro the neuronal Elav-GAL4 driver was used as a negative control [39]. For the four non-gene switch driver lines crosses to the Uro-RNAi and w1118, the genetic background flies used to outcross all driver and other transgenic lines, were performed in parallel. As expected, compared to the individual w1118 control crosses efficient Uro knockdown was achieved only when using the Da-GAL4, c42-GAL4 and Uro-GAL4 drivers (Fig 1E). Coinciding with the Uro knockdown, the shortened lifespan of flies consuming the HY diet was only observed with the driver lines actively reducing Uro expression but was absent in case of the Elav-GAL4 driver (Fig 1F). Importantly, flies with reduced Uro expression were significantly longer lived on low nutrient diets with 0.5% and 1% yeast extract compared to diets with a higher yeast extract content (Fig 1C and 1D, S1 Fig), demonstrating the importance of diet in UA mediated effects on lifespan.

Uro knockdown enhances UA accumulation in a diet-dependent manner in Drosophila

Next, using Uro knockdown flies (DaGS>Uro-RNAi #1 + RU486) in comparison to their isogenic control siblings (DaGS>Uro-RNAi #1 - RU486) metabolomic profiling verified the elevation of UA levels after feeding the HY diets for 14 days. The fivefold increase in UA levels in the Uro knockdown flies was accompanied by an almost complete loss of allantoin production (Fig 2A). In accordance with the qRT-PCR (Fig 1B), these data indicate on the metabolite level the efficiency of urate oxidase depletion (Fig 2A). An unbiased micro-CT scan revealed radiopaque masses (putative ‘concretions’) specifically in the abdominal area of Uro knockdown flies, but not in the control population (Fig 2B). This observation was further validated by optical microscopy of dissected animals revealing the presence of concretions in the Malpighian tubule and hindgut of Uro knockdown flies (Fig 2C). Due to the restricted expression pattern of the Uro gene to the excretory system both the Malpighian tubule and the hindgut represent the anatomical sites where UA is most likely expected to accumulate [40]. Furthermore, metabolomic analyses of isolated concretions (Fig 2D) confirmed UA as the predominant
Diet-dependent uric acid pathologies in *Drosophila*

![Diagram A](image1)

Relative metabolite level (
(HY + RU / HY - RU)

![Diagram B](image2)

DaGS>Uro-RNAi #1

![Diagram C](image3)

DaGS>Uro-RNAi #1

![Diagram D](image4)

Concretion (%)

![Diagram E](image5)

Relative concentration (%)

![Diagram F](image6)

Flies with concretion (%)

![Diagram G](image7)

Flies with concretion (%)
component, accounting for 95% of the metabolites measured (Fig 2E). We found enhanced UA aggregation with increasing age (Fig 2F). While DaGS>Uro-RNAi flies without RU486 supplementation formed almost no concretions, in Uro knockdown flies their appearance was observed as early as 4 days of feeding the HY diet. The proportion of flies forming UA deposits increased gradually over time with almost 70% of Uro knockdown flies forming concretions after 14 days (Fig 2F). In contrast, DaGS>Uro-RNAi #1 flies fed the LY diet, with or without RU486, hardly formed any concretion (Fig 2F). Very much like the lifespan phenotype, we investigated the concretion formation with the set of non-gene switch drivers used before. Again, in comparison to the w1118 control crosses performed in parallel a high rate of concretion formation was only detectable on a HY diet when the Uro-RNAi construct was expressed by the Da-GAL4, c42-GAL4 or Uro-GAL4 driver, thus causing efficient urate oxidase depletion and UA accumulation (Fig 2G). The w1118 or Uro-RNAi crosses with the Elav-GAL4 driver did not result in concretion formation underscoring the tissue-specificity of the effect. Using the high proportion (~ 70%) of Uro knockdown flies developing concretions after feeding the HY diet for 14 days as a reference, we analyzed the dietary yeast concentration as factor in promoting concretion formation. Like the lifespan attenuation, concretion formation started to increase significantly in Uro knockdown flies when the dietary yeast concentration reached 3% or more (S2A Fig). Concretion formation was also seen when comparing the control siblings and knockdown flies of the DaGS>Uro-RNAi #2 population (S2B Fig). However, DaGS>Uro-RNAi #2 flies reared on the HY diet without RU486 supplementation showed a much higher background in terms of concretion formation (cf. S2A and S2B Fig). In analogy to human pathophysiology the shortened lifespan and enhanced concretion formation of Uro knockdown flies match with the enhanced mortality in patients with gout as well as the age- and diet-dependent occurrence of crystalopathies in humans [16, 41–43].

The dietary purine content mediates the lifespan and concretion phenotypes of Uro knockdown flies

To determine what component of the HY diet triggered increased UA production and associated phenotypes, we systematically supplemented the ‘benign’ LY diet with various basic nutrient groups: purines, pyrimidines, proteins, or sugar. Of the components tested, we found that purines (metabolic precursors of UA), but not pyrimidines, proteins, or sugar led to a dose-dependent increase in concretion formation when added to the LY food (Fig 3A). Dietary purine supplementation above 10 mM caused a significantly higher concretion formation in Uro knockdown flies (DaGS>Uro-RNAi #1 with RU486) compared to their isogenic controls (DaGS>Uro-RNAi #1 without RU486). Interestingly, purine supplementation of 20 and 40 mM even caused concretion formation to rise in the DaGS>Uro-RNAi #1 flies without RU486 (Fig 3A), thereby showing the harmful and lithogenic potential of dietary purines. Given that the purine supplemented LY diet phenocopied the concretion formation seen with the HY diet, this strongly suggests an important role of dietary purines in promoting and maintaining UA aggregation and concretion formation.
Fig 3. Dietary purine mediates lifespan attenuation and UA concretion formation. (A) Concretion formation of DaGS>Uro-RNAi #1 flies reared on a HY or LY diet for 14 days. Where indicated, the latter was supplemented with increasing concentrations of protein (Prot), sucrose (Suc), purine (Pu) or pyrimidine (Py). Each food was generated in a - RU and + RU version to compare concretion formation of isogenic control and Uro knockdown siblings, respectively. (B) Survival curve of DaGS>Uro-RNAi #1 flies fed the LY diet without or with 40 mM purine supplementation (+ Pu) in presence or absence of RU.

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diet, we examined the impact of purines on lifespan. Intriguingly, addition of 40 mM purines to the LY diet (LY+Pu) recapitulated the lifespan effects seen with the HY diet (cf. Figs 3B and 1C). I.e., purine addition caused a) a shortening of the lifespan of control flies and b) this lifespan attenuation was more pronounced in Uro knockdown flies, which are unable to convert the increased UA load to allantoin (Fig 3B). Thus, altering the ability to utilize purines could be an effective way to prevent UA associated pathologies in flies and strengthen the relevance of the Uro knockdown model as translational research tool.

