

GENES AND DEATH: THE EVOLUTIONARY TRADE-OFF

Mutations, sexual reproduction, and death are the cornerstones of evolution. But is death a programmed necessity or an evolutionary accident?

Organisms like the Hydra (specifically strain 105) have seemingly evaded aging during their 50 years in controlled laboratory settings; however, in the wild, death remains an almost inevitable and universal law of nature.

It is often suggested that evolution selected "genes for death" to benefit the species, but from a Darwinian perspective, this is a contradiction: natural selection favors individual survival and reproduction, not self-destruction.

A recent study by Sanchez et al. in *Nature* (1) resolves this paradox through the lens of antagonistic pleiotropy. The researchers identified a genetic axis in mice, *Foxo1* and its effector *Trim63* (*MuRF1*), that acts as a double-edged sword: In Youth: These genes are essential for disease tolerance. During severe infection (sepsis), they protect the heart from pathological remodeling and multi-organ injury, ensuring the individual survives long enough to reproduce. In Old Age: The same mechanism becomes maladaptive. In older mice, *Foxo1* and *Trim63* switch roles, driving the very cardiac damage and pathogenesis that leads to death.

This confirms a key rule of evolution: traits that provide a massive survival advantage during our reproductive prime are favored by selection, even if those same traits become the "executioners" that make us decrepit later in life. Death, then, is not the goal, but the price we pay for the defenses that kept us alive when it mattered most.

1. <https://www.ncbi.nlm.nih.gov/pubmed/41535469>