# Lung Pre-cancer

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# **Contents**

1		oduction	3
	1.1	Lung Cancer	3
	1.2	Precancer	3
	1.3	Study Objectives	3
2	Mat	erials	3
-	2.1	List of IPNs	3
	2.1	2.1.1 Carcinoma in situ	3
		2.1.2 Adenocarcinoma <i>in situ</i>	3
		2.1.3 Atypical Adenomatous Hyperplasia	3
		2.1.4 Dysplasia	3
		2.1.5 Minimally Invasive Adenocarcinoma	3
	2.2	Data Structure & Count	4
	2.2	Data Structure & Count	_
3	Met	hods	4
	3.1	Workflows	4
4	Resu		4
	4.1	Quality Check with FastQC	4
	4.2	Quality Check with Sequenza	4
	4.3	Progressive Evolution at Single Nucleotide Level	4
	4.4	Macro-evolution at Chromosomal Level	4
	4.5	Selective Clonal Sweep during Neoplastic Evolution	4
	4.6	Cancer Gene Mutation during Cancer Evolution	4
	4.7	Distinct Drivers and Genetic Constraints in Multi-focal IPNs	4
5	Disc	eussion	4
6	Refe	erences	4
L	ist o	of Tables	
L	ist o	of Figures	
	1	Common cancer survival rates (Hong et al., 2021)	3
	2	Workflow for data pre-processing for variant discovery (Van der Auwera et al., 2013; DePristo et al., 2011)	4
	3	Somatic short variant discovery workflow (Van der Auwera et al., 2013; DePristo et al., 2011)	5
	4	Germline short variant discovery workflow (Van der Auwera et al., 2013; DePristo et al., 2011)	5
	5	RNA-seq short variant discovery workflow (Van der Auwera et al., 2013; DePristo et al., 2011)	6
	6	FastQC results with WES data	6
	7	FastOC results with WTS data	6

# 1 Introduction

## 1.1 Lung Cancer

Lung cancer is the most common form of cancer as 12.3 % of all cancers (Minna, Roth, & Gazdar, 2002).

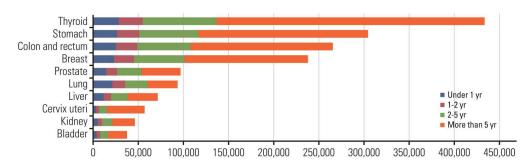


Figure 1: Common cancer survival rates (Hong et al., 2021)

#### 1.2 Precancer

# 1.3 Study Objectives

# 2 Materials

#### 2.1 List of IPNs

#### 2.1.1 Carcinoma in situ

Carcinoma in situ (CIS)

#### 2.1.2 Adenocarcinoma in situ

Adenocarcinoma in situ (AIS)

#### 2.1.3 Atypical Adenomatous Hyperplasia

Atypical adenomatous hyperplasia (AAH)

### 2.1.4 Dysplasia

#### 2.1.5 Minimally Invasive Adenocarcinoma

Minimally invasive adenocarcinoma (MIA)

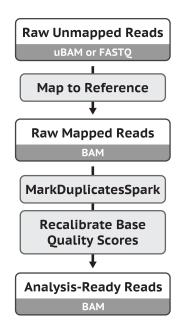


Figure 2: Workflow for data pre-processing for variant discovery (Van der Auwera et al., 2013; DePristo et al., 2011)

#### 2.2 Data Structure & Count

- 3 Methods
- 3.1 Workflows
- 4 Results
- 4.1 Quality Check with FastQC
- 4.2 Quality Check with Sequenza
- 4.3 Progressive Evolution at Single Nucleotide Level
- 4.4 Macro-evolution at Chromosomal Level
- 4.5 Selective Clonal Sweep during Neoplastic Evolution
- 4.6 Cancer Gene Mutation during Cancer Evolution
- 4.7 Distinct Drivers and Genetic Constraints in Multi-focal IPNs
- 5 Discussion

## 6 References

- DePristo, M. A., Banks, E., Poplin, R., Garimella, K. V., Maguire, J. R., Hartl, C., ... others (2011). A framework for variation discovery and genotyping using next-generation dna sequencing data. *Nature genetics*, 43(5), 491.
- Hong, S., Won, Y.-J., Lee, J. J., Jung, K.-W., Kong, H.-J., Im, J.-S., ... others (2021). Cancer statistics in korea: Incidence, mortality, survival, and prevalence in 2018. *Cancer Research and Treatment: Official Journal of Korean Cancer Association*, 53(2), 301.
- Minna, J. D., Roth, J. A., & Gazdar, A. F. (2002). Focus on lung cancer. Cancer cell, 1(1), 49-52.
- Van der Auwera, G. A., Carneiro, M. O., Hartl, C., Poplin, R., Del Angel, G., Levy-Moonshine, A., ... others (2013). From fastq data to high-confidence variant calls: the genome analysis toolkit best practices pipeline. *Current protocols in bioinformatics*, 43(1), 11–10.

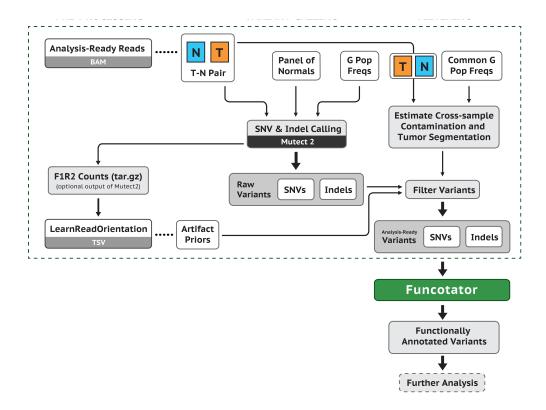


Figure 3: Somatic short variant discovery workflow (Van der Auwera et al., 2013; DePristo et al., 2011)

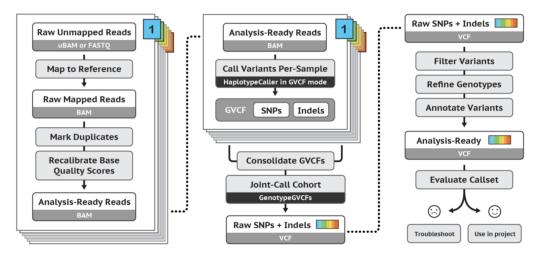


Figure 4: Germline short variant discovery workflow (Van der Auwera et al., 2013; DePristo et al., 2011)

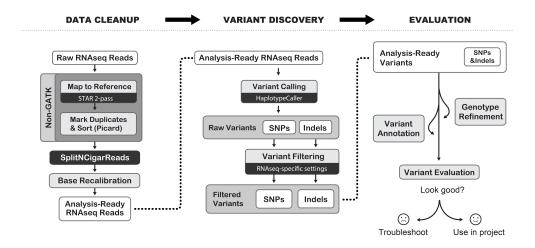


Figure 5: RNA-seq short variant discovery workflow (Van der Auwera et al., 2013; DePristo et al., 2011)

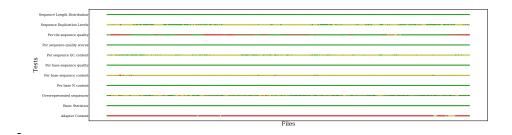


Figure 6: FastQC results with WES data

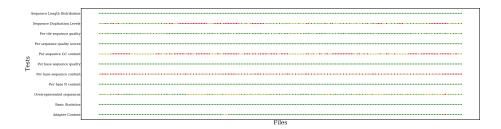


Figure 7: FastQC results with WTS data