Manipulation of purine homeostasis rescues the phenotypes of Uro knockdown flies

Purine homeostasis is controlled through orchestrated steps involving de novo synthesis, salvage, and degradation of purine intermediates (S3A Fig). Two drugs, allopurinol and methotrexate, are well-known to effectively perturb purine metabolism. Allopurinol is commonly used in the treatment of gout in humans and prevents purine degradation by inhibiting xanthine dehydrogenase (XDH), an enzyme that converts xanthine and hypoxanthine to UA. Methotrexate interferes with de novo purine synthesis by inhibiting dihydrofolate reductase (DHFR), an enzyme required for folate production (S3A Fig). Adding increasing amounts of allopurinol (Fig 4A) or methotrexate (Fig 4B) to the HY diet consumed by Uro knockdown flies caused a dose-dependent reduction of the concretion formation. At concentrations of 10 mM allopurinol or 50 μM methotrexate concretion formation was almost completely suppressed. We also determined if the effects of drug supplementation on food palatability and its consumption confounded our results. We compared food consumption of flies by a dye-colored food intake assay after them being fed a certain diet for 14 days. We did not observe a statistically significant change in food intake of Uro knockdown flies in case the HY diet was supplemented with allopurinol or methotrexate, when analyzed by ANOVA and Tukey’s multiple comparison post-test (S3B Fig). Next, we used a genetic strategy to verify known and identify novel targets that influence UA pathologies. Therefore, we generated a recombinant fly strain that harbors the DaGS driver and the Uro-RNAi #1 transgene, hereafter denoted by DaGS::Uro-RNAi. We confirmed that the recombinant DaGS::Uro-RNAi line when crossed to either w1118 or b35785 (a strain that carries a no target UAS-mCherry-RNAi construct) still forms concretions when fed the RU486 supplemented HY diet. Indeed, 60–65% of the progeny of both crosses (DaGS::Uro-RNAi x w1118 and DaGS::Uro-RNAi x b35785) form concretions after being fed the RU486 supplemented HY diet for 14 days (Fig 4C). Using a GAL4-RNAi line targeting expression of the GAL4 transcription factor itself as a positive control reduced concretion formation of DaGS::Uro-RNAi x GAL4-RNAi flies due to the suppression of urate oxidase depletion (Fig 4C). Hence, the recombinant DaGS::Uro-RNAi line represents a useful tool to study UA associated pathologies.

In agreement with the allopurinol and methotrexate treatments, knocking down the corresponding target genes, XDH and DHFR, by RNAi also reduced concretion formation in the recombinant DaGS::Uro-RNAi flies (Fig 4C). Those data were complemented by knockdown of another crucial gene involved in purine de novo synthesis, 5'-phosphoribosyl-1'-pyrophosphosphate synthetase (PRPS), whose knockdown by PRPS-RNAi reduced concretion formation of the DaGS::Uro-RNAi flies as efficiently as the DHFR-RNAi (Fig 4C, S3A Fig). Additionally, allopurinol and methotrexate were able to revert the short lifespan of DaGS::Uro-RNAi #1 flies when added to the RU486 supplemented HY diet (Fig 4D and 4E). Unlike allopurinol, methotrexate supplementation also triggered a lifespan extension of the control DaGS::Uro-RNAi #1 flies consuming the HY diet without RU486 (Fig 4E). Thus, the beneficial effect of methotrexate on lifespan is likely independent of the Uro expression level, yet, accentuates the
Diet-dependent uric acid pathologies in *Drosophila*

![Graph A](image1.png)

**A**

Flies with concretion (%)

| Allopurinol [mM] | 0 | 1 | 5 | 10 |
|------------------|---|---|---|----|
|                  | 0 | 75| 50| 25 |
| Error Bar        |   | **| ***|    |

![Graph B](image2.png)

**B**

Flies with concretion (%)

| Methotrexate [μM] | 0 | 0.2 | 1 | 5 | 50 |
|-------------------|---|-----|---|---|----|
|                   | 0 | 75  | 50| 25| 25 |
| Error Bar         |   | **| ***|    |    |

![Graph C](image3.png)

**C**

| Strain                        | Flies with concretion (%) |
|-------------------------------|---------------------------|
| w1118                         | 0                         |
| b35785                        | 75                        |
| GAL4-RNAi                     | 50                        |
| PRPS-RNAi                     | 25                        |
| DHFR-RNAi                     | 0                         |
| XDH-RNAi                      | 0                         |

**D**

Survival (%)

HY + AP

HY

RU

![Graph D](image4.png)

**E**

Survival (%)

HY + MTX

HY

RU

![Graph E](image5.png)
importance of the purine metabolism in aging (Fig 4E). Overall, over-expression of purine homeostasis rescued the phenotypes of Uro knockdown flies, thereby supporting the validity of the fly model for UA based pathologies and suggesting an important role for purine metabolism in UA homeostasis.

The transcription factor FoxO dampens concretion formation by reducing UA levels and ROS formation

Common genetic variations identified to date explain only 6–7% of the variance encountered in serum UA levels [44–47]. Furthermore, commonly prescribed medications such as allopurinol can result in serious adverse drug reactions or provide insufficient efficacy in a high percentage of users [48–50]. Thus, other genes remain to be identified that may represent better targets to lower the metabolic UA load. To identify previously unrecognized regulators of UA metabolism, we started by examining the ILS pathway for two reasons. Firstly, ILS is a conserved signaling cascade activated when flies are fed a HY diet [25, 51–54]. Secondly, a recent genome-wide association study identified polymorphisms in the human ILS gene IGF1R that were associated with serum UA concentration [47]. Thus, we examined components of the ILS pathway as putative targets controlling UA production and utilization. We used the recombinant DaGS::Uro-RNAi line to address the impact of perturbing ILS signaling. Both strategies the RNAi mediated knockdown of the insulin-like receptor (InR) gene or expression of a dominant-negative InR gene variant (InR-DN) markedly reduced Drosophila concretion formation when fed the HY diet (Fig 5A). Similarly, over-expression of the phosphatase and tensin homolog (PTEN), which acts as negative regulator of ILS signaling downstream of the InR, significantly reduced concretion formation compared to the DaGS::Uro-RNAi x w1118 flies (Fig 5A). Downstream of PTEN the kinase AKT integrates incoming ILS cues into metabolism by phosphorylating different effector molecules. Like inhibition of the InR, blocking ILS signaling by RNAi mediated depletion of AKT in Uro knockdown flies also reduced concretion formation (Fig 5A). One of the effector molecules inactivated by AKT phosphorylation is the evolutionarily conserved transcription factor FoxO [55]. Transgenic over-expression of FOXO in the background of the DaGS::Uro-RNAi flies showed a significant rescue of concretion formation with two different FOXO expression strains (Fig 5B). In contrast, knockdown of FOXO by two different RNAi constructs did not affect concretion formation (Fig 5B). In addition to over-expression and knockdown of FOXO, we examined the direct influence of UA levels (endogenous) FoxO activity using the Uro knockdown flies. Based on the study by Alic et al., we picked seven target genes that are up- or down-regulated by binding of FoxO and are part of either purine metabolism (Ade2, Ade5, Aprt, Veil) or the ILS pathway (AKT, InR, Thor) [56]. No significant change in the expression level of the seven genes was observed when comparing control and Uro knockdown flies (S4A Fig). Thus, activity of FoxO was not altered in response to an increased UA level. However, as FOXO manipulation has a strong effect on UA concretion, we propose it is upstream of UA formation pathways. To further validate the physiological relevance of FOXO over-expression we compared the metabolite levels of relevant
purine intermediates (including UA) in DaGS::Uro-RNAi x w1118 and DaGS::Uro-RNAi x FOXO flies. While early intermediates of purine synthesis such as IMP were not altered, purine degradation products were reduced in flies with the FOXO over-expression (Fig 5C). In particular, UA levels were significantly reduced threefold, thus explaining the reduced deposit formation in flies over-expressing FOXO.

