Organsisms of vastly differing morphologies, ecologies, and behaviors—such as fruit flies and humans—are now known to share a multitude of molecular, cellular, and developmental processes. Not only is there extensive similarity in the sequences of fly and human genes, but in addition, almost all of the proteins and major signal transduction pathways that control cell division and differentiation in mammals are also found in the fruitfly Drosophila melanogaster (Rubin et al. 2000; http://flybase.bio.indiana.edu/). Components in these pathways perform the same biochemical functions and act in the same order in both fruitfly and mammalian cells.

Evolutionary conservation is of considerable practical and theoretical importance to biologists. First, it provides a valuable source of data for the reconstruction of phylogeny (Salemi and Vandamme 2003). Evolutionary connections between organisms that were once hidden by morphology have now been exposed in genomic analyses. Second, the conservation of evolutionary processes or traits is a prime area of investigation in theoretical evolutionary biology (Gould 2002). What can, and cannot, be changed evolutionarily? In a constantly evolving world, how can any biological system or trait survive unchanged (Van Valen 1973)? Finally, conservation provides fundamental insights into how complex biological systems, such as immunity, are assembled, maintained, and altered in evolution.

Elements of Immunity

“Innate” immunity refers to the variety of physical, cellular, and molecular features that provide the first lines of defense against infections. The relatively quick innate immune responses operate along with slower but more targeted adaptive immune responses that generate antigen-specific mechanisms that eventually lead to the destruction and elimination of the pathogen.

In mammals, the skin and the epithelial lining of the mucosal tissues act as the primary nonspecific barriers, impeding infectious agents from entering the body. The mucous membrane barrier traps microorganisms, and the cilia present on the epithelial cells assist in sweeping the microbes towards the external openings of the respiratory and gastrointestinal tracts. If infectious agents gain entry into the body, internal innate immune responses become activated and rapidly eliminate the infection. Internal innate immune agents and responses include (amongst others) low pH of the stomach and vagina, proteolytic enzymes and bile in the small intestine, and phagocytosis.

Phagocytosis is a fundamental innate immune mechanism carried out by a number of different cell types, including macrophages. Specific macrophage subpopulations are associated with different tissues (alveolar macrophages in the lung, microglial cells in the central nervous system, etc.). Their main function is to consume microorganisms, other foreign substances, and old, dying cells.

Innate immunity is present from birth, and the information for innate immune responses is inherited. Cells in the mammalian innate immune system (e.g., macrophages) detect “microbial” by recognizing pathogen-associated molecular patterns (PAMPs; Janeway 1989). PAMPs are products of microbial metabolism that are conserved over evolution, distributed in a wide variety of pathogens, and not found in host cells. Lipopolysaccharide is an example of a PAMP and is found in bacteria, viruses, and fungi. Receptors, called pattern recognition receptors, are present on surfaces of host cells and recognize PAMPs. When activated, pattern recognition receptors induce intracellular signaling via the transcription factor NF-κB, resulting in the activation of genes involved in host defense.

Adaptive immunity is characterized by greater specificity than innate immunity, as the adaptive immune response can not only distinguish foreign cells from self, but can also distinguish one foreign antigen from another. Another hallmark of adaptive immunity is memory, which enables the body to remember specific adaptive responses in response to specific antigens. Immunological memory allows the body to make a greater and more rapid second response when the body is reinfected by the same pathogen. Immunological memory underlies both immunization and resistance to reinfection, conferring a tremendous evolutionary advantage to vertebrates. The adaptive immune response has nearly infinite flexibility: the T and B lymphocytes of the acquired immune system can rearrange the elements of their immunoglobulin and T-cell receptor genes to create billions of clones with distinct antigen receptors. In organisms where both innate and acquired immune systems are present, there is a clear interdependence between the two systems. For a fully functional immune system, these components must act in synergy.

Citation: Govind S, Nehm RH (2004) Innate immunity in fruit flies: A textbook example of genomic recycling. PLoS Biol 2(8): e276.

