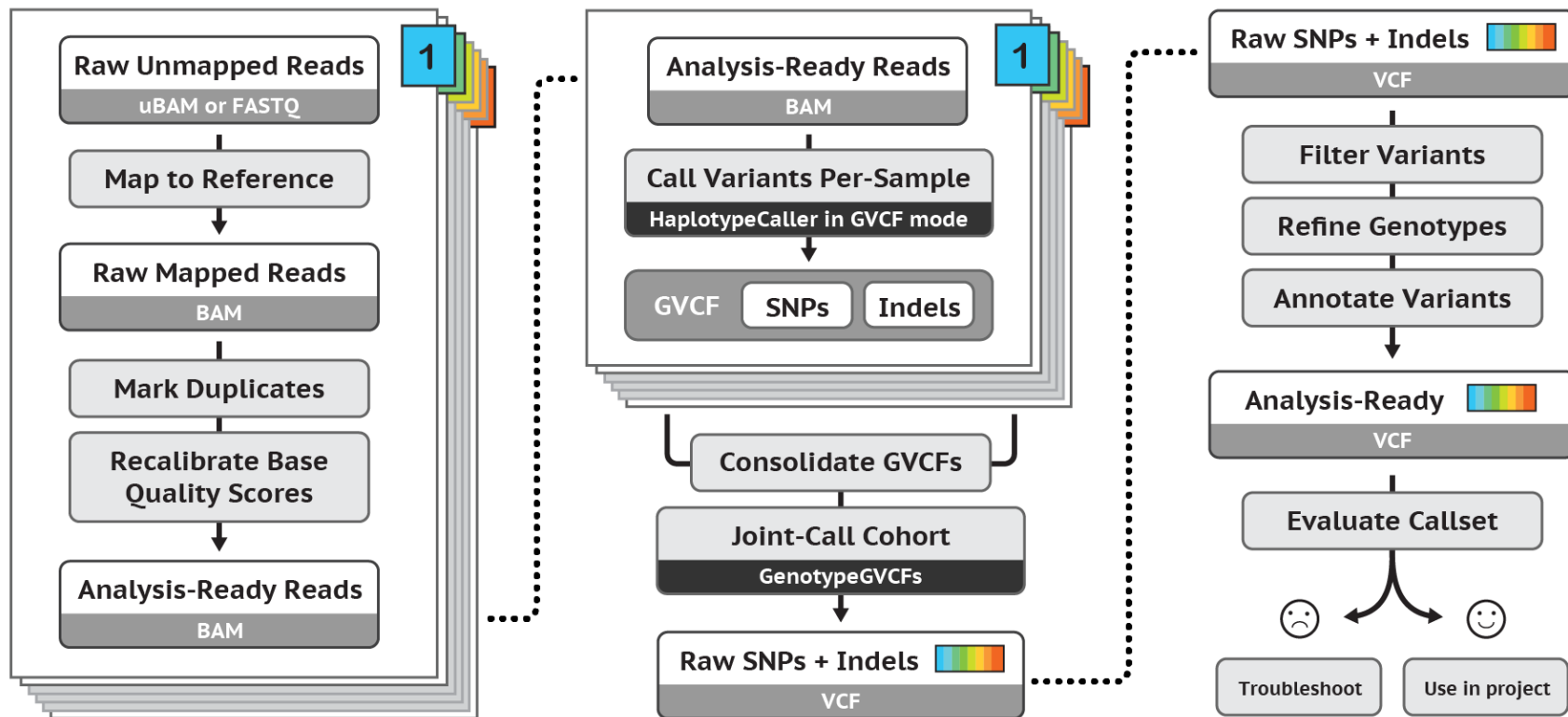




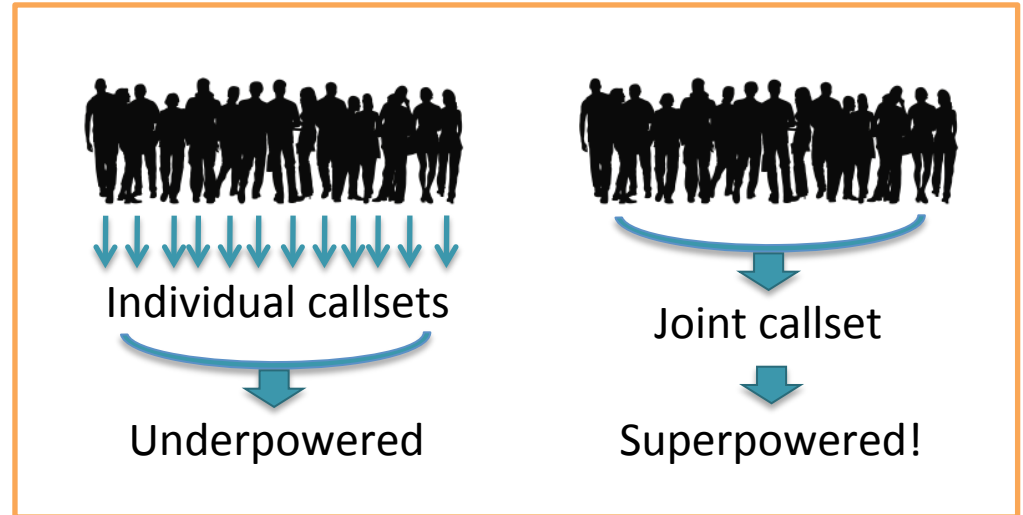
Germline Variant Discovery Highlights

Best Practices for Germline SNP & INDEL Discovery

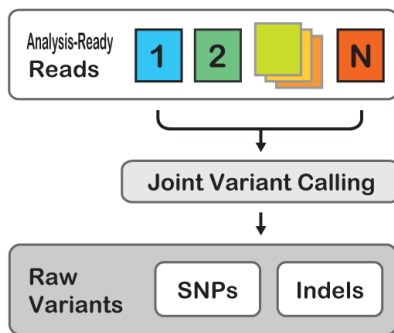


Joint analysis empowers discovery

- Single genome in isolation: almost never useful
- Family or population data add valuable information
 - rarity of variants
 - *de novo* mutations
 - ethnic background

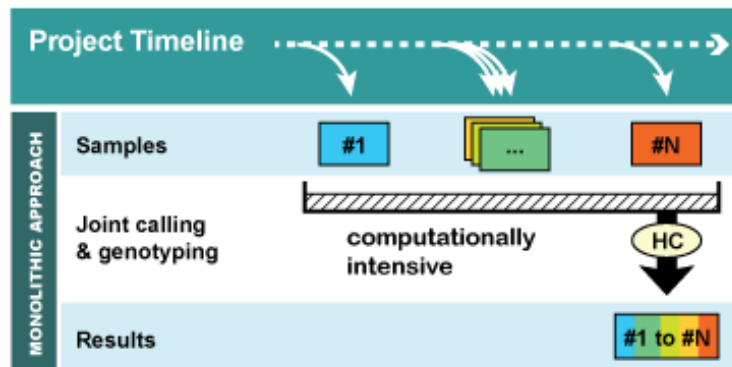


Traditional **multi-sample calling** approach : very inefficient



**Compute requirements
scale very badly with
number of samples!!!**

It gives us the right answers, but...