To determine whether the ILS pathway influences UA levels in a conserved manner between flies and humans, we assessed whether single nucleotide polymorphisms (SNPs) in different genes of the ILS pathway are associated with either serum UA (SUA) levels or gout in humans. Using data from 80,795 adult subjects from the Kaiser Permanente Research Program on Genes, Environment, and Health (S1 Table) [50], we found that SNPs in the AKT2 and FOXO3 genes were associated with either SUA levels or the incidence for gout (Table 1, S2 Table). This association was significant after adjusting for multiple testing of phenotypes and SNPs within each gene. In light of our fly studies identifying AKT inhibition (Fig 5A) and FOXO over-expression (Fig 5B) as suppressors of UA concretions the human SNP associations shown in Table 1 support the idea that modulation of the ILS signaling network could play a critical and eventually conserved role in affecting UA levels and associated pathology in both flies and humans.

Next, we addressed the underlying mechanism of the link between the ILS pathway and UA levels. UA has been proposed to play several physiological roles, depending on where it is

### Table 1. A human candidate-gene study (CGS) identifies SNPs in FOXO3 and AKT2 being associated with UA pathologies.

| Human gene | Phenotype | Gene threshold |
|------------|-----------|----------------|
|            | Gout      | SUA            |               |
| FOXO1      | 0.00611   | 0.04906        | 0.00058       |
| FOXO3      | 0.01596   | 0.00004        | 0.00061       |
| AKT1       | 0.07641   | 0.08287        | 0.00066       |
| AKT2       | 0.00003   | 0.01738        | 0.00066       |
| AKT3       | 0.00804   | 0.01533        | 0.00033       |
| PTEN       | 0.01080   | 0.00979        | 0.00064       |

P-values of single nucleotide polymorphisms (SNPs) within relevant genes of the ILS network that are significantly associated with gout or serum uric acid (SUA) levels in human subjects are indicated in bold. Significance of calculated p-values is set by the gene-specific threshold (last column). Supporting information such as description of the study population and effect sizes for significant SNPs is given in S1 Table and S2 Table as well as the material and methods section.

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acting. For instance, UA can act as an antioxidant in the blood, where it accounts for up to 60% of the antioxidative capacity [57]. In turn, in an intracellular setting such as in adipocytes, UA is considered a prooxidant and proinflammatory molecule [12, 58, 59]. Similarly, in Drosophila UA is considered a damage-associated molecular pattern (DAMP) able to trigger a so-called sterile inflammation, which typically results in the production of antimicrobial peptides [60]. Compared to their isogenic control siblings Uro knockdown flies depicted a two- to eightfold increase in the expression of the three antimicrobial peptides attacin A (Att A), diptericin (Dipt), and metchnikowin (Metch) after 14 days on the HY diet (S4B Fig). Increased expression of the antimicrobial peptides was seen for both crosses DaGS>Uro-RNAi #1 / #2 on the HY plus RU486 diet. Using the recombinant DaGS::Uro-RNAi line over-expression of FOXO diminished the high levels of antimicrobial peptides (S4C Fig). However, knockdown of the known NFκB paralogs Relish, Dorsal or the Dorsal-related immunity factor (Dif), whose gene products act as dimeric transcription factors activating expression of antimicrobial peptides, did not alleviate UA concretion formation (S4D Fig) [61, 62]. Work by Zhao et al. indicates that over-expression of individual antimicrobial peptides like diptericin increases the tolerance of flies resisting oxidative stress [63]. Thus, the high expression of antimicrobial peptides (S4B Fig) could represent a defense mechanism of Uro knockdown flies against oxidative stress rather than being a driver of UA accumulation. Consistent with this idea of intracellular UA generating a prooxidative milieu, Uro knockdown flies showed significantly elevated ROS levels in the cells of the hindgut as revealed by the doubling of the ROS sensitive DHE staining intensity (Fig 5D). Considering the well-documented role of FoxO proteins in cellular responses to combat oxidative stress [55, 64, 65], we speculated if FOXO over-expression could reduce the build-up of ROS found in Uro knockdown flies. Comparing ROS levels of DaGS::Uro-RNAi x w1118 flies to DaGS::Uro-RNAi x FOXO flies showed that an increased FoxO abundance indeed caused a reduction of ROS in the hindgut as well as Malpighian tubules (Fig 5E and 5F).

**Lifespan shortening and UA concretion formation of Uro knockdown flies are mediated by NOX**

The increase of ROS levels upon Uro knockdown (Fig 5D–5F) could stem from two sources: (1) inefficient expression of ROS combating genes, or (2) an overabundance of ROS-producing enzymes. From the eleven oxidative stress-related genes whose expression was examined in the Uro knockdown flies, only the ROS-producing NADPH Oxidase (NOX) showed a two-fold higher mRNA content (Fig 6A, S4E Fig). Elevated NOX expression was reversed completely by either FOXO over-expression or by introducing a NOX-RNAi element into DaGS::Uro-RNAi flies (Fig 6A, NOX mRNA). Of note, neither the FOXO over-expression nor the presence of the NOX-RNAi element altered efficiency of the Uro knockdown (Fig 6A, Uro mRNA). Next, we asked if reduction of NOX expression via RNAi could alleviate the concretion formation characteristically observed in Uro knockdown flies. As with FOXO over-expression, knocking down NOX expression in Uro knockdown flies significantly reduced UA accumulation, which was demonstrated with two different NOX-RNAi lines (Fig 6B). Further comparison of purine metabolite levels between Uro knockdown flies (DaGS::Uro-RNAi x w1118) and the Uro plus NOX double knockdown flies (DaGS::Uro-RNAi x NOX-RNAi) identified a significant drop of UA levels and depicted the interwoven nature of the NADPH oxidase with purine catabolism (Fig 6C). We than speculated whether the effect of NOX depletion extended to an improvement in lifespan. Like concretion formation, the recombinant DaGS::Uro-RNAi line also recapitulated the lifespan attenuation on a HY diet when crossed to w1118 (Fig 6D, S5A Fig) or the mCherry-RNAi strain b35785 (S5B Fig). The median lifespan of
Fig 6. NOX mediated ROS production triggers lifespan attenuation and UA concretion formation on a high yeast diet. (A) Levels of NOX mRNA upon modulation of FOXO. Recombinant DaGS::Uro-RNAi #1 flies were crossed to w1118 (w; no additional UAS-locus), or active UAS-transgenes triggering either over-expression of FOXO or inhibition of NOX (NOX-RNAi) and fed the HY + RU or HY - RU diet before relative mRNA levels of urate oxidase (Uro) and NADPH oxidase (NOX) were determined by qRT-PCR. (B) Concretion formation of recombinant DaGS::Uro-RNAi #1 flies crossed to w1118 or one of the two NOX-RNAi lines after 14 days of feeding the HY + RU diet. (C) Relative purine metabolite levels of fly homogenates from recombinant DaGS::Uro-RNAi #1 flies crossed to w1118 or NOX-RNAi #1. Metabolite levels were compared after 14 days of feeding flies the HY + RU or HY - RU diet. (D) Survival curves of flies from (C). Error bars represent the SE of multiple biological repeats. Supporting data is shown in S4 Fig and S5 Fig.

