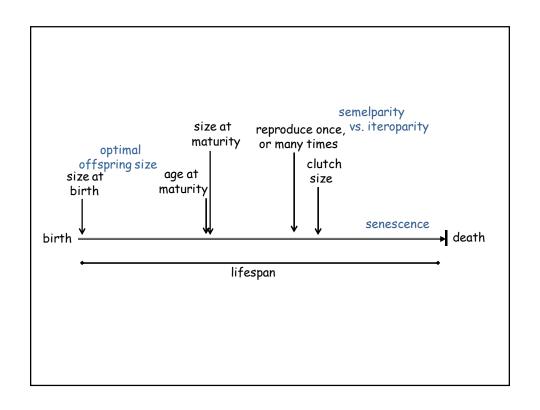


# Outline

senescence

what are life history traits?
trade-offs between life history traits





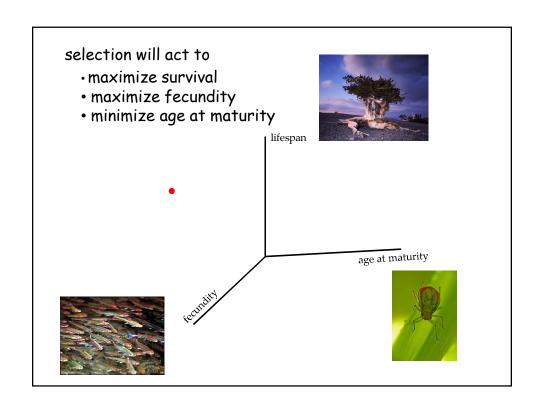
At what age and size should an individual become mature?

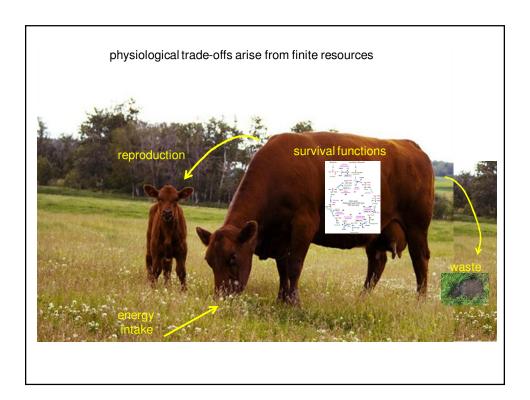
Should it reproduce once or many times?

How many offspring should it produce at one time?

What size should offspring be?

