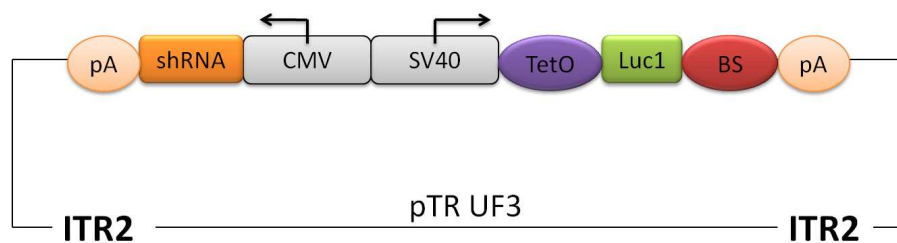


quick 'n' dirty cloning

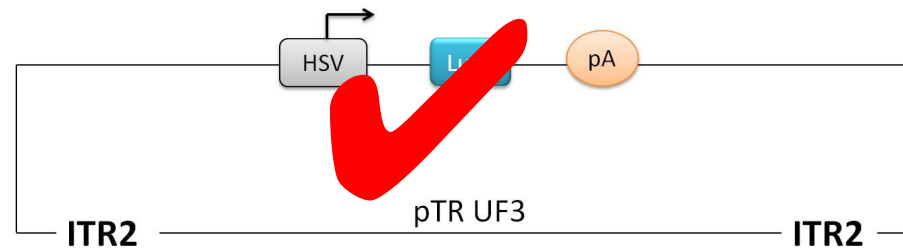
status quo
status quo

work progress

- first **promising** sequences for tuning plasmid
 - re-trial in parallel
 - plus: Phusion PCR strategy
- normalization construct **ready**
- ongoing cloning of TetR construct
- preparation of (endogenous + artificial) miRNA binding sites
- **problems** with cloning of shRNA into pcDNA5 for stable integration
 - re-do from the scratch
- likely: results by the end of this week!



“tuning plasmid“



“normalization plasmid“