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DaGS::Uro-RNAi x w1118 flies was reduced by 35% from 31 days without RU486 to 20 days in presence of RU486 (Fig 6D). This lifespan attenuation on the HY diet was partially rescued in DaGS::Uro-RNAi x NOX-RNAi flies using NOX-RNAi #1 or #2 (Fig 6D, S5F Fig).

To corroborate the NOX related findings, we used a pharmacological approach by inhibiting NOX activity and the associated ROS production using either apocynin (ACY; a NOX inhibitor) or vitamin C (Vit C; a ROS scavenger), respectively. Dietary ACY supplementation reduced the concretion formation of DaGS>Uro-RNAi flies on the HY plus RU486 diet in a dose-dependent manner. Concretion formation was reduced for both Uro-RNAi fly populations from about 70% without ACY supplementation down to 15% in presence of 5 mM ACY (Fig 7A). In terms of lifespan, a 5 mM ACY supplementation to the HY diet was able to extend the short lifespan of both Uro knockdown fly lines whereas the corresponding control siblings (DaGS>Uro-RNAi #1 / #2 without RU486) showed no change of survival in presence of ACY (Fig 7B and 7C). Thus, the ACY treatment very much mirrored the results obtained with the NOX-RNAi (cf. Figs 6B, 6D and 7A–7C). Last, we speculated if a water-soluble, i.e. membrane-permeable, ROS scavenger like Vit C could reduce the elevated ROS level found in Uro knockdown flies. Supplementation of the HY diet with increasing amounts of Vit C ranging from 10 to 100 mM gradually reduced concretion formation of Uro knockdown flies (Fig 7D). 50 mM Vit C addition to the HY diet also rescued the lifespan effect observed in flies with reduced urate oxidase levels (Fig 7E). Like the MTX treatment (Fig 4E), dietary Vit C supplementation also extended the median survival of control flies (Fig 7E). Worth mentioning, foods rich in Vit C are associated with a reduced risk for elevated UA and gout, due to its uricosuric effect [66, 67]. While Vit C seemed to extend the median lifespan of both the control (DaGS>Uro-RNAi #1 without RU486) and Uro knockdown flies (DaGS>Uro-RNAi #1 with RU486) by 16% and 53%, respectively, the impact of apocynin on lifespan was specific for the urate oxidase depleted animals (Fig 7B, 7C and 7E). In presence of RU486, ACY extended the median lifespan of DaGS>Uro-RNAi #1 flies by 47% (Fig 7B) and that of DaGS>Uro-RNAi #2 flies by 30% (Fig 7C).

In summary, our study argues the importance of purines as a dietary component that limits lifespan. Generally, mutants in genes that influence lifespan upon dietary restriction either extend lifespan upon rich nutrient conditions while failing to extend lifespan upon dietary restriction conditions or attenuate the maximum lifespan upon dietary restriction. However, Uro mutants belong to a “novel” class of genes that limit lifespan only upon rich conditions but have little or no influence upon dietary restriction. We hypothesize that mutants that display such a phenotype encode a gene that amplifies the cellular damage that takes place under rich nutrient conditions compared to dietary restriction. We speculate that the increase of UA is deleterious, and a lower UA level is partially responsible for the lifespan extension upon dietary restriction. The fly model established here will also be useful in dissecting the pathological effects of elevated UA on crystal formation as well as the underlying genetic factors driving crystal formation in the first place. The latter point is of particular interest, considering that only 5% of hyperuricemic people with SUA above 9 mg/dl develop gout [8, 9]. The genetic predisposition influencing UA accumulation is currently subject of genome-wide association studies and is complemented by our approach combining in vivo fly work and human candidate gene studies. Our data showing that genes such as AKT, FOXO and NOX are associated with UA levels and the incidence for deposit formation start shedding light on this important issue. In addition, the fly model will be beneficial to study the larger role of UA and UA deposits on influencing mortality as was underpinned by recent studies linking exceptional longevity and a reduced prevalence of age-related diseases with UA levels at the lower end of the human SUA spectrum [21, 22].
Figure A shows the effect of different concentrations of ACY on the formation of uric acid concretions in Drosophila. Figure B illustrates the survival rates under different conditions of HY with and without RU. Figure C presents similar data for DaGS>Uro-RNAi #2. Figure D and E show the influence of Vitamin C on uric acid concretions and survival rates, respectively. Figure F depicts the molecular mechanisms underlying diet-dependent uric acid pathologies, involving InR, AKT, FoxO, NOX, ROS, and purines.
Our work demonstrates that inhibition of Uro in flies can serve as a useful model to study UA induced pathologies related to lifespan and crystal formation. Our results also highlight the importance of ILS signaling, and its downstream effector NOX, as an important mediator of UA metabolism and related pathologies (Fig 7F). Future research using this model can help determine the role of the several candidate genes in humans that have been implicated in altering UA levels. Given the over-nutrition encountered in developed countries, the causal nexus uncovered here identifies potential new drug targets that could help regulate UA levels, ameliorating pathologies associated with hyperuricemia and extending human healthspan.

Material and methods

Contact for reagent and resource sharing

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Pankaj Kapahi (pkapahi@buckinstitute.org).

Experimental model and subject details

Fly strains used in this study. Fly strains purchased from Bloomington Drosophila Stock Center are indicated by the prefix “b”, strains from FlyORF, Zurich by “F”, strains from NIG-Fly, Mishima by “n”, and strains from VDRC, Vienna by “v”.