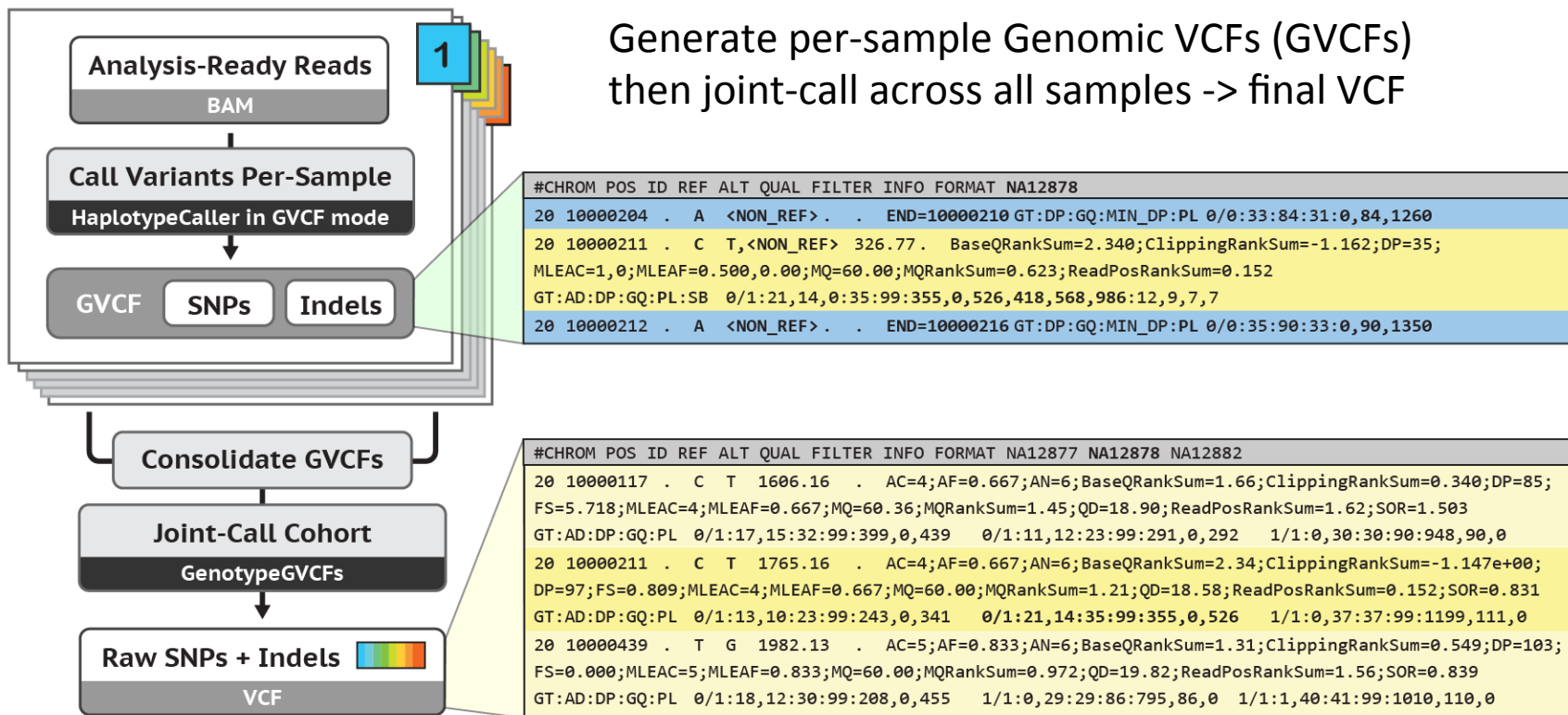


Want to add new samples?

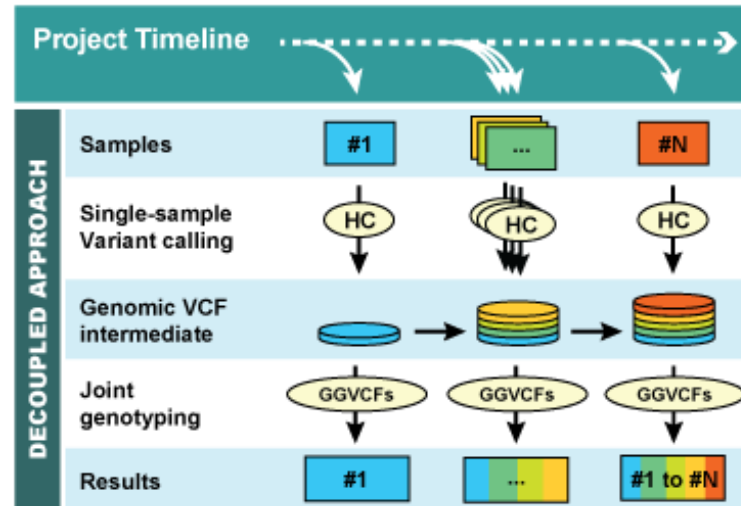
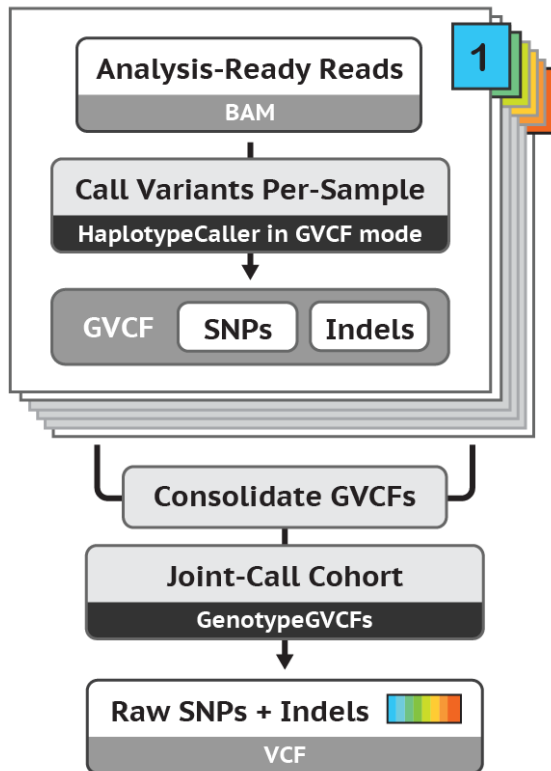
**Got to re-run pipeline from
scratch! The N+1 problem!**



Solution: the GVCf-based joint calling workflow



Same results as old approach - but scalable and incremental!



Scales linearly with number of samples!

Want to add a new sample? Make a GVCF for that sample then re-call the cohort at will!

Further refinements: filtering and more