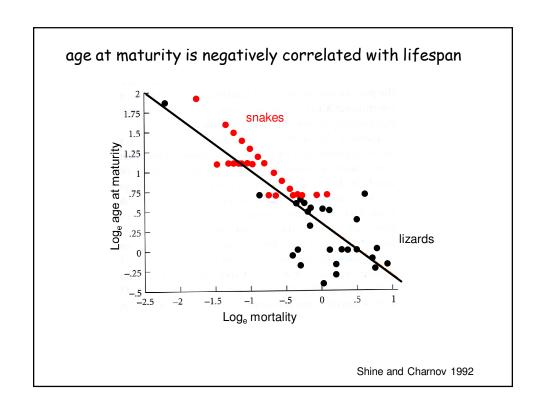


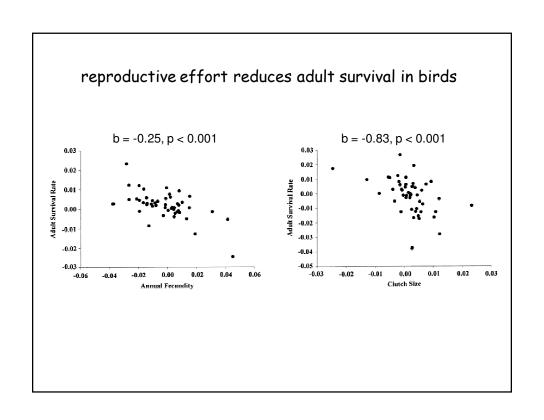


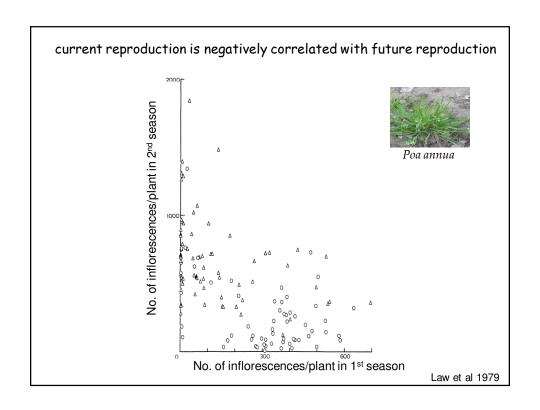


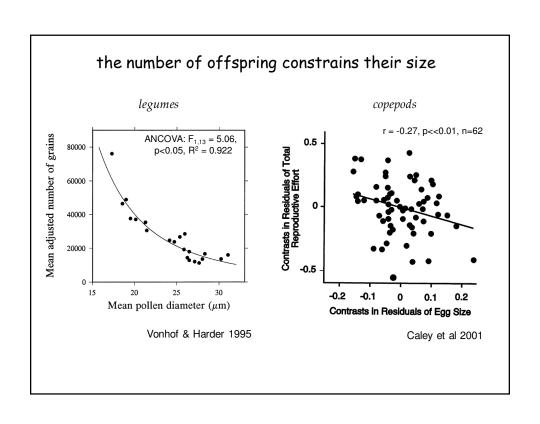
frequently observed trade-offs between life history characters

age at maturity, lifespan (-)
growth rate, size at maturity (+)
offspring size, clutch size (-)
clutch size, number of clutches (-)
lifespan, number of clutches (+)



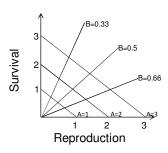






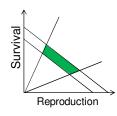
# why don't we always see an expected trade-off?

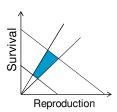
• resource acquisition vs. resource allocation



A = acquired energy

B = allocation to reproduction





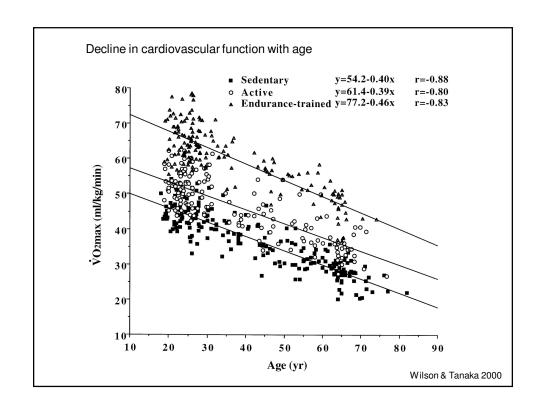
Van Noordwijk and de Jong 1986

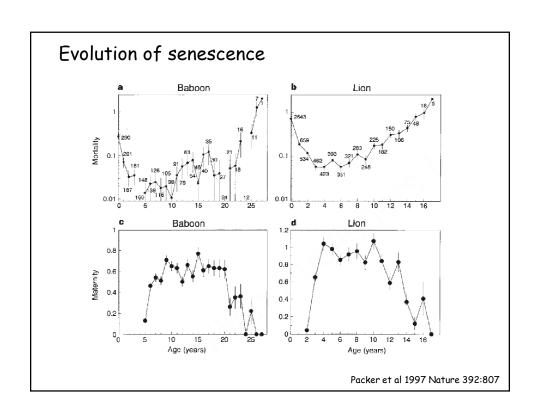
# Evolution of senescence



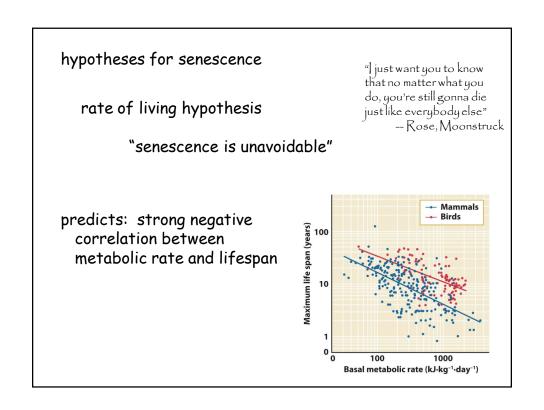
Senescence: greater susceptibility to injuries, disease and death as one grows older