GAL4 driver lines: daughterless gene switch (kindly provided by Linda Partridge), daughterless (b55851), c42 (b30835), Uro (b44416), Elav (b458)

UAS responder lines: Uro-RNAi #1 (n7171-R1), Uro-RNAi #2 (v330484), InR-RNAi (b31594), InR-DN (b8252), PTEN (kindly provided by Tiang Xu), AKT-RNAi (b31701), FOXO #1 (b9575), FOXO #2 (F000143), Rel-RNAi (b28943), Di-RNAi (b32934), NOX-RNAi #1 (b32902), NOX-RNAi #2 (b32433), GAL4-RNAi (b35784), PRPS-RNAi (b35619), DHR-RNAi (b35015), XDH-RNAi (v25175)

Other lines: w1118, b35785 (expresses a dsRNA for RNAi of mCherry under UAS control). We generated a recombinant fly line carrying both genetic elements (+/+; DaGS::Uro-RNAi #1/ DaGS::Uro-RNAi #1; +/+ ) on chromosome two. This line was crossed to UAS responder lines to introduce an additional transgenic element (including the no target UAS-mCherry-RNAi line b35785) or w1118 (Fig 4C, Fig 5A, 5C, 5E and 5F, Fig 6 and S4C, S4D and S5 Figs).

With the exception of Uro-RNAi #2 (v330484) all other fly strains were routinely outcrossed into the commonly used w1118 background to achieve genetic homogeneity amongst different strains and provide isogenization.

Fly husbandry, dietary and pharmacological manipulations. All fly lines were maintained on standard fly yeast extract medium containing 1.5% yeast, 5% sucrose, 0.46% agar, 8.5% of corn meal, and 1% acid mix (a 1:1 mix of 10% propionic acid and 83.6% orthophosphoric acid) prepared in distilled water. To prepare the media, cornmeal (85 g), sucrose (50 g), active dry yeast (16 g, "Saf-instant") and agar (4.6 g) were mixed in a liter of water and brought to boil under constant stirring. Once cooled down to 60°C 10 ml of acid mix was added to the media. The media were then poured in vials (~10 ml/vial) or bottles (50 ml/bottle) and

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allowed to cool down before storing at 4°C for later usage. These vials or bottles were then seeded with some live yeast just before the flies are transferred and used for maintenance of lab stocks, collection of virgins, or setting up crosses.

Experimental low and high yeast diets varied only with regard to the yeast type and content ranging from 0% to 8%. Diets declared as LY (low yeast content) and HY (high yeast content) contained 0.5% and 5% Baker’s yeast extract (212750 Bacto Yeast Extract, B.D. Diagnostic Systems, Sparks, MD), respectively. If not stated otherwise pharmacological treatments or dietary supplementations were mixed with the other ingredients during preparation of the food. As for the standard medium, the mix was brought to a boil under constant stirring and allowed to cool down to 60°C before adding the acid mix. For the induction of transgenic elements via the GAL4-UAS system when using a gene switch driver 200 μM RU486 (Mifepristone, Sigma) in 100% ethanol was added to the food while the isogenic controls received vehicle treatment. The media was then poured in vials (~5 ml/vial) and allowed to cool down before storing at 4°C for later usage. Further dietary supplementations and drug treatments used in our study are summarized in Table 2.

Fly rearing. Genetic crosses were obtained by pairing 15 young virgin females (carrying the GAL4 driver construct) with 4 young male flies (carrying the UAS responder construct) in new stock bottles. Flies were kept in the stock bottles for five days, after which the parents were removed, and the larvae could develop in a temperature (25°C) and humidity (60%) controlled designated fly room with a 12-hour light / dark cycle. Newly eclosed flies could mate for 2–3 days, to complete development post-eclosion, before they were grouped under light CO₂ anesthesia. Sorted females were then transferred to the appropriate media vials for subsequent analyses.

Method details

Lifespan analysis. Flies developed on standard fly 1.5% yeast extract medium were transferred to the necessary diet within 72 hours after eclosion. For survivorship analysis four to six media containing vials with 25 mated females were kept in a temperature (25°C) and humidity (60%) controlled designated fly room with a 12-hour light / dark cycle. Newly eclosed flies could mate for 2–3 days, to complete development post-eclosion, before they were grouped under light CO₂ anesthesia. The significance of change was determined using the log-rank test. Each lifespan was repeated at least once to generate independent, biological replicates.

| Dietary Additive | Ingredients | Vendor | Concentrations |
|------------------|-------------|--------|----------------|
| 10 mM Pu (Purines) | Adenine, Guanine | Sigma-Aldrich | 5 mM each |
| 20 mM Pu | Adenine, Guanine | Sigma-Aldrich | 10 mM each |
| 40 mM Pu | Adenine, Guanine | Sigma-Aldrich | 20 mM each |
| 20 mM Py (Pyrimidines) | Cytosine, Thymine | Sigma-Aldrich | 10 mM each |
| AP | Allopurinol | Sigma-Aldrich | 1, 5, 10 mM |
| MTX | Methotrexate | Sigma-Aldrich | 0.2, 1, 5, 50 μM |
| Vit C | Vitamin C | Sigma-Aldrich | 10, 50, 100 mM |
| ACY | Apocynin | Sigma-Aldrich | 1, 5 mM |
| 5% Protein | BSA, Casein | Cell Signaling Technology, Sigma-Aldrich | 2.5% (w/v) each |
| 15% Protein | BSA, Casein | s. above | 7.5% (w/v) each |
| 10% Sucrose | Sucrose | Mallinckrodt Chemicals | 10% (w/v) |
| 20% Sucrose | Sucrose | s. above | 20% (w/v) |

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allowed to cool down before storing at 4°C for later usage. These vials or bottles were then seeded with some live yeast just before the flies are transferred and used for maintenance of lab stocks, collection of virgins, or setting up crosses.

Experimental low and high yeast diets varied only with regard to the yeast type and content ranging from 0% to 8%. Diets declared as LY (low yeast content) and HY (high yeast content) contained 0.5% and 5% Baker’s yeast extract (212750 Bacto Yeast Extract, B.D. Diagnostic Systems, Sparks, MD), respectively. If not stated otherwise pharmacological treatments or dietary supplementations were mixed with the other ingredients during preparation of the food. As for the standard medium, the mix was brought to a boil under constant stirring and allowed to cool down to 60°C before adding the acid mix. For the induction of transgenic elements via the GAL4-UAS system when using a gene switch driver 200 μM RU486 (Mifepristone, Sigma) in 100% ethanol was added to the food while the isogenic controls received vehicle treatment. The media was then poured in vials (~5 ml/vial) and allowed to cool down before storing at 4°C for later usage. Further dietary supplementations and drug treatments used in our study are summarized in Table 2.

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Table 2. Dietary supplementations and drug treatments.

