# Long term stationary phase in SBW25, REL606 and MG1655: notes from the literature

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#### Preliminary notes

• REL606 is supposed to become senescent after 16 days (Książek 2010) (I have not found this in the paper, look deeper), and this has been confirmed by a previous experiment made by Jenna

#### Abbreviations

- LTSP: long term stationary phase
- GASP: growth advantage in stationary phase

### Bacterial aging (Książek 2010)

- Aging is a set of changes that impair the ability to maintain homeostasis
- In human somatic cells aging is manifested as inability to divide (replicative senescence)
  - The maximum number of times a cell can divide is given by the Hayflick limit
  - It does not depend on the cronological time
  - Senescence can be triggered with stressors (SIPS, stress induced premature senescence)
- Also S. cerevisiae shows senescence, probably mediated by the accumulation of cytoplasmic factors
- The main way of bacterial replication is binary fission, which is a simmetric process mediated by FtsZ
  - This simmetry has been traditionally considered as a proof that bacteria do not age
  - The simmetry of division in E. coli is not reflected at the level of cellullar organization
  - One of the daughter cells inherits pre-existing elements from the mother (old pole) while the other synthesizes these *de novo* (new pole)
  - The linear discendence of the old pole shows senescence, and its replicative lifespan is of about 100 divisions
  - The old pole elements are likely damaged DNA and proteins and fragments of cell wall

# (Westphal et al. 2018)

- Attenuation of RpoS is strongly selected at the begenning (after 1 day) in E. coli
- Mutations in *rho* and *rpoBC* (regulatory proteins) are favoured in LTSP
- The GASP phenotype does not appear in Terrific broth or Super borth (even a disadvantage appears!), but cells aged in them show GASP if transfeered to LB

# Differences between MG1655 and REL606 (Yoon et al. 2012)

- Only 4% of the genome is strain-specific
- It includes profages and recently transferred islands

- REL606 has an additional set of Type II secretion genes and D-arabinose utilization
- REL606 lacks the cluster fly for flagellar biosythesis and the very short patch repair system
- Different set of genes for the Qin prophage, O-antigen synthesis, catabolism of aromatic compounds, LPS oligosaccharide synthesis
  - REL606 has the hpa cluster for catabolism of 3- and 4-hydroxy phenyl acetic acid
  - MG1655 has the paa cluster for catabolism of phenyl acetic acid
- There are numerous gene disruptions caused by deletions, frameshifts, IS sequences
- The 2 strains grow similarly in LB but REL606 grows faster in minimal medium
- Negligible differences in the accumulation of byproducts in minimal vs complex medium
- At the trascription level, in REL606 highly expressed genes are those involved in replication, translation and nucleotide metabolism, while in MG1655 genes for motility, transcription and energy production
- Proteins that are more abundant in REL606 are those involved in amino acid biosynthesis and maltose metabolism
- On the contrary, in MG1655 are more abundant protein for amino acid degradation and stress-response
- REL606 releases more proteins in the medium in stationary phase
- REL606 is more susceptible to stressfull conditions caused by osmolarity, pH, salicylate and  $\beta$ -lactam antibiotics

#### References

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