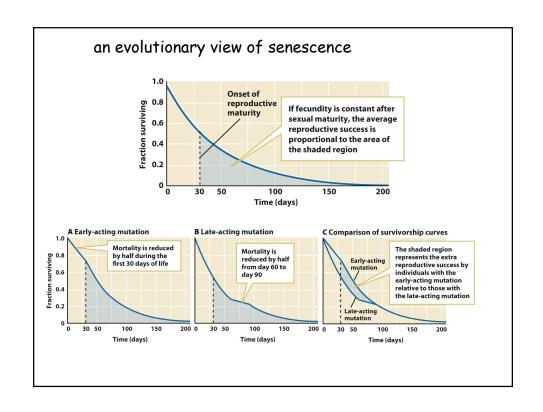
- neural degeneration
- reduced kidney filtrationdecreased respiratory capacity

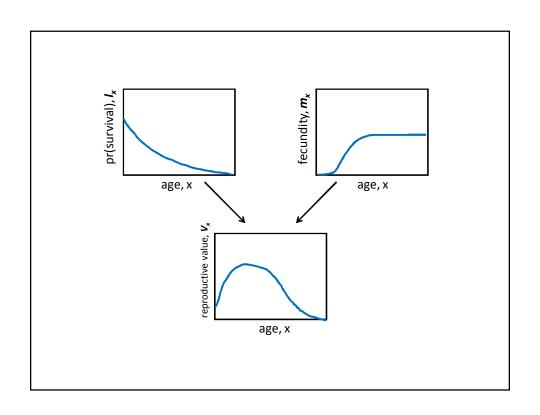




# hypotheses for senescence rate of living hypothesis "senescence is unavoidable" predicts: little/no remaining additive variance in the rate of senescence white the the the type under type under the type under the







### Mutation Accumulation hypothesis



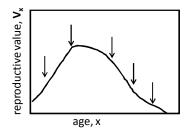
Medawar (1946):

"What is important from our point of view is that the contribution that each age-class makes to the ancestry of the future decreases with age."

reduced selection on older age classes

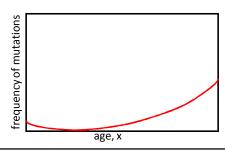
new, deleterious mutations build up in the genome early acting mutations are removed by selection late-acting ones are shielded from selection if post-reproductive

### Mutation Accumulation hypothesis



effect of deleterious mutations on fitness depends on the age at which they are expressed

weight by  $V_x$ 



mutations accumulate later in life because little detrimental effect on fitness

selection on interacting loci to shift time of expression

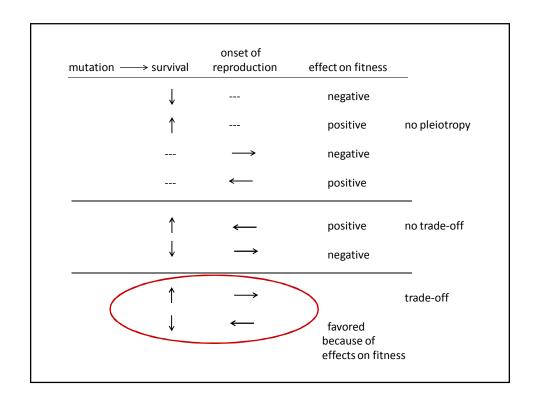
## Antagonistic Pleiotropy

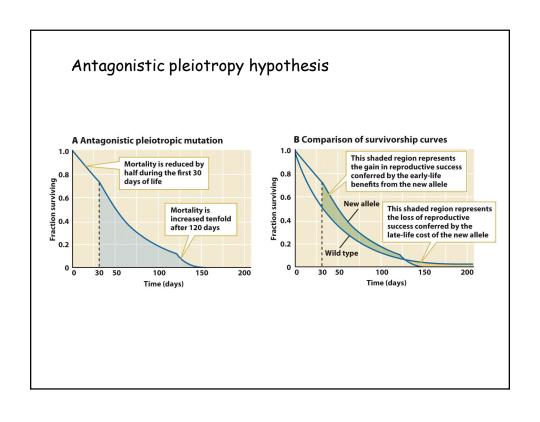


Williams (1957)