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| 40 mM Pu | Adenine, Guanine | Sigma-Aldrich | 20 mM each |
| 20 mM Py (Pyrimidines) | Cytosine, Thymine | Sigma-Aldrich | 10 mM each |
| AP | Allopurinol | Sigma-Aldrich | 1, 5, 10 mM |
| MTX | Methotrexate | Sigma-Aldrich | 0.2, 1, 5, 50 μM |
| Vit C | Vitamin C | Sigma-Aldrich | 10, 50, 100 mM |
| ACY | Apocynin | Sigma-Aldrich | 1, 5 mM |
| 5% Protein | BSA, Casein | Cell Signaling Technology, Sigma-Aldrich | 2.5% (w/v) each |
| 15% Protein | BSA, Casein | s. above | 7.5% (w/v) each |
| 10% Sucrose | Sucrose | Mallinckrodt Chemicals | 10% (w/v) |
| 20% Sucrose | Sucrose | s. above | 20% (w/v) |

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allowed to cool down before storing at 4°C for later usage. These vials or bottles were then seeded with some live yeast just before the flies are transferred and used for maintenance of lab stocks, collection of virgins, or setting up crosses.

Experimental low and high yeast diets varied only with regard to the yeast type and content ranging from 0% to 8%. Diets declared as LY (low yeast content) and HY (high yeast content) contained 0.5% and 5% Baker’s yeast extract (212750 Bacto Yeast Extract, B.D. Diagnostic Systems, Sparks, MD), respectively. If not stated otherwise pharmacological treatments or dietary supplementations were mixed with the other ingredients during preparation of the food. As for the standard medium, the mix was brought to a boil under constant stirring and allowed to cool down to 60°C before adding the acid mix. For the induction of transgenic elements via the GAL4-UAS system when using a gene switch driver 200 μM RU486 (Mifepristone, Sigma) in 100% ethanol was added to the food while the isogenic controls received vehicle treatment. The media was then poured in vials (~5 ml/vial) and allowed to cool down before storing at 4°C for later usage. Further dietary supplementations and drug treatments used in our study are summarized in Table 2.

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Dissection and scoring of concretion formation. For dissection and imaging assays adult female flies were dissected after 4, 7, 11, or 14 days. The excretory system, in particular the hindgut and Malpighian tubule area was checked and imaged for the presence of ectopic biomineralization. Flies were anesthetized by CO$_2$ on standard flypads (Cat # 59–108, Genesee Scientific, San Diego, CA) and dissected under a dissecting light microscope (SZ61, Olympus, Center Valley, PA) on a culture dish in droplets of distilled water utilizing fine forceps (Roboz ceramic, Gaithersburg, MD). The excretory system was imaged utilizing a Zeiss Stereo Microscope with external light source. While transparent posterior midguts and hindguts devoid of any sign of deposit formation were scored as negative, those with pale yellow/orange hard concretions were scored as concretion carrying specimens. The relative level of concretion formation was quantified by this binary approach. Per condition at least 15–20 flies were dissected and the number of flies with deposits was divided by the total number of flies dissected. Each experiment was repeated for a minimum of three independent, biological replicates.

Imaging of ectopic biomineral. Images of dissected guts and Malpighian tubules were captured with the Olympus SZX12 microscope. Isolated deposits extracted from the hindgut area were imaged with the Olympus BX51 microscope (Olympus Scientific Solutions Americas Corp. Waltham, MA).

For the micro-computer tomography (μCT) imaging fly specimens were scanned in 50% ethanol using a micro-XCT unit at 4X magnification (4.5 μm/voxel) with 1200 image projections. Scanning was performed with a 3 second exposure, source power of 40 W, 200 μA current, source distance of ~30 mm, detector distance of ~14 mm, camera binning 2, and an angle sweep from -93˚ to 93˚. Post-3D reconstruction, the acquired data was further processed and analyzed with Avizo software (Version 9.3.0; FEI, Hillsboro, Oregon).

Collection of deposits for mass spectrometry-based metabolomics. Flies were dissected in 50% methanol/water mix and the deposits from the hindgut area were isolated using fine forceps. Deposits of 25 flies carrying concretions were collected in an Eppendorf tube filled with 500 μl 50% methanol/water. The material was sedimented by centrifugation for 15 s with 5,000 rpm and the supernatant was aspirated. After two further wash steps with 500 μl 50% methanol/water deposits were frozen in liquid nitrogen and stored at -80˚C until further processing.

High-performance liquid chromatography (HPLC) mass spectrometry (MS). High-performance liquid chromatography (HPLC) was performed using an Agilent 1260 UHPLC system and connected to a Phenomenex Luna NH2 column (2 x 100 mm, 3 μm, 100 Å) and a SecurityGuard NH2 guard column 4 x 2 mm ID. Mass spectrometry (MS) was performed using a 5500 Triple-Quadrupole LC-MS/MS mass spectrometer from Sciex fitted with a Turbo VTM ion source. Sciex’s Analyst v1.6.1 [68] was used for all forms of data acquisition, development of HPLC method, and optimization of analyte-specific MRM (multiple reaction monitoring) transitions. Skyline v4.1 was used for LC-MS/MS data analysis. For whole fly analysis, 5 flies (in sextuplicates) per condition were flash-frozen over liquid nitrogen and subsequently homogenized ultrasonically using a Fisher Scientific’s 550 Sonic Dismembrator with 50 μl of an 8:2 mixture of methanol/water (v/v), containing 2.5 μM of 2-chloroadenosine as internal standard. Three 20 s pulses at amplitude setting 3 of the instrument (on ice) were sufficient to completely homogenize fly specimens. The homogenates were then vortexed for 5 times over a period of ~30 min (each 1 min long). Subsequently, the samples were centrifuged at 10,000 rpm for 10 min, the supernatant was filtered, and 3 μl of each was injected for HPLC-MRM analysis (vide infra) without any additional processing. For the analysis of fly deposits, samples were collected as described above and prepared for HPLC-MRM analysis (vide infra) as described above for whole flies.
Optimization of analyte-specific MRM transitions, such as determination of suitable precursor and product ions and optimal MS parameters for each transition (Q1, precursor → Q3, product) were achieved by isocratic flow injection of the 1–10 μM solution (final) for each standard, diluted in 80% methanol. The most intense transition was used as quantifier, whereas one or more additional transitions were used as qualifier for each compound (S3 Table). A final standard mixture of all compounds at 5 μM (containing the internal standard 2-chloroadenosine at 2.5 μM), were prepared prior to analysis and injected at the onset of each biological sample set.

Based on previous reports [69], the following HPLC program was developed: a solvent gradient of 20 mM ammonium acetate + 20 mM ammonium hydroxide (pH ~ 9.5) + 5% acetonitrile (aqueous)–acetonitrile (organic) was used with 0.4 ml/min flow rate, starting with an acetonitrile content of 85% for 1 min, which was decreased to 30% over 3 min and then to 0% over 7 min and held at 0% for 2 min. The HPLC column was subsequently reconstituted to its initial condition (acetonitrile content of 85%) over the next 1 min and re-equilibrated for 7 min. Metabolome extracts from whole flies or deposits were analyzed by HPLC-MRM with positive/negative switching of source ion modes [69]. Source conditions were as follows: curtain gas (CUR) 20, collision gas (CAD) 7, ion source gas 1 (GS1) 30, ion source gas 2 (GS2) 30, ionspray voltage (IS) ±4500 V, and source temperature (TEM) 450˚C. Quantification was based on integration of analyte-specific peaks obtained from HPLC-MRM runs.