Copyright: © 2004 Govind and Nehm. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abbreviation: PAMP, pathogen-associated molecular pattern

Shubha Govind and Ross H. Nehm are in the Department of Biology and The Graduate Center, City College of New York, New York, New York, United States of America. E-mail: sgovind@ccny.cuny.edu (SG)

DOI: 10.1371/journal.pbio.0020276
Innate Immunity in Drosophila

Because it lacks an adaptive immune response, Drosophila melanogaster serves as a wonderful model for studying aspects of the innate immune system that might otherwise be obscured by the actions of the adaptive immune response. Insects defend themselves against parasites and pathogens by invoking a multitude of innate immune responses (Figure 1; for more details, see recent reviews by Hoffmann and Reichhart [2002], Hultmark [2003], Brennan and Anderson [2004], Meister [2004], and Theopold et al. [2004]). Like humans, Drosophila protects itself against microbes and parasites via epithelial barriers: for example, epithelial cells of the trachea, gut, genital tract, and Malpighian tubules produce antimicrobial peptides (local response).

Once within the body cavity, microbes may be consumed by the phagocytic blood cells called plasmatocytes (Figure 1). Larger pathogens (such as eggs of parasitic wasps) are inactivated by encapsulation, an immune response carried out by specialized cells called lamellocytes (Figure 1). Lamellocytes differentiate in response to macroscopic pathogens, and their precursors are thought to reside in the larval lymph gland. The transcription factors (GATA, Friend-of-GATA, and Runx family proteins) and signal transduction pathways (Toll/NF-kB, Serrate/Notch, and JAK/STAT) that are required for specification and proliferation of blood cells during normal hematopoiesis, as well as during the hematopoietic proliferation that accompanies immune challenge, are conserved (Evans et al. 2003; Meister 2004). In this issue of PLoS Biology, Crozatier et al. (2004) identify the transcription factor Collier as being critical for the differentiation of lamellocytes in Drosophila. The mammalian ortholog of Collier (Early B-cell Factor) is involved in B-cell differentiation in mice.

In addition to triggering cellular immune responses, invading pathogens also activate humoral reactions. Microbes induce the rapid secretion of antimicrobial peptides from the cells of the fat body into the larval or adult body cavity (systemic response; Figure 1). A microbial infection initiates a zymogen cascade that plays a crucial role in the activation of the antimicrobial genes in the fat body. Infection or wounding also triggers a protein-cleaving cascade that results in the production of toxic intermediates and melanin around microbes or wound sites. This proteolytic cascade is similar to the vertebrate clotting cascade. Drosophila hemolymph also coagulates and participates in host defense and wound healing (Figure 1; Theopold et al. 2004). Given the evolutionary success of insects, this combination of defense mechanisms has proven to be extremely effective, allowing insects to thrive in septic environments.
NF-κB Activation: The Toll and Imd Pathways of Drosophila

The Drosophila genome encodes several members of the multifunctional Toll family of receptors (Beutler and Rehli 2002). Mutations in the Drosophila Toll gene (as well as in other components in the pathway) make the fly susceptible to fungal or gram-positive bacterial infections. However, Toll does not act as a pattern recognition receptor in the fly; instead its activation depends on the presence of the processed (active) form of the growth-factor-like polypeptide Spätzle. Processing of Spätzle depends on a serpin-controlled proteolytic cascade (Figure 2).

While components of the Drosophila Toll pathway were identified in earlier genetic screens for developmental mutants, those in the Imd pathway have been the focus of more recent studies, mainly in the context of Drosophila immunity (Hoffmann and Reichhart 2002; Hultmark 2003). The effector NF-κB transcription factor of the Imd pathway is Relish, which upon immune activation is cleaved by the Dredd caspase (Figure 2). Using a combination of the RNA interference approach of silencing gene function and a high-throughput cell culture assay, Foley and O’Farrell (2004) report the identification of two new conserved members of this Imd pathway. Sickie is a novel protein required for Relish activation, and Defense repressor 1 is a novel inhibitor of the Dredd caspase.

The impressive progress in our understanding of innate immunity in Drosophila is now guiding scientists to explore the immune system of other insects such as the mosquito, Anopheles gambiae, that spreads human malaria. Immune responses in this mosquito are linked to the elimination of the malarial parasites (Osta et al. 2004). A comparison of the immunity-related genes in Anopheles and Drosophila reveals the presence of the Toll signaling pathway in the mosquito genome, even though there are some differences in genes encoding pathogen recognition and signal transduction molecules (Christophides et al. 2002). A detailed and comparative view of the genetic mechanisms underlying their host defense will contribute to the identification of new targets for insecticide development, and provide opportunities for controlling the transmission of pathogens.