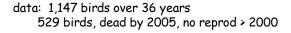
"Selection of a gene that confers an advantage at one age and a disadvantage at another will depend not only on the magnitudes of the effects themselves, but also on the times of the effects. An advantage during the period of maximum reproductive probability would increase the total reproductive probability more than a proportionately similar disadvantage later on would decrease it."

# Antagonistic Pleiotropy hypothesis trade-offs between early- and late- age effects fecundity age, x

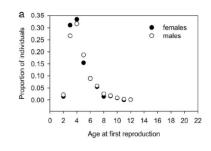


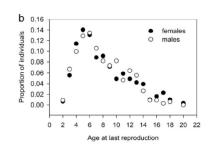


### evidence for antagonistic pleiotropy in mute swans

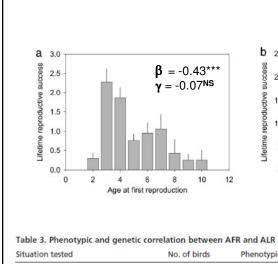


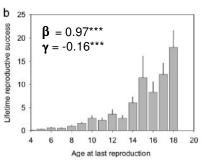






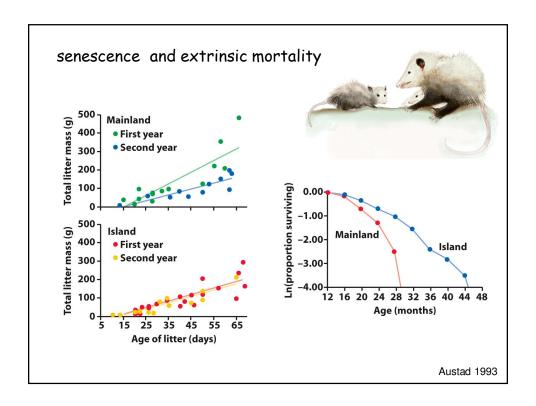
Charmantier et al 2006 PNAS





Situation tested	No. of birds	Phenotypic correlation $\pm$ SE	Genetic correlation $\pm$ SE
All breeding swans dead by 2005	648	0.193 ± 0.051***	0.555 ± 0.291*
All breeding swans dead by 2005, and with ALR > AFR	469	0.118 ± 0.060*	$0.560 \pm 0.404$
All breeding swans dead by 2005, and with age at death > ALR	272	0.191	0.495

Charmantier et al 2006 PNAS

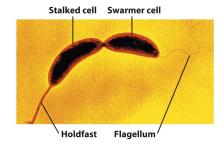


trade-off between investment in reproduction and investment in repair

somatic cells vs. germline cells
transcriptional and translational machinery
growth rate

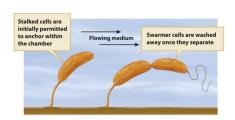
"disposable soma"

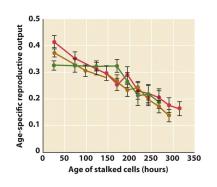
# a test of the reproduction-repair trade-off in bacteria



Caulobacter crescentus

starts as free-swimming swarmer
matures → stalked cell





Life history traits are those which are concerned with "decisions" about allocation of resources to maintenance, growth or reproduction

Life history traits often show trade-offs or correlations -- e.g., increased size of offspring may be related to a decreased number of offspring -- these trade-offs represent constraints on the evolution of life history characters

Iteroparity (multiple clutches) evolves when the relative survival rate of adults is high compared to juveniles. Low adult survival favors maximum investment in reproduction (semelparity).

Senescence evolves because selection on deleterious mutations changes as a function of age; mutations that enhance early reproduction (even to the detriment of lifespan) will be favored by selection