**Feeding assay.** After 14 days of feeding a particular diet, flies were switched to blue dye medium (15% sucrose, 1% agar, 1% FD&C Blue #1) for 2 hours. Flies were then frozen in liquid nitrogen. To prevent eye pigment from interfering with absorbance spectrum, heads were separated from the bodies and bodies were then homogenized in PBS buffer and centrifuged at 13,000 rpm for 25 minutes. Supernatants were transferred to a 96-well plate and analyzed for absorbance at 625 nm with use of SpectraMax M2 spectrophotometer [70].

**Dihydroethidium (DHE) staining for ROS measurements.** Intact guts with adhering Malpighian tubules were dissected from female flies in a droplet of PBS (137 mM NaCl, 2.7 mM KCl, 10 mM Phosphate, pH 7.4) on ice and subsequently stained for 5 min with 45 μM DHE (Thermo Fisher Scientific) dissolved in DMSO (Sigma Aldrich). After three 10 min wash steps with PBS the sample was fixed for 45 min using 4% formaldehyde (Sigma Aldrich) in PBS followed by another three 10 min wash steps. After a 15 min staining with 1 μg/ml DAPI (Thermo Fisher Scientific) dissolved in PBS the specimen was mounted onto a microscope slide using Mowiol (Sigma Aldrich). Confocal images were collected using a Zeiss LSM700 confocal system using an Excitation/Emission (nm) of 518/605. DHE intensity was quantified using Image J.

**Total RNA and cDNA preparation.** Total RNA was isolated from 5–10 females using the Quick-RNA MiniPrep Kit (Zymo Research) at room temperature. In brief, flies were anesthetized with CO₂ before homogenization using a Kontes Microtube Pestle Pestle Rods with Motor in an Eppendorf tube containing 200 μl RNA Lysis Buffer. After adding another 400 μl RNA Lysis Buffer the homogenate was centrifuged for 1 min with 12,000 rcf and 400 μl supernatant was transferred to the spin-away filter placed over a fresh collection tube for elimination of genomic DNA. 400 μl of 95% ethanol was added to the flow through and mixed thoroughly. Transfer mixture to Zymo-Spin IIIICG column in a collection tube and centrifuge for 1 min with 12,000 rcf. After adding 400 μl RNA Prep Buffer and a 30 sec centrifugation with 12,000 rcf two more wash steps with 700 μl and 400 μl RNA Wash Buffer follow. To ensure complete removal of the RNA Wash Buffer the last centrifugation is carried for 2 min with 12,000 rcf. The Zymo-Spin IIIICG column is placed in a collection tube and bound RNA is eluted with 30 μl DNAse/RNAsafe free water in a collection tube by centrifugation for 30 sec with 12,000
rcf. Quantity and quality of isolated RNA was determined using the NanoDrop 1000 Spectrophotometer (Thermo Scientific).

1 µg of total RNA in a volume of 16 µl was used per sample and cDNA was synthesized using 4 µl iScript Reverse Transcription Supermix for RT-qPCR (Bio-Rad) according to the manufacturer’s protocol. The RT-PCR reaction protocol included a priming step (5 min at 25˚C) followed by the reverse transcription (30 min at 42˚C) and inactivation of the reaction (5 min at 85˚C). If necessary, cDNA was stored at -20˚C before the quantitative real-time PCR.

**Quantitative real-time PCR (qRT-PCR).** Using 3 µl of the 1:10 diluted cDNA as a template, 2 µl of a primer pair (500 nM each) and 5 µl SensiFAST SYBR No-ROX Kit (BIOLINE) the qPCR was performed using the Light Cycler 480 Real-Time PCR System (Roche Applied Science). After pre-incubating the sample (95˚C for 2 min) to denature DNA forty PCR cycles of denaturing (95˚C for 5 s, ramp rate 4.8˚C/s), as well as annealing and extension (60˚C for 20 s, ramp rate 2.5˚C/s) followed. The specificity of amplicons was verified with a subsequent melting curve analysis. The data was analyzed by means of the ΔΔCt method and the values were normalized using β-tubulin as an internal control. The primer pairs used in this study are summarized in Table 3.

**Human candidate-gene study (CGS).** Data from the Kaiser Permanente Epidemiology Research on Adult Health and Aging (GERA) Cohort, a subset of the Research Program on Genes, Environment and Health (RPGEH), were gathered for analysis. Detailed methods on the genotyping, imputation, and ancestry calculations can be found in [50]. The study population consisted of 80,795 adult subjects of European ancestry. The average age of the cohort was

| Gene name | Forward Primer Sequence | Reverse Primer Sequence |
|-----------|-------------------------|-------------------------|
| CG9277 (β-Tubulin) | acatccgcccgcgtgccgc | aagaacctcgctgcctgaaca |
| CG7171 (Uro) | gggatggtgtgttaagagaa | tttcgacccggggagac |
| CG3143 (FOXO) | cgagatcgctcctcagag | aagatcgtcgccctactg |
| CG34399 (NOX) | cccttgccccgacatcatctct | tcgagtacacaaatgccgac |
| CG9127 (Adε2) | cgcttctggtctgagcagc | gctgtagactgtgtgtaacc |
| CG3989 (Adε5) | cccacctcagccactccata | gctgtagacgagccaaag |
| CG4006 (AKT) | atgacgcctcgacagagac | cctccgcgcacaaaaaacc |
| CG18402 (InR) | acaccttttacaaaccgca | ccctcgtctctcagtactg |
| CG8846 (Thor) | cccaccttcttttctctc | acagaccgcactgtctg |
| CG18315 (Aprt) | gattcgcgaagagggcaag | ttgctcgcctcaagggctgta |
| CG4827 (Veil) | ggtacctgagcagttagc | gacccaggcgcggtcact |
| CG10146 (Attacin A) | ccaatggtggtggtgacag | ggcacatcgaccgcatt |
| CG12763 (Diptericin) | cacagatttgagctgaatgg | ttcctcgctgctgctgagt |
| CG8175 (Metchnikowin) | ttgctggacgttggatttg | ttccgacgctgcaggtgac |
| CG6871 (Catalase) | tgacacaaacactccaaacg | ttgatcccaaggtgtctc |
| CG3131 (DUOX) | agcgcctgctgctcactacg | cgcgttcctctgctgact |
| CG5164 (Got-E1) | gggctacacatcagccgtgatg | tccatattcctcccgctggt |
| CG12013 (PHGPx) | ttacgctcgcaggtgagaagt | cgtttctgtactctccgtga |
| CG5826 (Prx3) | gaagactacaggcgccagtagcc | ggggcaacagatggg |
| CG11793 (SOD1) | caagggcaggagtttccctc | ctcaccccggagcacctc |
| CG8905 (SOD2) | aatttggctcggctcagac | tgatgcgcctgctcact |
| CG31884 (Trx-2) | acatggtgccagaggtgaagat | caagctgcagaaatcc |
| CG2151 (Trxr-1) | gatgacccagaaagggctgatg | gaaacctcgcagagggcaca |
| CG7642 (XDH) | ttggtgcctccccacctgggaag | ccttggtcgggtatccgagt |
approximately 63 years old at the time of sample collection for genotyping. Two phenotypes were used for this study: (1) Gout, defined as two ICD-9 codes for gout in the electronic health record; (2) mean serum UA levels. Association analyses were conducted using linear or logistic regression models in PLINK v1.90 adjusting for population structure using the top 10 principal components. In the model, concomitant diuretics and gender were included as covariates because of their strong correlation with all phenotypes. Maximum body mass index (BMI) correlated strongly with UA levels, and was therefore also included as a covariate in all analyses.