Concluding Remarks

The homologs of many genes involved in innate immune responses in flies and humans have also been found in mice, sharks, nematodes, and plants (e.g., Pujol et al. 2001; Nurnberger and Brunner 2002). In species studied to date, host defense appears to be mediated by homologous proteins. Taken together, these findings suggest that the regulatory mechanisms of host defense may be hard-wired in the genome much as DNA replication and cell division are. Protein motifs, domains, and signaling elements have, for millions of years, not only retained their ancestral biochemical features but have also continued to participate in similar physiological responses. It is crucial that our evolving knowledge of “genomic recycling” be used to enhance our understanding of the evolution of humans, not only in the context of “descendants of ancient apes,” but in the larger context of our fundamental unity and shared genetic history with all other species. This simple but fundamental idea has yet to be adopted by the majority of our students and teachers. Unless we do more to overcome resistance to the idea that humans share deep evolutionary connections with all animal life, students will become increasingly isolated from an understanding of, and participation in, the genomics and bioinformatics revolution that is transforming the biological and biomedical sciences.

References

Brennan CA, Anderson KV (2004) Drosophila: The genetics of innate immune recognition and response. Annu Rev Immunol 22: 457–483.
Beutler B, Rehli M (2002) Evolution of the TIR, tolls and TLRs: Functional inferences from computational biology. Curr Top Microbiol Immunol 270: 1–21.
Christophides GK, Zdobnov E, Barillas-Mury C, Birney E, Blandin S, et al. (2002) Immunity-related genes and gene families in Anopheles gambiae. Science 298: 159–165.
Crozatier M, Ubeda JM, Vincent A, Meister M (2004) Cellular immune response to parasitization in Drosophila requires the EBF orthologue Coller. PLoS Biol 2(8): e196 DOI: 10.1371/journal.pbio.0020196.
De Gregorio E, Spellman PT, Tzou P, Rubin GM, Lemaître B (2002) The Toll and Imd pathways are the major regulators of the immune response in Drosophila. EMBO J 21: 2568–2579.
Evans CJ, Banerjee U, Hartenstein V (2003) Thicker than blood: Conserved mechanisms in Drosophila and vertebrate hematopoiesis. Dev Cell 5: 673–690.
Foley E, O’Farrell PH (2004) Functional dissection of an innate immune response by a genome-
wide RNAi screen. PLoS Biol 2(8): e203 DOI: 10.1371/journal.pbio.0020203.

Gould SJ (2002) The structure of evolutionary theory. Cambridge (Massachusetts): Harvard University Press. 1433 p.

Hoffmann JA, Reichhart JM (2002) Drosophila innate immunity: An evolutionary perspective. Nature Immunol 3: 121–126.

Hultmark D (2003) Drosophila immunity: Paths and patterns. Curr Immunol Rev 15: 12–19.

Janeway CA Jr (1989) Approaching the asymptote? Evolution and revolution in immunology. Cold Spring Harb Symp Quant Biol 54: 1–13.

Meister M (2004) Blood cells of Drosophila: Cell lineages and role in host defense. Curr Immunol Rev 16: 10–15.

Nurnberger T, Brunner F (2002) Innate immunity in plants and animals: Emerging parallels between the recognition of general elicitors and pathogen-associated molecular patterns. Curr Opin Plant Biol 5: 318–324.

O斯塔 MA, Christophides GK, Kafatos FC (2004) Effects of mosquito genes on Plasmodium development. Science 305: 2030–2032.

Pujol N, Link EM, Liu LX, Kurz CL, Alleong G, et al. (2001) A reverse genetic analysis of components of the Toll signaling pathway in Caenorhabditis elegans. Curr Biol 11: 809–821.

Salemi M, Vandamme AM (2003) The phylogenetic handbook: A practical approach to DNA and protein phylogeny. Cambridge: Cambridge University Press. 450 p.

Rubin GM, Yandell MD, Wortman JR, Gabor Miklos GL, Nelson CR, et al. (2000) Comparative genomics of the eukaryotes. Science 287: 2204–2215.

Theopold U, Schmidt O, Soderhall K, Dushay MS (2004) Coagulation in arthropods: Defense, wound closure and healing. Trends Immunol 25: 289–294.

Van Valen L (1975) A new evolutionary law. Evol Theor 1: 1–30.