

lncRNA Literature Review

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- **AML WXS analysis**
- PAN cancer methylation
 - data downloading and preprocessing
 - STAD ($n = 479$), *PRAD* ($n = 304$)

- What we have?
 - 16 patients with matched normal, tumor and relapse sample, whole exome sequencing
- What's our hypothesis?
 - are there exist MRs that can predict relapse?
- What's been done?
 - potential MRs (from Yishai); Somatic mutations and CNVs of the 16 patients

Methods

Stochastic of finding mutations...

- Probability of at least 2 patients have the same mutation:
 - $M = \# \text{Mutations}; N = \# \text{ of patients}$
 - $1 - \frac{N! * N^M}{M^N}$
- Given the number of raw variants, p-value of at least 2 patients have 1 common mutation is $3.85 * 10^{-05}$

MRs associate with CNVs and Mutations

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MRs' association with Mutations using PrePPI

Mutated Genes with Minor Allele Frequency

Mutated Genes with Minor Allele Frequency (cont. . .)