SNPs within +/- 2kb of ILS genes (FOXO1, FOXO3, AKT1, AKT2, AKT3, PTEN) were analyzed. Lists of SNPs within genes were analyzed using NIH LDLink SNPclip tool [71], where SNPs in high linkage disequilibrium ($R^2 > 0.5$) were pruned. Gene-specific p-value thresholds were set as $0.05/(2$ phenotypes’ number of SNPs in each gene post-pruning). As a quality control step, top associated SNPs were checked for extreme HWE departures in our samples.

R v3.4.0 was used to create phenotypes and analyze associations between risk factors and phenotypes.

**Quantification and statistical analysis**

Bar graphs and scatter blots represent the mean ± standard error (SE), when indicated. Lifespan curves, and other graphs were visualized using GraphPad Prism 5 software. For statistical comparisons between the control and single treatment group a two-tailed student’s t-test was used; except for lifespan analysis where the log-rank test was performed. When multiple conditions were compared ANOVA was used followed by Tukey’s multiple comparison post-test comparing all pairs of conditions. Significant differences are indicated according to the p-value by asterisks with $p < 0.001$ (***), $0.01 < p < 0.05$ (**). Non significant differences are indicated by n.s.

**Supporting information**

**S1 Fig.** Uro knockdown attenuates *Drosophila* lifespan specifically on diets with a high yeast extract content. Average median lifespan of DaGS>*Uro*-RNAi #1 flies fed diets with the indicated yeast concentration in absence (- RU) or presence (+ RU) of RU486. The average median lifespan was deduced from individual Kaplan-Meier survival curves and is defined as the time point in days when 50% of the population is alive. Error bars represent the SE of multiple biological repeats. Lifespans for the 2, 3, 4, and 8% yeast concentration were not repeated. (TIF)

**S2 Fig.** Uro knockdown enhances UA concretion formation in the excretory system specifically on a HY diet. (A) Concretion formation of DaGS>*Uro*-RNAi #1 flies after 14 days of feeding a diet with the indicated yeast concentration supplemented without (- RU) or with (+ RU) RU486. (B) Concretion formation of DaGS>*Uro*-RNAi #2 flies reared on a HY - RU or HY + RU diet for 14 days. Error bars represent the SE of multiple biological repeats. (TIF)

**S3 Fig.** Overview of purine homeostasis, inhibitors thereof, and their impact on food consumption. (A) Shown are key enzymes (red), inhibitors (blue) thereof, and metabolites (black) of the tripartite purine metabolism orchestrated by purine *de novo* synthesis, salvage and degradation. (B) Using a colorimetric assay, the food intake of DaGS>*Uro*-RNAi #1 flies was compared after exposure to the indicated HY + RU diet supplemented without (-) or with 10 mM allopurinol (AP) or 50 μM methotrexate (MTX) for 14 days. AU = arbitrary units. (TIF)

**S4 Fig.** FoxO activity and antimicrobial peptide expression in Uro knockdown flies fed a high yeast diet. (A) Relative mRNA levels of transcripts encoding for FoxO target genes
(taken from [56]) were determined by qRT-PCR. Genes in the orange and blue area were previously determined as targets up-regulated and down-regulated by induction of FOXO expression, respectively. Genes of interest belong to purine metabolism (phosphoribosylformylglycinamide synthase (Adε2), PAICS bifunctional enzyme (Adε5), adenine phosphoribosyltransferase (Apfr), Veil or the ILS pathway (AKT, InR, Thor). DaGS>Uro-RNAi #1 flies were fed the high yeast diet with RU486 (HY + RU) or without RU486 (HY - RU) for 14 days before comparing gene expression levels in the Uro knockdown and control flies. (B) Relative mRNA levels of transcripts encoding for the antimicrobial peptides attacin A (Att A), diptericin (Dipt), and metchnikowin (Metch) were determined by qRT-PCR. DaGS>Uro-RNAi #1 or DaGS>Uro-RNAi #2 flies were fed the HY + RU or HY - RU diet for 14 days before comparing expression levels. (C) As in (B) the mRNA levels were determined by qRT-PCR from the recombinant DaGS::Uro-RNAi #1 flies crossed to w1118 (no additional UAS-locus), or strains with active UAS-transgenes triggering either over-expression of FOXO (FOXO #1, FOXO #2) or inhibition of NOX (NOX-RNAi #1, NOX-RNAi #2). Flies were fed the HY + RU or HY - RU diet for 14 days. (D) The recombinant DaGS::Uro-RNAi #1 line was crossed to w1118 (no additional UAS-locus), or active UAS-RNAi lines targeting one of the NFκB paralogs Relish (Rel-RNAi), Dif (Dif-RNAi) or Dorsal (Dl-RNAi). To measure concretion formation the flies were fed the HY + RU diet for 14 days prior to dissection. (E) Relative mRNA levels of transcripts encoding for indicated oxidative stress-related proteins were compared 4 and 14 days after feeding the HY + RU or HY - RU diet to DaGS>Uro-RNAi #1 flies. Error bars represent the SE.

S5 Fig. NOX depletion rescues the short lifespan of Uro knockdown flies. (A-E) Survival curves of recombinant DaGS::Uro-RNAi #1 flies crossed to w1118 (no additional UAS-locus), b35785 (carrying a no target UAS-mCherry-RNAi), or active UAS-RNAi lines targeting NOX (NOX-RNAi #1, NOX-RNAi #2) or the GAL4 transcription factor (GAL4-RNAi). Flies were fed the transgene activating diet HY + RU or control diet HY—RU. (F) Average median lifespan from multiple repeats of the fly strains shown in A-D fed the different HY diets. Error bars represent the SE.

S1 Table. Description of human candidate gene study (CGS) population. Serum uric acid (SUA) levels are not included in a typical doctors’ visit and are therefore missing for most healthy controls. Avg, average; max BMI, maximal body mass index in the evaluation period.

S2 Table. Top SNPs and associated effect sizes for each phenotype. SNPs reaching the gene-specific significance threshold are shown with their effect sizes and specific P-values. SNP, single nucleotide polymorphism; SE, standard error; OR, odds ratio; UA, uric acid.

S3 Table. Summary of the analyte-specific MRM (multiple reaction monitoring) transitions.

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